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Office of Water

Development FINAL Document for Effluent Limitations Guidelines and Standards for the Nonferrous Metals Manufacturing Point Source Category

Volume VIII Primary Columbium – Tantalum Secondary Tantalum Secondary Uranium

Printed on Recycled Paper

ORGANIZATION OF THIS DOCUMENT

This development document for the nonferrous metals manufacturing category consists of a general development document which considers the general and overall aspects of the regulation and 31 subcategory specific supplements. These parts are organized into 10 volumes as listed below.

The information in the general document and in the supplements is organized by sections with the same type of information reported in the same section of each part. Hence to find information on any specific aspect of the category one would need only look in the same section of the general document and the specific supplements of interest.

The ten volumes contain contain the following subjects:

- Volume I General Development Document
- Volume II Bauxite Refining Primary Aluminum Smelting Secondary Aluminum Smelting
- Volume III Primary Copper Smelting Primary Electrolytic Copper Refining Secondary Copper Refining Metallurgical Acid Plants
- Volume IV Primary Zinc Primary Lead Secondary Lead Primary Antimony
- Volume V Primary Precious Metals and Mercury Secondary Precious Metals Secondary Silver Secondary Mercury
- Volume VI Primary Tungsten Secondary Tungsten and Cobalt Primary Molybdenum and Rhenium Secondary Molybdenum and Vanadium
- Volume VII Primary Beryllium Primary Nickel and Cobalt Secondary Nickel Secondary Tin
- Volume VIII Primary Columbium and Tantalum Secondary Tantalum Secondary Uranium
- Volume IX Primary and Secondary Titanium Primary Zirconium and Hafnium
- Volume X Primary and Secondary Germanium and Gallium Primary Rare Earth Metals Secondary Indium

DEVELOPMENT DOCUMENT

for

EFFLUENT LIMITATIONS GUIDELINES AND STANDARDS

for the

NONFERROUS METALS MANUFACTURING POINT SOURCE CATEGORY

VOLUME VIII

Primary Columbium and Tantalum Secondary Tantalum Secondary Uranium

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May 1989

U.S. Environmental Protection Agency Office of Water Office of Water Regulations and Standards Industrial Technology Division Washington, D. C. 20460

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For detailed contents see detailed contents list in individual supplement.

NONFERROUS METALS MANUFACTURING POINT SOURCE CATEGORY

DEVELOPMENT DOCUMENT SUPPLEMENT

for the

Primary Columbium and Tantalum Subcategory

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SECTION I

SUMMARY

This document provides the technical basis for promulgating effluent limitations based on best practicable technology (BPT) and best available technology (BAT) for direct dischargers, pretreatment standards for indirect dischargers (PSES), pretreatment standards for new indirect dischargers (PSNS), and standards of performance for new source direct dischargers (NSPS) for the primary columbium-tantalum subcategory.

The primary columbium-tantalum subcategory consists of five plants. Of the five plants, three discharge directly to rivers, lakes, or streams; two discharge to publicly owned treatment works (POTW); and none achieve zero discharge of process wastewater.

EPA first studied the primary columbium-tantalum subcategory to determine whether differences in raw materials, final products, manufacturing processes, equipment, age and size of plants, and water usage, required the development of separate effluent limitations and standards for different segments of the subcategory. This involved a detailed analysis of wastewater discharge and treated effluent characteristics, including (1) the sources and volume of water used, the processes employed, and the sources of pollutants and wastewaters in the plant; and (2) the constituents of wastewaters, including toxic pollutants.

Several distinct control and treatment technologies (both inplant and end-of-pipe) applicable to the primary columbiumtantalum subcategory were identified. The Agency analyzed both historical and newly generated data on the performance of these technologies, including their nonwater quality environmental impacts (air quality and solid waste generation) and energy requirements. EPA also studied various flow reduction techniques reported in the data collection portfolios (dcp) and plant visits.

Engineering costs were prepared for each of the control and treatment options considered for the subcategory. These costs then used by the Agency to estimate the impact of were implementing the various options on the industry. For each control and treatment option that the Agency found to be most effective and technically feasible in controlling the discharge pollutants, the number of potential closures, number of of employees affected, and impact on price were estimated. These results are reported in a separate document entitled The Economic Impact Analysis of Effluent Limitations and Standards for the Nonferrous Smelting and Refining Industry.

Based on consideration of the above factors, EPA identified various control and treatment technologies which formed the basis

for BPT and selected control and treatment appropriate for each set of standards and limitations. The mass limitations and standards for BPT, BAT, NSPS, PSES, and PSNS are presented in Section II.

After examining the various treatment technologies, the Agency has identified BPT to represent the average of the best existing technology. Metals removal based on lime precipitation and sedimentation technology is the basis for the BPT limitations. Steam stripping is selected as the technology basis for ammonia limitations. To meet the BPT effluent limitations based on this technology, the primary columbium-tantalum subcategory is expected to incur a capital cost of \$680,000 (1982 dollars) and annual cost of \$1,140,000 (1982 dollars).

For BAT, the Agency has built upon the BPT basis by adding inprocess control technologies which include recycle of process water from air pollution control and metal contact cooling wastewater streams. Filtration is added as an effluent polishing step to the end-of-pipe treatment scheme. To meet the BAT effluent limitations based on this technology, the primary columbium-tantalum subcategory is estimated to incur a capital cost of \$830,000 and an annual cost of \$1,202,000.

The best demonstrated technology (BDT), which is the technical basis of NSPS, is equivalent to BAT. In selecting BDT, EPA recognizes that new plants have the opportunity to implement the best and most efficient manufacturing processes and treatment technology. As such, the technology basis of BAT has been determined as the best demonstrated technology.

The Agency selected the same technology as BAT for PSES. To meet the pretreatment standards for existing sources, the primary columbium-tantalum subcategory is estimated to incur a capital cost of \$1,030,000 and an annual cost of \$701,000. For PSNS, the Agency selected end-of-pipe treatment and in-process flow reduction control techniques equivalent to NSPS.

SECT - II

SECTION II

CONCLUSIONS

EPA has divided the primary columbium-tantalum subcategory into 11 subdivisions for the purpose of effluent limitations and standards. These subdivisions are:

(a) Concentrate digestion wet air pollution control,

(b) Solvent extraction raffinate,

(c) Solvent extraction wet air pollution control,

(d) Precipitation and filtration,

(e) Precipitation and filtration wet air pollution control,(f) Tantalum salt drying,

(g) Oxides calcining wet air pollution control,

(h) Reduction of tantalum salt to metal,

(i) Reduction of tantalum salt to metal wet air pollution control,

(j) Tantalum powder wash and wet air pollution control, and

(k) Consolidation and casting contact cooling.

BPT is promulgated based on the effluent concentrations achievable by the application of chemical precipitation and sedimentation (lime and settle) technology, along with preliminary treatment consisting of ammonia steam stripping for selected waste streams. The following BPT effluent limitations are promulgated:

(a) <u>Concentrate Digestion</u> <u>Wet Air Pollution Control</u> BPT

				. •
Pollutant or Pollutant Proper	ty		Maximum for Any One Day	Maximum for Monthly Average
•	·			- 5
Metri	c Units - m	ng/kg of	concentrate di	heted
English Uni	ts - lbs/mi	llion 1	os of concentra	te digested
Lead	· · · ·		0 610	
		. I	2.612	1.244
Zinc			9.080	3.794
Ammonia (as N)	_ •		829.000	364.500
Fluoride			217.700	124.400
TSS			255.000	121.300
pH	Within t	he range	e of 7.0 to 10.	0 at all times
				· · ·

4331

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - II

		مربع المراجع ال
Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
Metric Units - m English Units - lbs/mi	ng/kg of concentrate di llion lbs of concentra	gested te digested
Lead Zinc Ammonia (as N) Fluoride TSS pH Within	3.845 13.370 1,221.000 320.400 375.400 the range of 7.0 to 10	
(c) Solvent Extraction Wet	Air Pollution Control	ВРТ
Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
Metric Units - m English Units - lbs/mi	ng/kg of concentrate di llion lbs of concentra	gested te digested
Lead Zinc Ammonia (as N) Fluoride Total Suspended Solids pH	1.032 3.586 327.400 85.960 100.700 Within the rang at all	
(d) Precipitation and Filt	ration BPT	
	g/kg of concentrate di	
•	llion lbs of concentra	-
Lead Zinc Ammonia (as N) Fluoride TSS . pH Within	5.570 19.990 1,825.000 479.100 561.300 the range of 7.0 to 10	2.738 8.350 802.200 273.800 267.000 .0 at all times

(b) Solvent Extraction Raffinate BPT

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - II
(e) Precipitation and Filtration Wet Air Pollution Control BI
Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average
Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested
Lead 26.680 12.700 Zinc 92.730 38.740 Ammonia (as N) 8,466.000 3,722.000 Fluoride 2,223.000 1,270.000 TSS 2,604.000 1,239.000 pH Within the range of 7.0 to 10.0 at all times
(f) <u>Tantalum</u> <u>Salt</u> <u>Drying</u> BPT
Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average
Metric Units - mg/kg of tantalum salt dried English Units - lbs/million lbs of tantalum salt dried
Lead 25.430 12.110 Zinc 88.390 36.930 Ammonia (as N) 8,070.000 3,548.000 Fluoride 2,119.000 1,211.000 TSS 2,482.000 1,181.000 PH Within the range of 7.0 to 10.0 at all times
(g) Oxides Calcining Wet Air Pollution Control BPT
Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average
Metric Units - mg/kg of columbium-tantalum oxide dried English Units - lbs/million lbs of columbium-tantalum oxide drie
Lead 16.140 7.685 Zinc 56.100 23.440 Ammonia (as N) 5,122.000 2,252.000 Fluoride 1,345.000 768.500 TSS 1,576.000 749.200 pH Within the range of 7.0 to 10.0 at all times

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - II

Pollutant or Pollutant Property		Maximum for Ionthly Average
Metric Units - mg/kg English Units - lbs/milli	of tantalum salt red on lbs of tantalum sa	
Lead Zinc Ammonia (as N) Fluoride TSS pH Within the	69.750 242.500 22,140.000 5,813.000 6,809.000 range of 7.0 to 10.0	3,322.000 3,239.000
(i) <u>Reduction</u> of <u>Tantalum</u> <u>Sal</u> <u>Control</u> BPT	t to Metal Wet Air Po	ollution
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day M	Nonthly Average
Metric Units - mg/kg	of tantalum salt red	luced
English Units - lbs/milli	on lbs of tantalum sa	lt reduced
Lead	0.858	0.409
Zinc	2.983	1.246
Ammonia (as N)	272.400	119.700
Fluoride	71.510	40.860
TSS	83.770	39.840
pH Within the	range of 7.0 to 10.0	at all times
(j) <u>Tantalum</u> Powder Wash BPT		
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day M	Nonthly Average
Metric Units - mg/kg	of tantalum powder w	ashed
English Units - lbs/millio	n lbs of tantalum pow	der washed
Lead	8.582	4.087
Zinc	29.830	12.470
Ammonia (as N)	2,724.000	1,198.000
Fluoride	715.200	408.700
TSS	837.800	398.500
pH Within the	range of 7.0 to 10.0	at all times

(h) <u>Reduction of Tantalum Salt to Metal</u> BPT

SECT - II

(k) Consolidation and Casting Contact Cooling BPT

Pollutant o Pollutant P			Maximum Any One		Maximum for onthly Average	3
	ts - 1bs				or consolidate talum cast or	Đ
Lead Zinc Ammonia (as Fluoride TSS pH	N)	Within the		0.000 0.000 0.000 0.000 0.000 0.000 0 to 10.0	0.000 0.000 0.000 0.000 0.000 at all times	

BAT is promulgated based on the effluent concentrations achievable by the application of chemical precipitation, sedimentation, and multimedia filtration (lime, settle, and filter) technology and in-process flow reduction control methods, along with preliminary treatment consisting of ammonia steam stripping for selected waste streams. The following BAT effluent limitations are promulgated:

(a) Concentrate Digestion Wet Air Pollution Control BAT

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day Mo	onthly Average
Metric Units - mg/kg English Units - lbs/millior		
Lead	0.174	0.081
Zinc	0.635	0.261
Ammonia (as N)	82.910	36.450
Fluoride	21.770	12.440
(b) Solvent Extraction Raffinat	се ВАТ	1

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg English Units - lbs/million		
Lead	2.564	1.190
Zinc	9.338	3.845
Ammonia (as N)	1,221.000	536.500
Fluoride	320.400	183.100

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - II

Pollutant or		um for		um for
Pollutant Property	Any O	ne Day	Monthly	Average
Metric Unit	s - mg/kg of conce	ntrate di	igested	
English Units - 1	os/million lbs of	concentra	ate digest	tea
Lead		0.069		0.032
Zinc		0.251		0.103
Ammonia (as N)		32.790		14.420
Fluoride		8.610		4.920
(d) Precipitation and	Filtration BAT			
Pollutant or	Maxim	um for	Maxim	um for
Pollutant Property			Monthly	
		ntrate di	gested	
Metric Unit:	s - mg/kg of conce			
	s - mg/kg of conce os/million lbs of			zed
English Units - 11 Lead		concentra 3.833		1.780
English Units - 1 Lead Zinc	os/million lbs of	concentra 3.833 13.960	ate digest	1.780 5.750
English Units - 11 Lead Zinc Ammonia (as N)	os/million lbs of 1,	concentra 3.833 13.960 825.000	ate digest	1.780 5.750 302.200
	os/million lbs of 1,	concentra 3.833 13.960	ate digest	1.780 5.750
English Units - 1 Lead Zinc Ammonia (as N) Fluoride	os/million lbs of 1,	concentra 3.833 13.960 825.000 479.100	ate digest	1.780 5.750 302.200 273.800
English Units - 1 Lead Zinc Ammonia (as N) Fluoride (e) <u>Precipitation</u> and	ps/million lbs of 1, <u>Filtration Wet Ai</u>	concentra 3.833 13.960 825.000 479.100 <u>r Polluti</u>	ate digest	1.780 5.750 302.200 273.800 01 BAT
English Units - 1 Lead Zinc Ammonia (as N) Fluoride (e) <u>Precipitation</u> and Pollutant or	os/million lbs of 1, <u>Filtration Wet Ai</u> Maxim	concentra 3.833 13.960 825.000 479.100	ate digest	1.780 5.750 302.200 273.800 <u>ol</u> BAT
English Units - 1 Lead Zinc Ammonia (as N) Fluoride (e) <u>Precipitation and</u> Pollutant or Pollutant Property <u>Metric Units</u>	os/million lbs of 1, <u>Filtration Wet Ai</u> Maxim	concentra 3.833 13.960 825.000 479.100 <u>r Polluti</u> um for ne Day ntrate di	ate digest	1.780 5.750 302.200 273.800 01 BAT 1m for Average
English Units - 1 Lead Zinc Ammonia (as N) Fluoride (e) <u>Precipitation and</u> Pollutant or Pollutant Property <u>Metric Units</u> English Units - 1	ps/million lbs of l, <u>Filtration Wet Ai</u> Maxim Any O s - mg/kg of conce	concentra 3.833 13.960 825.000 479.100 <u>r Polluti</u> um for ne Day ntrate di concentra	ate digest	1.780 5.750 302.200 273.800 <u>ol</u> BAT
English Units - 1 Lead Zinc Ammonia (as N) Fluoride (e) <u>Precipitation and</u> Pollutant or Pollutant Property <u>Metric Units</u> English Units - 1 Lead	ps/million lbs of l, <u>Filtration Wet Ai</u> Maxim Any O s - mg/kg of conce	concentra 3.833 13.960 825.000 479.100 <u>r Polluti</u> um for ne Day ntrate di	ate digest	1.780 5.750 302.200 273.800 <u>ol</u> BAT im for Average ced 0.826
English Units - 1 Lead Zinc Ammonia (as N) Fluoride (e) <u>Precipitation and</u> Pollutant or Pollutant Property <u>Metric Units</u>	ps/million lbs of 1, <u>Filtration Wet Ai</u> Maxim Any O 5 - mg/kg of conce ps/million lbs of	concentra 3.833 13.960 825.000 479.100 <u>r Polluti</u> um for ne Day ntrate di concentra 1.778	ate digest ton Contro Maximu Monthly lgested ate digest	1.780 5.750 302.200 273.800 <u>ol</u> BAT

(c) Solvent Extraction Wet Air Pollution Control BAT

	11 1 ⁵				
PRIMARY COLUMBIUM AND	TANTALUM	SUBCATEGO	ORY	SECT -	II
		· · · · · · ·		•	
(f) Tantalum Salt Drying	ВАТ		ав.		
$(1) \underline{10110010m} \underline{D010} \underline{D1} \underline{D1}$					
······································	•	·			
Pollutant or		Maximum i	for	Maxi	mum for
Pollutant Property	, ³ • • •	Any One I			y Avera
			1		<i>j</i> 110010
Metric Units -	ma/ka of	tantalum	salt	dried	
English Units - 1bs	/million	lbs of tar	ntalun	n salt d	ried
Lead		16	.950		7.87
Zinc			.750		25.43
Ammonia (as N)		8,070		3	,548.00
Fluoride		2,119			,211.00
		27110	• 0,00	<u>т</u>	/ 211.00
······································					·····
(g) Oxides Calcining Wet	Air Poll	ution Cont	trol	BAT	
					· · ·
	Carl Contract Contract	· · · ·			
			• <u> </u>	· · · · · · · · · · · · · · · · · · ·	
	1 1	Maximum 1	for	Maxi	mum for
Pollutant or		Maximum 1 Any One I			mum for v Avera
		Maximum f Any One I		Maxi Monthl	
Pollutant or Pollutant Property	q of colu	Any One I	Day	Monthl	y Avera
Pollutant or		Any One I	Day talum	Monthl oxide d	y Avera
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio		Any One I mbium-tant columbiur	Day talum m-tant	Monthl oxide d	y Avera ried ide dri
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead		Any One I mbium-tant columbiur 1	Day talum m-tant .076	Monthl oxide d	y Avera ried ide dri 0.50
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio		Any One I mbium-tant columbium 1 3	Day talum m-tant .076 .919	Monthl oxide d	y Avera ried ide dri 0.50 1.61
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead		Any One I mbium-tant columbium 1 3	Day talum m-tant .076	Monthl oxide d	y Avera ried ide dri 0.50
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc		Any One I mbium-tant columbiur 1 3 512	Day talum m-tant .076 .919	Monthl oxide d alum ox	y Avera ried ide dri 0.50 1.61
Pollutant or Pollutant Property Metric Units - mg/ko English Units - 1bs/millio Lead Zinc Ammonia (as N)		Any One I mbium-tant columbiur 1 3 512	Day talum m-tant .076 .919 .200	Monthl oxide d alum ox	y Avera ried ide dri 0.50 1.61 225.20
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride	on lbs of	Any One I mbium-tant columbium 1 3 512 134	Day talum m-tant .076 .919 .200 .500	Monthl oxide d alum ox	y Avera ried ide dri 0.50 1.61 225.20
Pollutant or Pollutant Property Metric Units - mg/ko English Units - 1bs/millio Lead Zinc Ammonia (as N)	on lbs of	Any One I mbium-tant columbium 1 3 512 134	Day talum m-tant .076 .919 .200	Monthl oxide d alum ox	y Avera ried ide dri 0.50 1.61 225.20
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride	on lbs of	Any One I mbium-tant columbium 1 3 512 134	Day talum m-tant .076 .919 .200 .500	Monthl oxide d alum ox	y Avera ried ide dri 0.50 1.61 225.20
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalur</u>	on lbs of	Any One I mbium-tant columbium 1 3 512 134 <u>Metal</u> H	Day talum m-tant .076 .919 .200 .500 BAT	Monthl oxide d alum ox	y Avera ried ide dri 0.50 1.61 225.20 76.84
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalur</u> Pollutant or	on lbs of	Any One I mbium-tant columbiur 1 3 512 134 <u>Metal</u> I Maximum f	Day talum m-tant .076 .919 .200 .500 BAT	Monthl oxide d alum ox Maxii	y Avera ried ide dri 0.50 1.61 225.20 76.84
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalur</u>	on lbs of	Any One I mbium-tant columbium 1 3 512 134 <u>Metal</u> H	Day talum m-tant .076 .919 .200 .500 BAT	Monthl oxide d alum ox Maxii	y Avera ried ide dri 0.50 1.61 225.20 76.84
Pollutant or Pollutant Property Metric Units - mg/ke English Units - lbs/millie Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalu</u> Pollutant or Pollutant Property	on lbs of <u>m Salt to</u>	Any One I mbium-tant columbiur 1 3 512 134 <u>Metal</u> I Maximum f Any One I	Day talum m-tant .076 .919 .200 .500 BAT for Day	Monthl oxide d alum ox Maxin Monthl	y Avera ried ide dri 0.50 1.61 225.20 76.84
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalur</u> Pollutant or Pollutant property Metric Units - r	on lbs of <u>m Salt to</u> ng/kg of	Any One I mbium-tant columbiur 1 3 512 134 <u>Metal</u> H Maximum f Any One I tantalum s	Day talum m-tant .076 .919 .200 .500 BAT for Day salt r	Monthl oxide d alum ox Maxi Monthl educed	y Avera ried ide dri 0.50 1.61 225.20 76.84 mum for y Avera
Pollutant or Pollutant Property Metric Units - mg/ke English Units - lbs/millie Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalu</u> Pollutant or Pollutant Property	on lbs of <u>m Salt to</u> ng/kg of	Any One I mbium-tant columbiur 1 3 512 134 <u>Metal</u> H Maximum f Any One I tantalum s	Day talum m-tant .076 .919 .200 .500 BAT for Day salt r	Monthl oxide d alum ox Maxi Monthl educed	y Avera ried ide dri 0.50 1.61 225.20 76.84 mum for y Avera
Pollutant or Pollutant Property Metric Units - mg/kd English Units - lbs/millid Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalu</u> Pollutant or Pollutant or Pollutant Property Metric Units - m English Units - lbs/m	on lbs of <u>m Salt to</u> ng/kg of	Any One I mbium-tant columbiur 1 3 512 134 <u>Metal</u> I Maximum f Any One I tantalum s bs of tant	Day talum m-tant .076 .919 .200 .500 BAT for Day salt r talum	Monthl oxide d alum ox Maxi Monthl educed	y Avera ried ide dri 0.50 1.61 225.20 76.84 mum for y Avera duced
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalur</u> Pollutant or Pollutant or Pollutant Property Metric Units - m English Units - lbs/m	on lbs of <u>m Salt to</u> ng/kg of	Any One I mbium-tant columbiur 1 3 512 134 <u>Metal</u> I Maximum f Any One I tantalum s bs of tant 46.	Day talum m-tant .076 .919 .200 .500 BAT for Day salt r talum .500	Monthl oxide d alum ox Maxi Monthl educed	y Avera ried ide dri 0.50 1.61 225.20 76.84 mum for y Avera duced 21.59
Pollutant or Pollutant Property Metric Units - mg/kd English Units - lbs/millid Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalur</u> Pollutant or Pollutant or Pollutant Property Metric Units - m English Units - lbs/m Lead Zinc	on lbs of <u>m Salt to</u> ng/kg of	Any One I mbium-tant columbium 1 3 512 134 <u>Metal</u> I Maximum f Any One I tantalum s bs of tant 46. 169.	Day talum m-tant .076 .919 .200 .500 BAT for Day salt r talum .500 .400	Monthl oxide d alum ox Maxi Monthl educed salt re	y Avera ried ide dri 0.50 1.61 225.20 76.84 mum for y Avera duced 21.59 69.75
Pollutant or Pollutant Property Metric Units - mg/ko English Units - lbs/millio Lead Zinc Ammonia (as N) Fluoride (h) <u>Reduction of Tantalur</u> Pollutant or Pollutant or Pollutant Property Metric Units - m English Units - lbs/m	on lbs of <u>m Salt to</u> ng/kg of	Any One I mbium-tant columbiur 1 3 512 134 <u>Metal</u> I Maximum f Any One I tantalum s bs of tant 46.	Day talum m-tant .076 .919 .200 .500 .500 BAT for Day salt r talum .500 .400 .000	Monthl oxide d alum ox Maxin Monthl educed salt re	y Avera ried ide dri 0.50 1.61 225.20 76.84 mum for y Avera duced 21.59

(i) <u>Reduction of Tantalum Salt to Metal Wet Air Pollution</u> <u>Control BAT</u>

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
 		- 2
Metric Units - mg English Units - lbs/mi	/kg of tantalum salt i llion lbs of tantalum	salt reduced
Lead	0.572	0.266
Zinc	2.084	0.858
Ammonia (as N)	272.400	119.700
Fluoride	71.510	40.860
(j) <u>Tantalum</u> <u>Powder</u> <u>Wash</u>	BAT	
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg English Units - lbs/mil	/kg of tantalum powder lion lbs of tantalum p	
Lead	5.721	2.656
Zinc	20.840	8.582
Ammonia (as N)	2,724.000	1,198.000
fluoride	715.200	408.700
(k) <u>Consolidation</u> and <u>Cast</u>	ing Contact Cooling	BAT
Pollutant or	Maximum for	Maximum for
Pollutant Property		Monthly Average
Metric Units - mg/kg of col English Units - lbs/million consolidate	lbs of columbium or t	
Lead	0.000	0.000
Linc	0.000	0.000
Namonio (og NI)	0.000	0.000
Ammonia (as N) Fluoride	0.000	0.000

NSPS are promulgated based on the effluent concentrations achievable by the application of chemical precipitation, sedimentation, and multimedia filtration (lime, settle, and filter) technology and in-process flow reduction control methods, along with preliminary treatment consisting of ammonia steam stripping for selected waste streams. The following effluent standards are promulgated for new sources: PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - II

(a)	Concer	ntrate	Digestion	<u>Wet</u>	<u>Air</u>	Polluti	on <u>Con</u>	trol NS	PS	
	utant d utant 1		гу			Maximum Any One				m for Average
	Englis		: Units - s - 1bs/r							ed
Lead Zinc Ammor Fluo TSS pH	nia		Withir	n the	range	8	0.174 0.635 2.910 1.770 9.330 0 to 1	0_0 at		0.081 0.261 36.450 12.440 7.464 times
(b)	Solver	nt Exti	action Ra	fin	ate Na	SPS				
	utant d utant I		-y		-	Maximum Any One				m for Average
	Englis	Metric sh Unit	: Units - ts - 1bs/n	mg/ko nillio	g of o on lba	concent s of co	rate d ncentr	igested ate dig	est	ed
Lead Zinc Ammor Fluo: TSS pH	nia (as ride	5 N)	Withir	n the	range	1,22 32 13	0.400 7.300	•	1 1	83.100 09.900
(c)	Solver	nt Extr	action We	et Ai	r <u>Pol</u>	Lution (Contro	1 NSPS		
	utant d utant H		У			Maximum Any One				m for Average
	Englis	Metric sh Unit	: Units - :s - 1bs/m	mg/ko nillio	g of o on lba	concenti s of con	rate d ncentr	igested ate dig	est	ed
Lead Zinc Ammon Fluo: TSS pH	nia (as ride	5 N)	Withir	1 the	range	32).069).251 2.790 3.610 3.690) to 1	0.0 at		0.032 0.103 14.420 4.920 2.952 times

(d) Precipitation and Filtration NSPS

Pollutant or Pollutant Property	Maximum for Maximum for Marimum for Marimum Marimum Marimum Marimum Marimum Marimum Marimum Marimum Marimum Mar	Maximum for hthly Average
Metric Units - mo	g/kg of concentrate digest	ted
English Units - lbs/mil	llion lbs of concentrate o	ligested
Lead	3.833	1.780
Zinc	13.960	5.750
Ammonia (as N)	1,825.000	802.200
Fluoride	479.100	273.800
TSS	205.400	164.300
pH Within t	the range of 7.0 to 10.0 a	at all times
(e)Precipitation and Filtrat	tion Wet Air Pollution Con	ntrol NSPS
Pollutant or	Maximum for M	Maximum for
Pollutant Property	Any One Day Mor	hthly Average
Metric Units - mo	g/kg of concentrate digest	ted
English Units - 1bs/mil	llion lbs of concentrate o	ligested
Lead	1.778	0.826
Zinc	6.478	2.668
Ammonia (as N)	846.600	372.200
Fluoride	222.300	127.000
TSS	95.270	76.210
pH Within t	che range of 7.0 to 10.0 a	at all times
(f) <u>Tantalum Salt</u> Drying NS	SPS	
Pollutant or	Maximum for M	Maximum for
Pollutant Property	Any One Day Mor	hthly Average
	y/kg of tantalum salt drie illion lbs of tantalum sal	
Lead	16.950	7.871
Zinc	61.750	25.430
Ammonia (as N)	8,070.000	3,548.000
Fluoride	2,119.000	1,211.000
TSS	908.200	726.500
pH Within t	che range of 7.0 to 10.0 a	at all times

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - II Oxides Calcining Wet Air Pollution Control NSPS (q) Maximum for Maximum for Pollutant or Monthly Average Pollutant Property Any One Day Metric Units - mg/kg of columbium-tantalum oxide dried English Units - lbs/million lbs of columbium-tantalum oxide dried Lead 1.076 0.500 1.614 Zinc 3.919 Ammonia (as N) 512.200 225.200 Fluoride 134.500 76.840 TSS 57.630 46.110 Within the range of 7.0 to 10.0 at all times рH Reduction of Tantalum Salt to Metal NSPS (h) Maximum for Maximum for Pollutant or Pollutant Property Any One Day Monthly Average Metric Units - mg/kg of tantalum salt reduced English Units - lbs/million lbs of tantalum salt reduced Lead 46.500 21.590 Zinc 169.400 69.750 Ammonia (as N) 22,140.000 9,732.000 Fluoride 5,813.000 3,322.000 1,993.000 TSS 2,491.000 Within the range of 7.0 to 10.0 at all times Hα (i) Reduction of Tantalum Salt to Metal Wet Air Pollution Control NSPS Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average Metric Units - mg/kg of tantalum salt reduced English Units - lbs/million lbs of tantalum salt reduced 0.572 Lead 0.266 Zinc 2.084 0.858 Ammonia (as N) 272.400 119.700 Fluoride 71.510 40.860 TSS 30.650 24.520 рH Within the range of 7.0 to 10.0 at all times

4341

(j) <u>Tantalum</u> <u>Powder</u> <u>Wash</u> NSPS

Pollutant or Pollutant Property	Maximum f Any One D	or Maximum for ay Monthly Average
Metric Units - mg/kc	of tontolum n	oudor upchod
English Units - lbs/millio		
Lead	5.	721 2.656
Zinc	20.	8408.5820001,198.000200408.700
Ammonia (as N)	2,724.	000 1,198.000
Fluoride	715.	200 408.700
TSS pH Within the	300	500 245.220 to 10.0 at all times
(k) <u>Consolidation</u> and <u>Casting</u>	<u> Contact</u> Cooli	ng NSPS
Pollutant or	Maximum f	or Maximum for
Pollutant Property	Any One D	ay Monthly Average
Metric Units - mg/kg of columk English Units - lbs/million lk		
consolidated		of current oube of
Lead		000 0.000
Zinc		000 0.000
Ammonia (as N)		000 0.000
Fluoride TSS		000 0.000
		000 0.000 to 10.0 at all times
-		
PSES are promulgated based of		
achievable by the application sedimentation, and multimedia		
filter) technology and in-pr		
methods, along with prelimi		
ammonia steam stripping for s	selected waste	streams. The
following effluent standards a	re promulgated	:
(a) <u>Concentrate</u> <u>Digestion</u> Wet	Air Pollution	Control PSES
Pollutant or	Maximum f	or Maximum for
Pollutant Property	Any One D	
Metric Units - mg/k English Units - lbs/milli		
Lead	0.	174 0.081
Zinc		635 0.261
Ammonia (as N)		910 36.450
Fluoride		770 12.440

(b) Solvent Extraction Raffinate PSES PSES

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Averag
Metric Units - mg/	kg of concentrate d	igested
English Units - 1bs/mill	Lion lbs of concentra	ate digested
Lead	2.564	1.190
Zinc	9.338	3.845
Ammonia (as N) Fluoride	1,221.000 320.400	536.500 183.100
riuoride	520.400	103.100
(c) Solvent Extraction Wet A	Air Pollution Control	L PSES
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/	kg of concentrate di	igested
English Units - lbs/mill	lion lbs of concentra	ate digested
Lead	0.069	0.032
Zinc	0.251	0.103
Ammonia (as N)	32.790	14.420
Fluoride	8.610	4.920
(d) Precipitation and Filtra	tion PSES	
Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
Foliutant Floperty	Any One Day	Monthly Average
Metric Units - ma	'kg of concentrate di	dested.
English Units - 1bs/mill		
Lead	3.833	1.780
Zinc	13.960	5.750
Ammonia (as N)	1,825.000	802.200
Fluoride	479.100	273.800
		م_{ور}
· · · · ·		

(e)Precipitation and Filtration We	et Air Pollution	<u>Control</u> PSES
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg of	concentrate di	gested
English Units - lbs/million 1	lbs of concentra	ate digested
Lead	1.778	0.826
Zinc	6.478	2.668
Ammonia (as N)	846.600	372.200
Fluoride	222.300	127.000
(f) <u>Tantalum</u> Salt Drying PSES P	SES	
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg o	f tantalum salt	dried
English Units - lbs/million	lbs of tantalu	m salt dried
Lead	16.950	7.871
Zinc	61.750	25.430
Ammonia (as N)	8,070.000	3,548.000
Fluoride	2,119.000	1,211.000
(g) Oxides Calcining Wet Air Pol	lution Control	PSES
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg of col	umbium-tantalum	n oxide dried
English Units - lbs/million lbs c	of columbium-tan	ntalum oxide dried
Lead	1.076	0.500
Zinc	3.919	1.614
Ammonia (as N)	512.200	225.200
Fluoride	134.500	76.840

(h) <u>Reduction of Tantalu</u>	um <u>Salt</u>	to <u>Meta</u>	1 PSES		· · · · · · · · · · · · · · · · · · ·
Pollutant or Pollutant Property			mum for One Day		imum for ly Average
Metric Units - English Units - 1bs/	mg/kg c /million	of tanta 1bs of	lum salt tantalu	reduced m salt r	educed
Lead Zinc Ammonia (as N) Fluoride			46.500 169.400 ,140.000 ,813.000	- 	21.590 69.750 9,732.000 3,322.000
(i) <u>Reduction of Tantalu</u> <u>Control</u> <u>PSES</u>	<u>ım Salt</u>	<u>to</u> <u>Meta</u>	<u>l Wet Ai</u>	r <u>Pollut</u>	ion
Pollutant or Pollutant Property		State 1 1 1	mum for One Day		imum for ly Average
Metric Units - English Units - 1bs/					
Lead Zinc Ammonia (as N) Fluoride			0.572 2.084 272.400 71.510		0.266 0.858 119.700 40.860
(j) <u>Tantalum</u> Powder <u>Wash</u>	n PSES				
Pollutant or Pollutant Property		Any	mum for One Day	Month	imum for ly Average
Metric Units - English Units - 1bs/m	mg/kg o nillion	f tanta. lbs of	lum powd tantalum	er washe powder	d washed
Lead Zinc Ammonia (as N) Fluoride		2	5.721 20.840 724.000 715.200		2.656 8.582 1,198.000 408.700
		· · ·			

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(k) <u>Consolidation and Casting Contact Cooling PSES</u>

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg of columbium English Units - lbs/million 1 cast or con	bs of columbium	
Lead	0.000	0.000
Zinc	0.000	0.000
Ammonia (as N)	0.000	0.000
Fluoride	0.000	0.000

PSNS are promulgated based on the effluent concentrations achievable by the application of chemical precipitation, sedimentation, and multimedia filtration (lime, settle, and filter) technology and in-process flow reduction control methods, along with preliminary treatment consisting of ammonia steam stripping for selected waste streams. The following effluent standards are promulgated for new sources:

(a) Concentrate Digestion Wet Air Pollution Control PSNS

Pollutant or	Maximum for	Maximum for	
Pollutant Property	Any One Day	Monthly Average	
Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested			
Lead	0.174	0.081	
Zinc	0.635	0.261	
Ammonia (as N)	82.910	36.450	
Fluoride	21.770	12.440	

(b) Solvent Extraction Raffinate PSNS

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested

Lead	2.564	1.190
Zinc	9.338	3.845
Ammonia (as N)	1,221.000	536.500
Fluoride	320.400	183.100

(c) <u>Solvent</u> <u>Extraction</u>	<u>Vet Air I</u>	Pollution	Control	PSNS	
Pollutant or Pollutant Property		Maximum Any One			imum for Ly Average
Metric Units – English Units – 1bs/	- mg/kg c /million	f concent lbs of con	rate di ncentra	gested te dige	ested
Lead Zinc Ammonia (as N) Fluoride		32	0.069 0.251 2.790 3.610	· · ·	$0.032 \\ 0.103 \\ 14.420 \\ 4.920$
(d) <u>Precipitation</u> and Fi	ltration	PSNS		· .	
Pollutant or Pollutant Property		Maximum Any One			mum for y Average
Metric Units - English Units - 1bs/	mg/kg o million	f concentr lbs of cor	ate die Icentra	gested te dige	sted
Lead Zinc Ammonia (as N) Fluoride		13 1,825	3.833 960 .000 .100		l.780 5.750 802.200 273.800
(e) Precipitation and Fil	tration N	Wet <u>Air</u> Po	llutior	<u>Contr</u>	ol PSNS
Pollutant or Pollutant Property	· · · ·	Maximum Any One			mum for y Average
Metric Units - English Units - 1bs/r	mg/kg of million]	concentr Lbs of con	ate dig centrat	ested e dige	sted
Lead Zinc Ammonia (as N) Fluoride		6 846	.778 .478 .600 .300	۰ ۲۰۰۰	0.826 2.668 372.200 127.000

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - II

Tantalum Salt Drying PSNS (f)

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg of English Units - lbs/million		
Lead	16.950	7.871
Zinc	61.750	25.430
Ammonia (as N)	8,070.000	3,548.000
Fluoride	2,119.000	1,211.000

Oxides Calcining Wet Air Pollution Control PSNS (g)

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg of colu English Units - lbs/million lbs of		
Lead	1.076	0.500
Zinc	3.919	1.614
Ammonia (as N)	512.200	225.200
Fluoride	134.500	76.840

Reduction of Tantalum Salt to Metal PSNS (h)

Pollutant or	Maximum	Maximum for
Pollutant Property	Any One	Monthly Average

Metric Units - mg/kg of tantalum salt reduced English Units - lbs/million lbs of tantalum salt reduced

Lead	46.500	21.590
Zinc	169.400	69.750
Ammonia (as N)	22,140.000	9,732.000
Fluoride	5,813.000	3,322.000
		_, 000

PRIMARY COLUMBIUM AND TAN	TALUM SUBCATEGORY SEC	T - II
(i) <u>Reduction of Tantalum Sa</u> <u>Control</u> PSNS	<u>lt to Metal Wet Air Poll</u>	ution
	·	
Pollutant or		aximum for
Pollutant Property	Any One Day Mon	thly Average
Metric Units - mg/k	g of tantalum salt reduc	ed
English Units - lbs/mill	ion lbs of tantalum salt	reduced
	0 530	0.000
Lead	0.572	0.266
Zinc	2.084	0.858
Ammonia (as N)	272.400	119.700
Fluoride	71.510	40.860
(j) Tantalum Powder Wash PSN	G	
(j) <u>lancatum Fowder</u> wash FSN	3	
Pollutant or		aximum for
	Mayimim ror M	
Pollutant Property Metric Units - mg/k	Any One Day Mon g of tantalum powder was	thly Average
Pollutant Property Metric Units - mg/k English Units - lbs/milli	Any One Day Mon g of tantalum powder was on lbs of tantalum powde	thly Average hed r washed
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721	thly Average hed r washed 2.656
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840	thly Average hed r washed 2.656 8.582
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N)	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000	thly Average hed r washed 2.656 8.582 1,198.000
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N)	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840	thly Average hed r washed 2.656 8.582
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200	thly Average hed r washed 2.656 8.582 1,198.000
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u>	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g Contact Cooling PSNS	thly Average hed r washed 2.656 8.582 1,198.000 408.700
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u>	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g Contact Cooling PSNS Maximum for Ma	thly Average hed r washed 2.656 8.582 1,198.000 408.700
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u> Pollutant or	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g Contact Cooling PSNS	thly Average hed r washed 2.656 8.582 1,198.000 408.700
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u> Pollutant or Pollutant Property Metric Units - mg/kg of colum	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g <u>Contact Cooling</u> PSNS Maximum for Ma Any One Day Mon	thly Average hed r washed 2.656 8.582 1,198.000 408.700 aximum for thly Average consolidate
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u> Pollutant or Pollutant Property	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g <u>Contact Cooling</u> PSNS Maximum for Ma Any One Day Mon	thly Average hed r washed 2.656 8.582 1,198.000 408.700 aximum for thly Average consolidate
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u> (k) <u>Consolidation and Castin</u> Pollutant or Pollutant or Pollutant Property Metric Units - mg/kg of colum English Units - lbs/million 1 consolidated	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g <u>Contact Cooling</u> PSNS Maximum for Ma Any One Day Mon bium or tantalum cast or bs of columbium or tanta	thly Average hed r washed 2.656 8.582 1,198.000 408.700 aximum for thly Average consolidate lum cast or
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u> Pollutant or Pollutant or Pollutant Property Metric Units - mg/kg of colum English Units - lbs/million 1 consolidated	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g <u>Contact Cooling</u> PSNS Maximum for Ma Any One Day Mon bium or tantalum cast or bs of columbium or tantal	thly Average hed r washed 2.656 8.582 1,198.000 408.700 aximum for thly Average consolidate lum cast or 0.000
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u> Pollutant or Pollutant or Pollutant Property Metric Units - mg/kg of colum English Units - lbs/million 1 consolidated Lead Zinc	Any One Day Mon g of tantalum powder was on 1bs of tantalum powde 5.721 20.840 2,724.000 715.200 g Contact Cooling PSNS Maximum for Ma Any One Day Mon bium or tantalum cast or bs of columbium or tanta 0.000 0.000	thly Average hed r washed 2.656 8.582 1,198.000 408.700 408.700 consolidate lum cast or 0.000 0.000
Pollutant Property Metric Units - mg/k English Units - lbs/milli Lead Zinc Ammonia (as N) Fluoride (k) <u>Consolidation and Castin</u> Pollutant or Pollutant or Pollutant Property Metric Units - mg/kg of colum English Units - lbs/million 1 consolidated	Any One Day Mon g of tantalum powder was on lbs of tantalum powde 5.721 20.840 2,724.000 715.200 g <u>Contact Cooling</u> PSNS Maximum for Ma Any One Day Mon bium or tantalum cast or bs of columbium or tantal	thly Average hed r washed 2.656 8.582 1,198.000 408.700 aximum for thly Average consolidate lum cast or 0.000

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PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - III

SECTION III

SUBCATEGORY PROFILE

section of the primary columbium-tantalum supplement This describes the raw materials and processes used in manufacturing primary columbium and tantalum salts and in subsequent production of the respective metals. It also presents a profile of the columbium and tantalum plants identified in this study. While chemists refer to periodic table element number 41 as niobium (Nb), in American metallurgy it is known as columbium, and this name will be used in this report.

DESCRIPTION OF PRIMARY COLUMBIUM AND TANTALUM PRODUCTION

The processes used at a columbium and tantalum production facility depend largely upon the raw material used and the plant's final product. Four basic operations from ore or slag to metal must be performed: physical and chemical breakup of the slag to form columbium and tantalum salts (digestion); ore or separation of the columbium and tantalum salts from each other and from the various impurities present; reduction of the salt to the respective metal; and fabrication of the metals into some consistent form, e.g., ingots, bars, or plates. Some plants perform the first two operations, and some the last two; some perform all four operations. A typical plant in the first category is shown in Figure III-1 (page 4359).

RAW MATERIALS

Ore concentrates and slags are the chief raw materials for the production of columbium and tantalum. Ore concentrates are +derived principally from the minerals columbite, tantalite, pyrochlore, and ferroniobium, these minerals having a relatively high concentration of the desired metals. Columbium and tantalum ores are not currently mined in the United States, rather concentrates are imported from Canada, Brazil, and Australia. Columbium-tantalum ore concentrates contain approximately 50 to 60 percent columbium pentoxide (Nb₂O₅) or tantalum pentoxide (Ta₂O₅). Slags from foreign tin production have also been found to be a reliable source. There are several sources of tin slags, but the primary source is Thailand and Malaysia. These slags generally contain eight to 14 percent Nb205 and Ta205. Columbium and tantalum are usually found together, and are somewhat difficult to separate.

DIGESTION OF ORE OR SLAG

The ore or slag is first pulverized to approximately the consistency of talcum powder. Then, columbium and tantalum (along with some impurities) are leached from the powder by either hydrofluoric acid, sulfuric acid, or by chlorine gas.

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - III

Treatment of the ore or slag powder with chlorine gas at 500 to 1,000C evolves the volatile pentachlorides of columbium and tantalum, as well as the chlorides of various other substances. These are removed by selective condensation and the columbium and tantalum chlorides are separated by distillation. This process is completely anhydrous and generates no wastewater streams. The process has been used in the past, but is not now in use on a commercial scale due to the difficulty in separating the tantalum chloride and columbium chloride by distillation.

In the leaching process, aqueous hydrofluoric acid (sulfuric acid may also be used in conjunction with hydrofluoric acid) dissolves columbium, tantalum, and impurity metals from the powder, forming the fluoride salts of these metals. Unreacted concentrate or slag (gangue) is removed by settling and decantation and disposed of as a low level radioactive waste. Leaching is a more advantageous method for digesting the ore or slag because of the ease with which the columbium and tantalum fluorides can be separated by solvent extraction. Acid mist generated in the leaching process may be controlled by wet scrubbers. The scrubber liquor produced is a source of wastewater.

SEPARATION OF SALTS

Separation and purification of the columbium and tantalum fluorides is most economically achieved using solvent extraction. Methyl isobutyl ketone (MIBK) is the most commonly used solvent. Separation of columbium and tantalum by this method hinges on the different solubilities that the fluoride salts of the two metals exhibit in MIBK as a function of hydrofluoric acid normality in the feed. For instance, tantalum shows a great affinity for the organic (MIBK) phase at low normalities, while the normality must be substantially increased for columbium to show a similar affinity. Usually, a low normality feed stream is contacted with MIBK, whereupon tantalum salt of high purity is extracted. Additional hydrofluoric acid is then added to increase the normality of the aqueous phase, (the columbium-laden stream) which is then contacted with fresh MIBK, extracting the columbium salt.

The raffinate from this step is a source of wastewater. The columbium and tantalum are next extracted from MIBK by deionized water. Following extraction, the MIBK raffinate stream is recycled. Wet air pollution control used to control solvent extraction air fumes is a source of wastewater.

Columbium and tantalum salts are precipitated from the deionized water, usually by the addition of ammonia (to precipitate columbium) and potassium fluoride (to precipitate an intermediate tantalum salt). Ammonia is also used to precipitate high purity tantalum oxides. Fluoride fumes generated during precipitation may be controlled by a wet scrubber. The crystal precipitates are filtered from the aqueous mother liquor, which is then discarded. The crystals are then washed with water and dried. Columbium oxide precipitates are calcined in a kiln using wet

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

SECT - III

scrubbers to control gaseous fumes. A pure columbium pentoxide is achieved when calcined at 850°C in an oxidizing atmosphere. Tantalum salts are also dried, but wet scrubbers are not normally used; however, the water vapor may be condensed, captured, and discharged.

REDUCTION OF SALT TO METAL

A number of methods exist for the reduction of columbium and tantalum salts to metal. They include sodium reduction, aluminothermic reduction, carbon reduction, and electrolysis.

Sodium reduction is a popular method for producing both columbium and tantalum from their salts. In this process, sodium reduces the columbium or tantalum to metal. Layers of the salt are are alternated with layers of sodium in a reaction vessel, then capped with sodium chloride to prevent oxidation of the reduced The reaction mixture is often ignited electrically, but metal. once ignited, the exothermic reaction is self-sustaining. Wet scrubbers are often used to control the gaseous emissions from the reaction vessel. After cooling, the columbium or tantalum metal-containing material is crushed, and any iron picked up from the reaction vessel is removed magnetically. The remaining metal powder is further purified by leaching with water, followed by nitric or hydrochloric acid.

The aluminothermic reaction also may be used on both columbium and tantalum salts. This method also may be used on certain ferrocolumbium ores which do not require digestion and separation of columbium and tantalum salts. The salt (or ore) is mixed with aluminum powder. Potassium chlorate is added to provide additional reaction heat, and magnesium is added to properly ignite the mixture. Columbium and tantalum are reduced to metal while aluminum is oxidized.

Carbon reduction takes place through a two-step route known as the Balke process and can be used on both columbium and tantalum salts. Its predominant use, however, is the reduction of the metal oxides. The metal oxide is first mixed with fine carbon and heated under vacuum to 1,800C, where a metal carbide and carbon monoxide are formed. The carbide is then mixed with more oxide and reacts to form the pure metal and more carbon monoxide. No known wastewater is generated during this process.

Electrolytic reduction of tantalum is sometimes practiced using fused salt techniques. Potassium fluotantalate (K_2TaF_7) , the crystal which was precipitated by potassium fluoride in the separation of salts step, is electrolyzed to yield the pure tantalum metal, which is then separated from the cathode by pulverizing both metal and cathode acid, leaching out the cathode material (usually carbon).

CONSOLIDATION AND CASTING

Both columbium and tantalum show a tendency to lose their

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - III

metallic characteristics, particularly malleability and ductility, when even small amounts of impurities are present in the metal matrix. Therefore, special techniques must be used to further purify the metals and work them into their desired form. Some of the more popular processes include electron beam melting, cold-crucible arc melting, and simultaneous compacting and resistance heating.

Electron beam melting is currently the most common method of consolidation. A beam of high voltage, low current electrons is focused onto the crude metal and the top of a retractable ingot contained in a water-cooled copper cylinder. The beam melts the crude metal, and the falling molten globules form a pool on top of the ingot. The process is continuous, with the ingot being lowered as the molten metal solidifies. Most impurities boil out of the pool into the high vacuum environment required by the electron beam and are removed.

Arc melting occurs in much the same way as electron beam melting, except that a low voltage, high current arc of electricity melts the crude metal.

Simultaneous compaction and direct resistance heating is the oldest process used and is somewhat undesirable, as the metal must be processed two or three times to reach sufficient purity. The metal is typically compacted at about 6,900 atmospheres and heated to 1,400 to 1,500°C for several hours. It is then rolled and sintered at 2,300°C. Several rolling and sintering steps may be required.

PROCESS WASTEWATER SOURCES

In summary, the major uses of water in primary columbium and tantalum processing are:

- 1. Concentrate digestion wet air pollution control,
- 2. Solvent extraction,
- 3. Solvent extraction wet air pollution control,
- 4. Precipitation and filtration of metal salt,
- 5. Precipitation and filtration wet air pollution control,
- 6. Tantalum salt drying
- 7. Oxides calcining wet air pollution control,
- 8. Reduction of tantalum salt to metal,
- 9. Reduction of tantalum salt to metal wet air pollution control,
- 10. Tantalum powder wash and wet air pollution control, and
- 11. Consolidation and casting contact cooling.

OTHER WASTEWATER SOURCES

There may be other wastewater streams associated with the production of primary columbium-tantalum. The principal wastewater stream is maintenance and cleanup water. This stream is not considered as part of this rulemaking. EPA believes that

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

SECT - III

the flows and pollutant loadings associated with this wastewater stream are insignificant relative to the wastewaters selected and it is best handled by the appropriate permit authority on a case-by-case basis under the authority of Section 402 of the Clean Water Act.

AGE, PRODUCTION, AND PROCESS PROFILE

All five of the columbium-tantalum plants identified in this study were built in the 20-year period just after World War II (Table III-1, page 4356). Average plant production is approximately 450 tons per year, as shown in Table III-2 (page 4357).

Figure III-2 (page 4360) depicts the geographic locations of the plants comprising the columbium-tantalum subcategory of the nonferrous category. The plants are scattered, with half the plants located in the New England area and the rest in the Midwest or the West.

Table III-3 (page 4358) lists the major production processes presently used in the columbium-tantalum subcategory. Also shown is the number of plants discharging from these processes.

Table III-1

INITIAL OPERATING YEAR SUMMARY OF PLANTS IN THE COLUMBIUM-TANTALUM SUBCATEGORY BY DISCHARGE TYPE

		Initial O (Plant A	perating Year ge in Years)		
Type of Plant Discharge	1983-1969 0-15	1968-1959 15-25	1958-1949 25-35	Before 1949 35+	Total
Direct	0	1	2	0	3
Indirect	0	_1	1	0	2
Total	0	2	3	0	5

PRIMARY COLUMBIUM AND TANTALUM

PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

TABLE III-2

PRODUCTION RANGES FOR THE COLUMBIUM - TANTALUM SUBCATEGORY

Production Ranges for 1976 (tons/yr)	Number of Plants
Less than 450	3
More than 450	<u>2</u>
TOTAL	5

TABLE III-3

PRODUCTION PROCESSES UTILIZED BY THE COLUMBIUM - TANTALUM SUBCATEGORY

Process		of Plants Process	Number of Plants Generating Wastewater
Digestion	,	3	· _
-Air pollution Control		3	3
Extraction		3	-
Solvent Extraction Raff Air Pollution Control	inate	3 2	3 2
Precipitation and Filtrat	ion	3	-
Precipitation and Filtr	ation	3	3
wastewater Air Pollution Control		2	2
Salt Drying		2	2
Oxide Calcining		4	_
Air Pollution Control	: .	4	4
Reduction of Tantalum Sal	t	4	-
Reduction Wastewater Air Pollution Control	•	3 2	3 2
Tantalum Powder Wash		1	. 1
Consolidation and Casting	I	4	-
Contact Cooling Water		1	1

Through reuse or evaporation practices, a plant may "generate" a wastewater from a particular process but not discharge it.

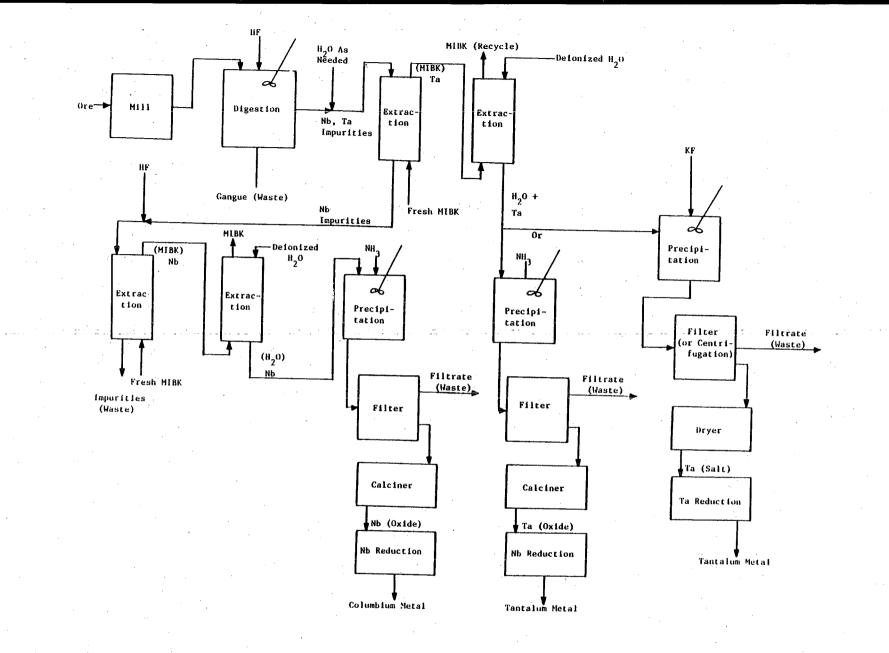


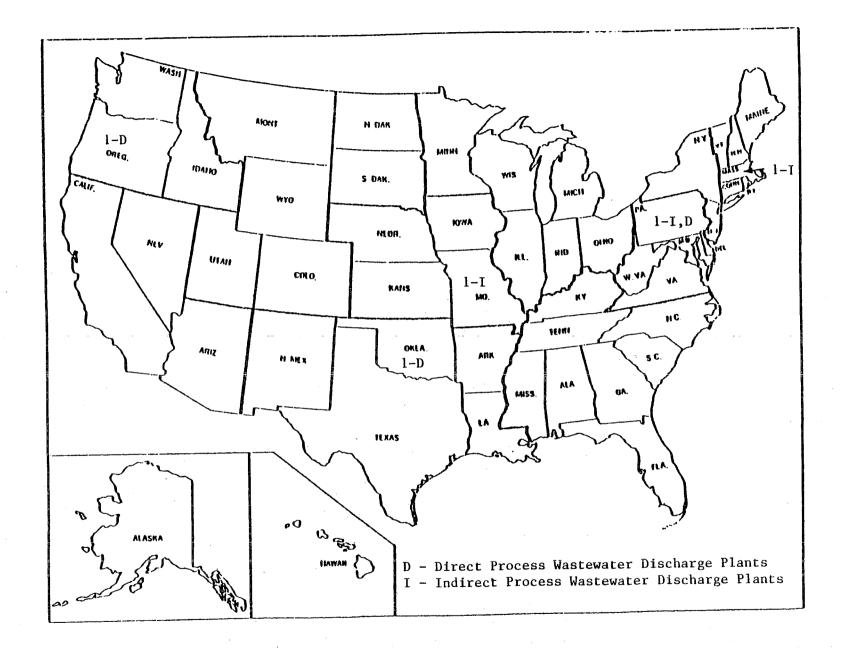
Figure III-1

PRIMARY COLUMBIUM-TANTALUM PRODUCTION PROCESS

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PRIMARY COLUMBIUM AND TANTALUM SECT -

III



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Figure III-2

GEOGRAPHIC LOCATIONS OF THE COLUMBIUM AND TANTALUM METAL PRODUCTION PLANTS

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SECTION IV

SUBCATEGORIZATION

This section summarizes the factors considered during the designation of the subdivisions of the primary columbium-tantalum subcategory.

FACTORS CONSIDERED IN SUBDIVIDING THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

The factors listed for general subcategorization were each evaluated when establishing the primary columbium-tantalum subcategory and its subdivisions. In the discussion that follows, the factors will be described as they pertain to this particular subcategory.

rationale for considering subdivision of the primarv The columbium-tantalum subcategory is based primarily on the production process used. Within this subcategory, a number of different operations are performed, which may or may not have a water use or discharge, and which may require the establishment of separate effluent limitations and standards. While primary columbium-tantalum is still considered a single subcategory, a more thorough examination of the production processes, water use and discharge practices, and pollutant generation rates has illustrated the need for limitations and standards based on a specific set of waste streams. Limitations and standards will be based on specific flow allowances for the following subdivisions:

- 1. Concentrate digestion wet air pollution control,
- 2. Solvent extraction raffinate,
- 3. Solvent extraction wet air pollution control,
- 4. Precipitation and filtration of metal salt,
- 5. Precipitation and filtration wet air pollution control,
- 6. Tantalum salt drying
- 7. Oxides calcining wet air pollution control,
- 8. Reduction of tantalum salt to metal,
- 9. Reduction of tantalum salt to metal wet air pollution control,
- 10 Tantalum powder wash and wet air pollution control, and
- 11. Consolidation and casting contact cooling.

These subdivisions follow directly from differences within the two distinct production stages of primary columbium and tantalum: production of salts from ore concentrates and slags, and the reduction of salts to produce the metals. Plants processing primary columbium and tantalum fall into three categories: plants which perform the ore-to-salt operation, plants which perform the salt-to-metal operation, plants which perform and both А review of the sampling data operations. shows that significantly different wastewater volumes and characteristics are produced by the two manufacturing processes. Therefore, 11

subdivisions of the primary columbium-tantalum subcategory are necessary.

OTHER FACTORS

The other factors considered in this evaluation were shown to be inappropriate bases for subcategorization. Air pollution control methods, treatment costs, nonwater quality aspects and total energy requirements are functions of the selected subcategorization factors--metal product, raw materials, and production processes. Therefore, they are not independent factors and will not affect the method of subcategorization. As discussed in Section IV of the General Development Document, certain other factors such as plant age, plant size, and the number of employees were also evaluated and were determined to be inappropriate for use as bases for subcategorization of this subcategory.

PRODUCTION NORMALIZING PARAMETERS

The effluent limitations and standards developed in this document establish mass limitations on the discharge of pollutants. To allow these regulations to be applied to plants with various production capacities, the mass of pollutant discharge must be related to a unit of production. This factor is known as the production normalizing parameter (PNP). In general, the amount of product or intermediate produced by a particular manufacturing process is used as the PNP. This is based on the principle that the amount of water generated is proportional to the amount of product made. The PNPs for the ll primary columbium-tantalum subdivisions are shown below:

1.	Concentrate digestion wet air pollution control	kkg of concentrate digested
2.	Solvent extraction raffinate	kkg of concentrate digested
3.	Solvent extraction wet air pollution control	kkg of concentrate digested
4.	Precipitation and filtration of metal salt	kkg of concentrate digested
5.	Precipitation and filtration wet air pollution control	kkg of concentrate digested
6.	Tantalum salt drying	kkg of tantalum salt dried
7.	Oxides calcining wet air	kkg of columbium or tantalum oxide calcined (dried)
8.	Reduction of tantalum salt to metal	kkg of tantalum salt reduced

4.363

- 9. Reduction of tantalum salt to metal wet air pollution control
- 10. Tantalum powder wash and wet air pollution control
- 11. Consolidation and casting contact cooling

kkg of tantalum salt reduced

- kkg of tantalum powder washed
- kkg of columbium or tantalum consolidated or cast

Other production normalizing parameters were considered; however, they were rejected. In the proposed mass limitations for this subcategory, the production normalizing parameter for each subdivision was the quantity of product recovered. It has been demonstrated to the Agency, however, that choosing PNPs in this manner for the first five subdivisions is not appropriate. Ore concentrates and tin slags contain significantly different columbium-tantalum values, and thus require different amounts of processing chemicals to recover equivalent amounts of metals. Production normalized flow rates presented in Section V verify this concept. A much better correlation between flow and production results when the PNP is changed to the mass of raw material rather than the mass of product.

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

SECTION V

WATER USE AND WASTEWATER CHARACTERISTICS

This section describes the characteristics of wastewater associated with the primary columbium-tantalum subcategory. Data used to quantify wastewater flow and pollutant concentrations are presented, summarized, and discussed. The contribution of specific production processes to the overall wastewater discharge from primary columbium-tantalum plants is identified whenever possible.

The two principal data sources used in preparation of this document are data collection portfolios (dcp) and field sampling collection portfolios contain information results. Data wastewater flows. A significant amount of data regarding has been collected from comments on the proposed also mass limitations and specific data requests to verify comments.

In order to quantify the pollutant discharge from primary a field sampling columbium-tantalum plants, program was Wastewater samples were collected in two phases: conducted. screening and verification. The first phase, screen sampling, was to identify which toxic pollutants were present in the wastewaters from production of the various metals. Screening samples were analyzed for 125 of the 126 toxic pollutants and other pollutants deemed appropriate. Because the analytical standard for TCDD was judged to be too hazardous to be made generally available, samples were never analyzed for this There is no reason to expect that TCDD would be pollutant. present in columbium-tantalum wastewater. A total of 10 plants selected for screen sampling in the nonferrous were metals manufacturing category with one of those plants in the primary columbium-tantalum subcategory. In general, the samples were for three classes of pollutants: toxic analyzed organic pollutants, toxic metal pollutants, and criteria pollutants both conventional and includes nonconventional (which pollutants).

described in Section IV of this supplement, the primary Ag columbium-tantalum subcategory has been further subdivided into subdivisions or building blocks, each representing a major 11 source of wastewater in the subcategory. Differences in the wastewater characteristics associated with these subdivisions are expected. For this reason, wastewater to be streams corresponding to each subdivision are addressed separately in the discussions that follow.

WASTEWATER SOURCES, DISCHARGE RATES, AND CHARACTERISTICS

The wastewater data presented in this section were evaluated in light of production process information compiled during this study. As a result, it was possible to identify the principal wastewater sources in the primary columbium-tantalum subcategory. They are:

- 1. Concentrate digestion wet air pollution control,
- 2. Solvent extraction raffinate,
- 3. Solvent extraction wet air pollution control,
- 4. Precipitation and filtration of metal salt,
- 5. Precipitation and filtration wet air pollution control,
- 6. Tantalum salt drying,
- 7. Oxides calcining wet air pollution control,
- 8. Reduction of tantalum salt to metal,
- 9. Reduction of tantalum salt to metal wet air pollution control,
- 10. Tantalum powder wash and wet air pollution control, and
- 11. Consolidation and casting contact cooling.

Data supplied by dcp responses were evaluated, and two flow-toproduction ratios were calculated for each stream. The two ratios, use and wastewater water discharge flow, are differentiated by the flow value used in calculation. Water use is defined as the volume of water or other fluid (e.g., solvent extraction raffinate) required for a given process per mass of product and is therefore based on the sum of recycle and make-up flows to a given process. Wastewater flow discharged after pretreatment or recycle (if these are present) is used in calculating the production normalized flow -- the volume of wastewater discharged from a given process to further treatment, disposal, or discharge per mass of columbium or tantalum produced. Differences between the water use and wastewater flows associated with a given stream result from recycle, evaporation, and carry-over on the product. The production values used in calculation correspond to the production normalizing parameter, PNP, assigned to each stream, as outlined in Section IV. The production normalized flows were compiled and statistically analyzed by stream type. Where appropriate, an attempt was made to identify factors that could account for variations in water This information is summarized in this section. A similar use. analysis of factors affecting the wastewater values is presented in Sections X, XI, and XII where representative BAT, BDT, and pretreatment discharge flows are selected for use in calculating the effluent limitations and standards. As an example, reduction of tantalum salt to metal air scrubbing flow is related to the reduction production. As such, the discharge rate is expressed in liters of scrubber wastewater per metric ton of metal reduced.

In order to quantify the concentrations of pollutants present in wastewater from primary columbium-tantalum plants, wastewater samples were collected at four of the five primary columbium-tantalum plants. Diagrams indicating the sampling sites and contributing production processes are shown in Figures V-1 through V-4 (pages 4420 - 4423).

The raw wastewater sampling data for the primary columbiumtantalum subcategory are presented in Tables V-2, V-4, V-6, V-8, and V-12, (pages 4373, 4379, 4384, 4388, and 4394 respectively). PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

Treated wastewater sampling data are shown in Tables V-18 through V-20 (pages 4411, 4414 and 4417). Tables V-15 through V-17 (pages 4402, 4404, and 4407) show miscellaneous raw wastewater for plants A, B, and C. These data were not used for the wastewater is characterizations discussed below because the wastewater is combined with nonprocess water or is generated by nonscope processes. Where no data are listed for a specific day of sampling, the wastewater samples for the stream were omitted. If the analysis did not detect a pollutant in a waste stream, the pollutant was omitted from the table.

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The data tables include some wastewater samples measured at concentrations not considered quantifiable. The base neutral extractables, acid fraction extractables, and volatile organics are generally considered not quantifiable at concentrations at or Below this concentration, the data 0.010 mg/l. is below considered too susceptible to random error to be quantitatively accurate. However, these data are useful in that they indicate the presence of a particular pollutant. The pesticide fraction is considered nonquantifiable at concentrations equal to or less than 0.005 mg/l. Nonquantifiable results are designated in the tables with an asterisk (double asterisk for less than or equal to 0.005 mg/l).

These detection limits shown on the data tables are not the same in all cases as the published detection limits for these pollutants by the same analytical methods. The detection limits used were reported with the analytical data and hence are the appropriate limits to apply to the data. Detection limit variation can occur as a result of a number of laboratoryspecific, equipment-specific, and daily operator-specific factors. These factors can include day-to-day differences in machine calibration, variation in stock solutions, and variation in operators.

The statistical analysis of data includes some samples measured at concentrations considered not quantifiable. Data reported as asterisk are considered as detected but below quantifiable an concentrations, and a value of zero is used for averaging. Toxic nonconventional, and conventional pollutant data organic, reported with a "less than" sign are considered as detected, but further quantifiable. A value of zero is also used for not If a pollutant is reported as not detected, it is averaging. excluded in calculating the average. Finally, toxic metal values reported as less than a certain value were considered as not detected, and a zero was used in the calculation of the average. For example, three samples reported as ND, *, and 0.021 mg/l have an average value of 0.010 mg/l.

The method by which each sample was collected is indicated by number, as follows.

1	one-time grab	
2	24-hour manual composite	
3	24-hour automatic composit	e

4	48-hour	manual composite
C	10 hour	automotic commonit

- 5 48-hour automatic composite
- 6 72-hour manual composite
- 7 72-hour automatic composite

In the data collection portfolios, all of the columbium-tantalum plants indicated that the toxic organic pollutants were known or believed to be absent from their wastewater. The majority of the metals were believed to be absent as summarized below:

	Known Present	Believed Present	Believed Absent	Known Absent
Antimony	1	-	2	1
Arsenic	1	·	2	1
Beryllium	-	·	3	1
Cadmium	 .	-	2	2
Chromium	1	· 1	1	1
Copper	_	1	2	1
Lead	-	· _	2	2
Mercury	-	1	1	2
Nickel	1	1	1	1
Selenium		· <u> </u>	4	1
Silver	-	-	2	2
Zinc	-		³ 3	1
			1	

CONCENTRATE DIGESTION WET AIR POLLUTION CONTROL

The first step in the production of primary columbium and tantalum is the digestion of ore concentrates and slags with hydrofluoric acid. The process solubilizes columbium and tantalum, along with various other metals which require removal. Three of the five columbium-tantalum plants use wet scrubbers on their concentrate digestion process. Water use and discharge rates are shown in liters per metric ton of columbium-tantalum concentrate digested in Table V-1 (page 4372).

Table V-2 (page 4373) summarizes the raw wastewater sampling data for the toxic and selected conventional and nonconventional pollutants. The wet scrubber liquor is strongly acidic (pH of approximately 2.0), containing suspended solids, fluorides, and at treatable concentrations (see Table metals V-2). some Insoluble ganque impurities are removed by filtration. On-site disposal of gangue impurities is required because it is radioactive. The waste gangue slurry is typically contained in a holding pond, the overflow from which is acidic and contains quantifiable concentrations of metals, fluorides, and suspended solids.

SOLVENT EXTRACTION RAFFINATE

The digested solution containing columbium and tantalum is contacted with an organic solvent such as methyl isobutyl ketone (MIBK) in a two step multistage extraction process, resulting in

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the extraction and separation of columbium and tantalum. Three plants discharge this wastewater as shown in Table V-3 (page 4378). The impurities from digestion remaining in the raffinate typically include treatable concentrations of organics, fluorides, metals, suspended solids, and oil and grease. The sampling data from an extraction raffinate waste stream are presented in Table V-4 (page 4379).

SOLVENT EXTRACTION WET AIR POLLUTION CONTROL

After extraction, the organic streams bearing columbium and tantalum are often contacted with deionized water to strip the columbium and tantalum from the organic phase. The organic solvent is then recycled to the first extraction process. Two plants use wet scrubbers to control air emissions from extraction operations. One of these plants uses the same scrubber for air pollution control of concentrate digestion and solvent extraction. The water use and discharge rates for the two plants are presented in Table V-5 (page 4383) in liters per metric ton of columbium-tantalum concentrate digested. This wastewater is acidic and contains concentrations of toxic organics and metals, fluorides, and suspended solids as shown in Table V-6 (page 4384).

PRECIPITATION AND FILTRATION OF METAL SALT

Precipitation of pure metal salts from the aqueous phase may be accomplished by ammonia addition to form columbium and tantalum All three plants reporting this waste stream discharge oxides. as shown in Table V-7 (page 4387). The filtrate wastewater it typically contains treatable concentrations of ammonia, fluoride, metals, and suspended solids. Ammonia stripping is frequently practiced to recover ammonia from the filtrate prior to discharge of the waste stream. Tantalum may also be recovered by treatment the solubilized tantalum salt with hydrofluoric acid of and potassium fluoride to precipitate potassium fluotantalate (K₂TaF₇). This precipitate also requires filtration and washing, leaving a filtrate effluent stream containing measurable concentrations of potassium, fluorides, and chlorides (see Table V-8) (page 4388).

PRECIPITATION AND FILTRATION WET AIR POLLUTION CONTROL

Gaseous fumes emitted during precipitation of columbium-tantalum are controlled with wet scrubbers at two plants. The scrubber liquor reflects the precipitation supernatant and contains fluoride and ammonia. One of the plants pretreats the scrubber liquor with ammonia steam stripping prior to central treatment. The water use and discharge rates for this scrubbing operation are presented in Table V-9 (page 4391).

TANTALUM SALT DRYING

Following filtration, potassium fluotantalate precipitates are usually dried to yield purified salts. Two of the five

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columbium-tantalum plants capture steam in drying operations. 4392). Water discharge rates are shown in Table V-10 (page reflects Wastewater associated with this process the precipitation process used. For example, treatable concentrations of fluoride may be present when potassium fluoride is used as the reagent for precipitation. Table V-12 (page 4394) shows data from combined wastewater from the tantalum salt drying, reduction of salt to metal, and reduction of salt to metal scrubber waste streams.

OXIDES CALCINING WET AIR POLLUTION CONTROL

Columbium-tantalum oxides are precipitated from solution using ammonia and then dried and calcined in a kiln. Four plants reported using wet scrubbers to control emissions from this process as shown in Table V-11 (page 4393). Scrubber liquor blowdown will contain a large quantity of ammonia and is steam stripped at two plants.

REDUCTION OF TANTALUM SALT TO METAL WASTEWATER

Reduction processes vary somewhat in the columbium-tantalum subcategory. Of the several reduction techniques discussed in Section III, only two were reported in practice by plants in the The first of these, sodium columbium-tantalum subcategory. reduction, appears to be the dominant technique. The process requires extensive washing of the product metal with water or a combination of water and acid. The production normalized discharge rates are shown in Table V-13 (page 4400). This waste stream typically contains treatable concentrations of fluoride (see Table V-12, page 4394), as well as toxic metals, chloride, and oil and grease. The other reduction process used, aluminothermic reduction, is reported to generate no wastewater. The waste streams are some times passed through a cyclone to recover valuable columbium and tantalum solids. In addition, water is used for sizing at one of the plants surveyed. However, this wastewater stream is combined with washing operations and is not further considered as a separate wastewater stream.

REDUCTION OF TANTALUM SALT TO METAL WET AIR POLLUTION CONTROL

Reduction process emissions are frequently controlled with wet scrubbers. The resulting discharge is similar to the reduction washing streams. This waste stream may also be passed through a cyclone to recover columbium and tantalum solids, if present. Water use and discharge rates are presented in Table V-14 (page 4401) in liters per metric ton of tantalum metal reduced. Sampling data for this waste stream are contained in Table V-12 (page 4394). This waste water contains many of the same pollutants found in reduction of salt to metal wastewater.

TANTALUM POWDER WASH AND SCRUBBER

Following reduction, tantalum powder may be acid washed to impart

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certain surface characteristics. This waste stream should contain very little toxic metals or fluoride. However, it will be characterized by a low pH. This waste stream should not be confused with the reduction of salt to metal wastewater. Although an acid medium may be used and achieve similar surface characteristics, reduction of salt to metal wastewater is generated primarily to leach away potassium fluoride, sodium fluoride, and sodium chloride salts. Therefore, tantalum powder wash is only applicable to those plants that carry out a separate wash step from reduction of salt to metal. The only plant with this stream reported a once through water usage of 20,433 l/kkg of tantalum powder washed.

CONSOLIDATION AND CASTING CONTACT COOLING

Only one of the plants surveyed practiced direct contact cooling of metal castings. This plant recycles 100 percent of the water used for this operation, resulting in zero discharge. No sampling data were available for this waste stream.

TABLE V-1

WATER USE AND DISCHARGE RATES FOR CONCENTRATE DIGESTION WET AIR POLLUTION CONTROL

(l/kkg of columbium-tantalum concentrate digested)

Plant Code	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge Rate
507	0	6219	6129
509	0	100191	100191
519*	80	93800	13132

* Same scrubber used for solvent extraction wet air pollution control.

Table V-2

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA CONCENTRATE DIGESTION SCRUBBER RAW WASTEWATER

Concentrations (mg/1, Exce					cept as No	ept as Noted)		
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
Toxi	c Pollutants(a)	•						
4.	benzene	22 114	2 2	*	0.042 *	* *	ND ND	0.021
6.	carbon tetrachloride	22 114	2 2	ND	ND 0.074	0.017	ND ND	0.017 0.037
7.	chlorobenzene	22	2	-	ND	ND	*	*
10.	1,2-dichloroethane	22 114	2 2	ND	0.156 0.04	0.086 ND	0.062 *	0.101 0.02
14.	1,1,2-trichloroethane	22 114	2 2	*	ND <0.032	ND ND	ND ND	<0.032
23.	chloroform	22 114	2 2	0.422	0.156 0.046	0.135 0.034	0.017 ND	0.103 0.040
30.	1,2- <u>trans</u> -dichloroethylene	22 114	2 2	ND	ND ND	ND 0.484	ND ND	0.484
38.	ethylbenzene	22 114	2 2	ND	0.057 ND	* ND	* ND	0.019
44.	methylene chloride	22 114	2 2	ND	88.4 ND	ND ND	ND ND	88.4

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA CONCENTRATE DIGESTION SCRUBBER RAW WASTEWATER

				Conce	ntrations	(mg/1, Ex	cept as N	oteď)
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
48.	dichlorobromomethane	22 114	2 2	0.025	ND 0.038	ND ND	ND ND	0.038
51.	chlorodibromomethane	22 114	2	*	ND <0.089	ND *	ND ND	· · · · ·
66.	bis(2-ethylhexyl) phthalate	22	7		0.48			0.48
68.	di-n-butyl phthalate	22	7		80.0	×		0.08
85.	tetrachloroethylene	22 114	2 2	*	0.157 ND	ND ND	ND ND	0.157
87.	trichloroethylene	22 114	2 2	ND	0.235 ND	* ND	ND ND	0.118
106.	PCB-1242	22	7		<0.015		` .	<0.015
109.	PCB-1232	22	7		<0.015			<0.015
113.	toxaphene	22	7		**			**
114.	antimony	22	7		2.9	ан А		2.9
115.	arsenic	22	7		0.003			0.003

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA CONCENTRATE DIGESTION SCRUBBER RAW WASTEWATER

a de la composition Na seguidad de la composition		Stream	Sample	Concentrations (mg/1, Exc				ept as Noted)		
	Pollutant	_Code_	Typet	Source	Day 1	Day 2	Day 3	Average		
117.	beryllium	22	7		0.18			0.18		
118.	cadmium	22	7		40	н С. С. С		40		
119.	chromium	22	7		1			1		
120.	copper	22	7		300		•	300		
121.	cyanide	22	7		0.003	0.004	0.002	0.003		
122.	lead	22	7		900	· · · ·		900		
123.	mercury	22	7	A. 100	0.063			0.063		
124.	nickel	22	7		5			5		
125.	selenium	22	7		<0.002	an an Arras Ar		<0.002		
127.	thallium	22	7	• • • •	<0.05			<0.05		
128.	zinc	22	7		1,000	n an	. 1	,000		

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA CONCENTRATE DIGESTION SCRUBBER RAW WASTEWATER

			Concentrations (mg/l, Except as Noted)					
Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
Nonconventionals			•	·				
ammonia	22	6			87.7	18.5	53.1	
chemical oxygen demand (COD)	22	7		2,030			2,030	
fluoride	22	7		24,000	2,800		13,400	
phenols (total; by 4-AAP method)	22 114	1 2		0.0			0.018 1 0.015	
total organic carbon (TOC)	22	7		236			236	
Conventionals					· · ·			
oil and grease	22 114	1		9 5	8 8	10 4	9 5.7	
total suspended solids (TSS)	22	6		823			823	
pH (standard units)	22 114	1 1		3. 2.4		2.0		

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA CONCENTRATE DIGESTION SCRUBBER RAW WASTEWATER

			Conce	entrations	(mg/1, Ex	cept as N	loted)
Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average

Samples taken from stream 114 were not analyzed for toxic metals. (a)

(b) Reported together.

tSample type: Note: These numbers also apply to subsequent sampling data tables in this section.

- 4377
- 1 one-time grab
- 2 24-hour manual composite 3 24-hour automatic composite
- 4 48-hour manual composite 5 48-hour automatic composite
- 6 72-hour manual composite
- 7 72-hour automatic composite

*Indicates less than or equal to 0.01 mg/l. **Indicates less than or equal to 0.005 mg/1. SECT

TABLE V-3

WATER USE AND DISCHARGE RATES FOR SOLVENT EXTRACTION RAFFINATE

(l/kkg of columbium-tantalum concentrate digested)

Percent Recycle	Production Normalized Water <u>Use</u>	Production Normalized Discharge Rate
0	10971	10971
0	7338	7338
0	9463	9463
	Recycle 0	Percent RecycleNormalized Water01097107338

Table V-4

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA SOLVENT EXTRACTION RAFFINATE RAW WASTEWATER

				Concentrations (mg/1, Except as Noted)					
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
<u>Toxic</u>	Pollutants(a)								
1.	acenaphthene	25	1		ND	0.017	ND	0.017	
4.	benzene	25 114	2 2		<0.046 *	<0.049 *	<0.051 ND	<0.049 *	
6.	carbon tetrachloride	25 114	2 2	ND	ND 0.074	ND *	ND ND	0.037	
7.	chlorobenzene	25 114	2 2	ND	ND ND	1.001 ND	0.034 ND	0.522	
8.	1,2,4-trichlorobenzene	25	1	- -	ND	0.051	*	0.026	
10.	1,2-dichloroethane	25 114	2 2	ND	ND 0.04	0.135 ND	ND *	0.02	
14.	1,1,2-trichloroethane	25 114	2 2		ND <0.032	ND ND	ND ND	<0.032	
15.	1,1,2,2-tetrachloroethane (b) 25 114	2 2		ND ND	ND ND	<20.39 ND	<20.39	
23.	chloroform	25 114	2 2	0.422	0.203 0.046	0.24 0.034	0.181 ND	0.208 0.040	

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA SOLVENT EXTRACTION RAFFINATE RAW WASTEWATER

	Concentrations (mg/1, Except as Noted)								
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
30.	1,2- <u>trans</u> -dichloroethylene	25 114	2 2	ND	ND ND	ND 0.484	ND ND	0.484	
38.	ethylbenzene	25 114	22	ND	ND ND	ND ND	0.04 ND	0.04	
48.	dichlorobromomethane	25 114	2 2	0.025	ND 0.038	ND ND	ND ND	· · ·	
51.	chlorodibromomethane	25 114	2 2	*	ND <0.089	ND *	7.08 ND	7.08	
54.	isophorone	25	- 1		ND	0.029	ND	0.029	
56.	nitrobenzene	25	1.		ND	0.1	ND	0.1	
66.	bis(2-ethylhexyl) phthalate	25	1	•	0.036	0.02	*	0.019	
68.	·	25	1		0.012	0.042	*	0.018	
71.	dimethyl phthalate	25	1		ND	ND	0.012	0.012	
85.	tetrachloroethylene	25 114	2 2	*	<0.245 ND	ND ND	0.138 ND	0.138	
87.	trichloroethylene	25 114	2 2	*	ND ND	<0.259 *	<0.27 ND	<0.265 *	

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA SOLVENT EXTRACTION RAFFINATE RAW WASTEWATER

				Concentrations (mg/1, Except as Noted)					
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
107.	PCB-1254	25	1		**	**	**	**	
109.	PCB-1248	25	1	·	**	**	**	**	
113.	toxaphene	25	1		<0.02	<0.02	<0.02	<0.02	
114.	antimony	25	1		20	4	30	18	
115.	arsenic	25	1 1		45	10	25	26.7	
117.	beryllium	25	1		<0.2	<0.2	<0.2	<0.2	
118.	cadmium	25	1		<0.2	<0.2	<0.2	<0.2	
119.	chromium	25	1 40	1	,000	1,000	1,000	1,000	
120.	copper	25	1		50	10	70	43.3	
121.	cyanide	25	1	· .	0.016	0.033	0.011	0.020	
122.	lead	25	1		200	300	1,000	500	
123.	mercury	25	1		0.01	0.007	-	0.011	
124.	nickel	25	1		<0.5	<0.5	<0.5	<0.5	
125.	selenium	25	1		70	35	30	45	

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA SOLVENT EXTRACTION RAFFINATE RAW WASTEWATER

			Conc	entration	s (mg/1, Ex	cept as N	oted)
Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
127. thallium	25	1		1.14	1.18	0.83	1.05
128. zinc	25	1		500	400	400	433
Nonconventionals		• • ••		_ . _ .			
chemical oxygen demand (COD)	25	1	. 1	3,000	11,000 10),000 1	1,330
phenols (total; by 4-AAP method)	25 114	1 2		0.02 0.01	0.008 0.024	0.014 0.011	0.014 0.015
Conventionals							
oil and grease	25	1		39	20	, 7	22
total suspended solids (TSS)	25	1	· .	12,600	18,300 1	7,500	6,130
pH (standard units)	114	1		2.4	1.9	2.0	

(a) Samples from both streams were not analyzed for the acid fraction of toxic organic pollutants.

(b) Questionable analysis.

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PRIMARY COLUMBIUM AND TANTALUM SECT I. <

TABLE V-5

WATER USE AND DISCHARGE RATES FOR SOLVENT EXTRACTION WET AIR POLLUTION CONTROL

(1/kkg of columbium-tantalum concentrate digested)

<u>Plant</u> Code	Percent Recycle	Production Normalized <u>Water</u> Use	Production Normalized Discharge Rate
507	0	2456	2456
519*	80	93800	13132

* Same scrubber used in concentrate digestion

Table V-6

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA SOLVENT EXTRACTION SCRUBBER RAW WASTEWATER

				Conce	ntrations	(mg/l, Ex	cept as No	oted)
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
Toxic	Pollutants							
4.	benzene	22	2		0.042	*	ND	0.021
6.	carbon tetrachloride	22	2	-	ND	0.017	. ND	0.017
7.	chlorobenzene	22	2		ND	ND	*	*
10.	1,2-dichloroethane	22	2		0.156	0.086	0.062	0.101
23.	chloroform	22	2		0.156	0.135	0.017	0.103
38.	ethylbenzene	22	2		0.057	*	*	0.019
44.	methylene chloride	22	2		88.4	ND	ND	88.4
66.	bis(2-ethylhexyl) phthalate	22	7		0.48			0.48
85.	tetrachloroethylene	22	2	·	0.157	ND	ND	0.157
87.	trichloroethylene	22	2		0.235	*	ND	0.118
106.	PCB-1242	22	7	· .	<0.015	. •		<0.015
109.	PCB-1232	22	7		<0.015			<0.015

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA SOLVENT EXTRACTION SCRUBBER RAW WASTEWATER

					Concentrations (mg/l, Except as Noted)				
• • •	Pollutant		Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
113.	toxaphene		22	7		**		,	**
114.	antimony		22	7		2.9			2.9
115.	arsenic		22	·	nin in the second	0.003		n in star	0.003
117.	beryllium	· · · ·	22	7		0.18			0.18
118.	cadmium		22	7		40	Ч., с.		40
119.	chromium		22	· 7		1. 1. 1. 1. 1.		с. У. 	·1
120.	copper		22	7		300			300
121.	cyanide	· · ·	22	7	ана. Алагана	0.003	0.004	0.002	0.003
122.	lead		22	7		900	· · ·		900
123.	mercury	·	22	7	· · ·	0:063			0.063
124.	nickel		22	7		5			5
125.	selenium		22	7		<0.002			<0.002
127.	thallium		22	7	. •	<0.05	- -		<0.05
128.	zinc		22	7	1	,000		- 1	,000

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PRIMARY COLUMBIUM AND TANTALUM SECT -

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA SOLVENT EXTRACTION SCRUBBER RAW WASTEWATER

	A . ¹	0 1	Concentrations (mg/l, Except as Noted)					
Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
Nonconventionals								
ammonia	22	6			87.7	18.5	53.1	
chemical oxygen demand (COD)	22	7		2,030	- · . -	·- · ·	2,030	
fluoride	22	7		24,000	28,000		13,400	
phenols (total; by 4-AAP method)	22	1		0.0	0.028	0.01	0.018	
total organic carbon (TOC)	22	7		236			236	
Conventionals								
oil and grease	22	1		9	8	10	9	
total suspended solid (TSS)	22	6		823	· · ·		823	
pH (standard units)	22	1		3.1	i ·			

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PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

TABLE V-7

SECT - V

WATER USE AND DISCHARGE RATES FOR PRECIPITATION AND FILTRATION OF COLUMBIUM-TANTALUM SALT

(1/kkg of columbium-tantalum concentrate digested)

Plant Code	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge Rate
507	0	18419	18419
509	0	9735	9735
519	0	12912	12912

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA PRECIPITATION AND FILTRATION RAW WASTEWATER

		Concentrations (mg/1, Except as Noted)						
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
Toxic	Pollutants(a)							
4.	benzene	117	2	*	*	*		*
15.	1,1,2,2-tetrachloroethane	117	2	*	*	ND		ND
66.	bis(2-ethylhexyl) phthalate	117	2	0.049	1.158		-	1.158
85.	tetrachloroethylene	117	2	*	*	ND		*
87.	trichloroethylene	117	2	*	*	<0.023		*
108.	PCB-1221	117	2	**	**			**
112.	PCB-1016	117	2	**	**			**
113.	toxaphene	117	2	ND	**			**
114.	antimony	117	2	<0.1	<0.1			<0.1
115.	arsenic	117	2	<0.01	0.04			0.04
117.	beryllium	117	2	<0.001	<0.1			<0.1
	-	117	2	<0.002	2			2
118. 119.	cadmium chromium	117	2	<0.005	<0.5			<0.5

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA PRECIPITATION AND FILTRATION RAW WASTEWATER

S two on	Com 1 c	Conce	oted)			
_Code	Sample Typet	Source Day 1		Day 2	Day 3	Average
117	2	<0.006	0.8			0.8
117	1	•	0.005	0.009		0.007
117	2	<0.02	<2		. •	:<2
117	· · · · · 2- · ·	0.0001	<0.0001	•••••••		<0.0001
117	2	<0.005	<0.5			<0.5
117	2	<0.01	<0.01	н		<0.01
117	2	<0.02	0.07	· .		0.07
117	2	<0.1	<0.1			<0.1
117	2	<0.06	<6			<6
		· · ·	• •	•	i F	• • •
117	2	<0.1	1,450	34	4.2	496.1
117	2		22	. · ·		22
117	2	1.6	3,525			3,525
117	2	, , , , , , , , , , , , , , , , , , , ,	0.008	0.009	0.00	0.007
117	2	, ,	1			· 1
	117 117 117 117 117 117 117 117 117 117	CodeTypet117211711172	StreamSample TypetSource1172 $\langle 0.006$ 11711172 $\langle 0.02$ 1172 $\langle 0.005$ 1172 $\langle 0.005$ 1172 $\langle 0.01$ 1172 $\langle 0.02$ 1172 $\langle 0.02$ 1172 $\langle 0.11$ 1172 $\langle 0.11$ 1172 $\langle 0.11$ 1172 $\langle 1.6$ 11721.611721.6	StreamSample TypetSourceDay 11172 $\langle 0.006$ 0.811710.0051172 $\langle 0.02$ $\langle 2$ 11720.0001 $\langle 0.0001$ 1172 $\langle 0.005$ $\langle 0.5$ 1172 $\langle 0.02$ $\langle 0.01$ 1172 $\langle 0.02$ 0.07 1172 $\langle 0.1$ $\langle 0.1$ 1172 $\langle 0.1$ $\langle 0.1$ 1172 $\langle 0.1$ $\langle 0.1$ 1172 $\langle 0.1$ $\langle 22$ 1172 $\langle 0.1$ $1,450$ 1172 $\langle 0.1$ $3,525$ 1172 0.008	StreamSample TypetSourceDay 1Day 21172 $\langle 0.006$ 0.8 1171 0.005 0.009 1172 $\langle 0.02$ $\langle 2$ 1172 0.0001 $\langle 0.0001$ 1172 $\langle 0.005$ $\langle 0.5$ 1172 $\langle 0.005$ $\langle 0.01$ 1172 $\langle 0.02$ 0.07 1172 $\langle 0.11$ $\langle 0.11$ 1172 $\langle 0.11$ $\langle 0.11$ 1172 $\langle 0.11$ $\langle 0.11$ 1172 $\langle 0.16$ $\langle 6$ 1172 $\langle 0.13$ $1,450$ 34 1172 1.6 $3,525$ 1172 0.008 0.009	CodeTypetSourceDay 1Day 2Day 31172 $\langle 0.006$ 0.8 1171 0.005 0.009 1172 $\langle 0.02$ $\langle 2$ 1172 0.0001 $\langle 0.0001$ 1172 $\langle 0.005$ $\langle 0.5$ 1172 $\langle 0.005$ $\langle 0.5$ 1172 $\langle 0.02$ 0.07 1172 $\langle 0.11$ $\langle 0.11$ 1172 $\langle 0.06$ $\langle 6$ 1172 $\langle 0.11$ $1,450$ 34 1172 22 1172 1.6 $3,525$ 1172 0.008 0.009 0.006

PRIMARY COLUMBIUM AND TANTALUM SECT

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA PRECIPITATION AND FILTRATION RAW WASTEWATER

		_	Conce	(mg/1, Exc	Except as Noted)		
Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
Conventionals	,			•			
oil and grease	117	1		12	6		9
total suspended solids (TSS)	117 .	2		27,890	•		27,,890
pH (standard units)	117	. 1		10.5	9.5	10.2	2

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(a) Samples from stream 117 were not analyzed for the toxic organic acid fraction.

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

TABLE V-9

- 1a ...

WATER USE AND DISCHARGE RATES FOR PRECIPITATION AND FILTRATION WET AIR POLLUTION CONTROL

(1/kkg of columbium-tantalum concentrate digested)

<u>Plant</u> Code	Percent <u>Recycle</u>	Production Normalized Water Use	Production Normalized Discharge Rate
509	0	114116	114116
519	0	12910	12910

TABLE V-10

WATER USE AND DISCHARGE RATES FOR TANTALUM SALT DRYING

(l/kkg of tantalum salt dried)

Plant Code	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge Rate
509	0	17493	17493
509*	0	60544	60544

* Water use represents tantalum salt drying and oxides calcining.

TABLE V-11

WATER USE AND DISCHARGE RATES FOR OXIDES CALCINING WET AIR POLLUTION CONTROL

(1/kkg of columbium-tantalum oxide calcined)

<u>Plant</u> Code	Percent <u>Recycle</u>	Production Normalized Water Use	Production Normalized Discharge Rate
507	89	30366	30366
509	0	50966	50966
519	100	NR	NR
4225	0	33935	33935

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA METAL SALT DRYING SCRUBBER, REDUCTION OF SALT TO METAL, AND REDUCTION OF SALT TO METAL SCRUBBER RAW WASTEWATER

				Concentrations (mg/1, Except as Noted)					
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
<u>Toxic Po</u>	llutants(a,b)								
4. be	nzene	23 113	2 2	*	ND *	ND *	ND *	*	
10. 1,	2-dichloroethane	23 113	2 2	ND	0.039	0.016 0.026	ND 0.021	0.022 0.023	
12. he	exachloroethane	23 113	7 7	ND	ND 0.023			0.023	
14. 1,	1,2-trichloroethane	23 113	2 2	*	ND <0.011	ND ND	ND ND	<0.011	
15. 1,	1,2,2-tetrachloroethane	23 113	2 2	ND	<0.029 ND	ND ND	ND ND	<0.029	
23. ch	nloroform	23 113	2 2	0.422	ND ND	0.018 0.032	0.012	0.015 0.061	
30. 1,	,2- <u>trans</u> -dichloroethylene	23 113	2 2	ND	0.015 ND	ND 0.26	nd Nd	0.015 0.26	
35. 2,	4-dinitrotoluene	23 113	7 7	ND	ND <0.016	• *		<0.016	
36. 2,	,6-dinitrotoluene	23 113	7 7	ND	ND <0.016			<0.016	

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA

METAL SALT DRYING SCRUBBER, REDUCTION OF SALT TO METAL, AND REDUCTION OF SALT TO METAL SCRUBBER RAW WASTEWATER

	ė.		Conc	centrations	cept as N	ept as Noted)	
Pollutant	Stream <u>Code</u>	Sample Typet	Source	Day 1	Day 2	Day 3	Average
38. ethylbenzene	23 113	2 2	ND	ND ND	* ND	ND ND	*
44. methylene chloride	23 113	2 2	ND	ND ND	ND ND	ND ND	
47. bromoform	23 113	2 2	ND	ND *	0.134 ND	ND 0.021	0.134 0.011
51. chlorodibromomethane	23 113	2 2	*	ND <0.031	0.02 ND	ND ND	0.02 <0.031
54. isophorone	113	7	ND	*			*
66. bis(2-ethylhexyl) phthalate	23 113	7 7	0.049	0.015 0.06			0.015 0.06
67. butyl benzyl phthalate	23 113	7 7		* ND			*
68. di-n-butyl phthalate	23 113	7 7	0.011	*			*
70. diethyl phthalate	23 113	7 7	ND	0.017 ND		2 	0.017 ND
71. dimethyl phthalate	23 113	7 7	ND	0.039 *		•	0.039

PRIMARY COLUMBIUM AND TANTALUM SECT

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA METAL SALT DRYING SCRUBBER, REDUCTION OF SALT TO METAL, AND REDUCTION OF SALT TO METAL SCRUBBER RAW WASTEWATER

			-	Concentrations (mg/1, Except as Noted)				
r	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average
81.	phenanthrene	23 113	7	*	ND *			*
85.	tetrachloroethylene	23	2		<0.013	ND	ND	<0.013
87.	trichloroethylene	23 113	2 2	*	ND *	* *	ND *	*
108.	PCB-1221	113	7	**	0.0516			0.0516
112.	PCB-1016	113	7	**	0.0336			0.0336
114.	antimony	23 113	7 7	<0.1	4.5 <0.1	:		4.5 <0.1
115.	arsenic	23 113	7 7	<0.01	0.45 0.05			0.45 0.05
117.	beryllium	23 113	7 7	<0.001	<0.02 0.02		·.	<0.02 0.02
118.	cadmium	23 113	7	<0.002	<0.2 <0.02			<0.2 <0.02
119.	chromium	23 113	7 7	<0.005	<0.24 1		· · · · ·	<0.24 1

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA METAL SALT DRYING SCRUBBER, REDUCTION OF SALT TO METAL, AND REDUCTION OF SALT TO METAL SCRUBBER RAW WASTEWATER

	C true om		Concentrations (mg/l, Except as Noted)					
Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
120. copper	23 113	7 7	0.066	<1 0.2			<1 0.2	
121. cyanide	-23	7		<0.001	0.004	0.003	0.0035	
122. lead	23 113	7 7 7	<0.02	0.004 10 1	0.002	0.018	0.008 10 1	
123. mercury	23 113	7 7	0.0001	0.0028 0.0018			0.0028 0.0018	
124. nickel	23 113	7 7	<0.005	1 · · · · · · · · · · · · · · · · · · ·	· · · · ·		1	
125. selenium	23 113	7 7	<0.01	0.018 <0.01			0.018 <0.01	
126. silver	23 113	7 7	<0.02	0.06			0.06	
127. thallium	23 113	7 7	<0.1	ND <0.1	- 1 		ND <0.1	
128. zinc	23 113	7 7	<0.06	6 <0 . 6			6 <0.6	

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PRIMARY COLUMBIUM AND TANTALUM SECT -

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA METAL SALT DRYING SCRUBBER, REDUCTION OF SALT TO METAL, AND REDUCTION OF SALT TO METAL SCRUBBER RAW WASTEWATER

			Concentrations (mg/l, Except as Noted)						
Pollutant	Stream Code	Sample Typet	Source Day 1 Day 2 Day 3 Aver	age					
Nonconventionals									
ammonia	23	2	25,700 18,500 16,900 20,377						
chemical oxygen demand (COD)	23 113		21 16.5 27.8 21.7 195 195	-					
chloride	113	7	<5 1,110 1,110						
fluoride	23 113	7 7	21,00021,0001.63,0003,000						
phenols (total; by 4-AAP method)	23 113	2 2	0.0280.0230.010.00.0050.0200.0050.0						
total organic carbon (TOC)	23 113	7 7	12 12 51 51						
Conventionals									
oil and grease	23 113	1 1	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	}					

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA METAL SALT DRYING SCRUBBER, REDUCTION OF SALT TO METAL, AND REDUCTION OF SALT TO METAL SCRUBBER RAW WASTEWATER

	A .	Sample Typet	Concentrations (mg/1, Except as Noted)					
Pollutant	Stream Code		Source	Day 1	Day 2	Day 3	Average	
total suspended solids (TSS)	23 113	7 7		181 656			181 656	
pH (standard units)	23 113	1	· · · · ·	6.8 6.1	8.3	8.2		

(a) Stream 113 was not analyzed for the toxic organic acid extractables.

(b) Stream 23 was not analyzed for the toxic organic acid extractable or pesticide fractions.

PRIMARY COLUMBIUM AND TANTALUM SECT <

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - V

TABLE V-13

WATER USE AND DISCHARGE RATES FOR REDUCTION OF TANTALUM SALT TO METAL

(1/kkg of tantalum salt reduced)

Plant Code	Percent Recycle	Production Normalized Water <u>Use</u>	Production Normalized Discharge Rate
507	0	272542	272542
513	0	170696	170696
519	0	54975	54975

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - V

TABLE V-14

WATER USE AND DISCHARGE RATES FOR REDUCTION OF TANTALUM SALT TO METAL WET AIR POLLUTION CONTROL

(l/kkg of tantalum metal reduced)

<u>Plant</u> Code	Percent <u>Recycle</u>	1 	Production Normalized Water <u>Use</u>	Production Normalized <u>Discharge</u> <u>Rate</u>
513	0	5 - 1 - 1	37826	37826
519	0		2043	2043

PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT A

		Stream	Sample	Cor	ncentrations	(mg/1, Ex	cept as Noted)	
	Pollutant	Code	Туре	Source	Day 1	Day 2	Day 3	Average
Toxic	Pollutants(a)				ł			
						·		10 01
115.	arsenic	50	2	<0.01	<0.01			<0.01
		51	2 2 2	<0.01	0.01			0.01
		52	2	<0.01	<0.01			<0.01
117.	beryllium	5Ô	2	-	<0.001			<0.001
11/•	Deryrrian	51	$\overline{2}$		<0.01		-	<0.01
	· · · ·	52	2 2 2		0.004			0.004
110	a admir m	50	2		0.009			0.009
118.	cadmium	51	. 2		<0.02			<0.02
		52	. 2 . 2 2		0.04			0.04
110	- hard and the second second	50	2		0.006			0.006
119.	chromium	50	2 2 2		0.15			0.15
		52	2		0.1			0.1
		JZ.	2		0.1			
120.	copper	50	2		1.0			1
1201	copper	51	2 2 2		0.79			0.79
		52	2		0.2			0.2
100	load	50	2		<0.02			<0.02
122.	lead	51	2		0.22			0.22
-		52	2 2 2		0.06			0.06
		50	2		0.0001			0.0001
123.	mercury	50	۲ 2		0.004	,	· · · · ·	0.004
		52	2 2		0.0001			0.0001
		77	L		0.0001			

PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT A

Pollutant	Stream Sample Code Type	Concentrations (mgSourceDay 1Day	/1, Except as Noted) y 2 Day 3 Average
<u>Conventionals</u>	•		<u> </u>
oil and grease	51 1 52 1	121 6	121 6
total suspended solids (TSS)	50 2 51 2 52 2	<1 1,012 52	<1 1,012 52
pH (standard units)	50 1 51 1 52 1	5.45 11.0 2.1	

(a) No samples were analyzed for toxic organic pollutants, cyanide, or asbestos.

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT B

				Concentrations (mg/1, Except as Noted)							
	Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average			
Toxic	Pollutants(a)										
1.	acenaphthene	116	7	ND	*			*			
4.	benzene	116	2	*	*	*	*	*			
8.	1,2,4-trichlorobenzene	116	2	ND	0.265			0.265			
- 10.	1,2-dichloroethane	- 116	2	ND .	ND .	0.017	ND	0.017			
23.	chloroform	116	2	0.422	*	ND	*	*			
30.	1,2-trans-dichloroethylene	116	2	ND	ND	0.094	0.017	0.056			
35.	2,4-dinitrotoluene	116	7	ND	*			*			
36.	2,6-dinitrotoluene	116	7	ND	*			*			
56.	nitrobenzene	116	7	ND	0.163			0.163			
66.	bis(2-ethylhexyl) phthalate	116	7	0.049	1.158			1.158			
71.	dimethyl phthalate	116	7	ND	*		•	*			
81.	phenanthrene	116	7	*	*			*			
	•	116	2	*	ND	ND	*	*			
87.	trichloroethylene	116	7	**	**			**			
108.	PCB-1221		7	**	**			**			
112.	PCB-1016	116	1								

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT B

		Stream Sample		Concentrations (mg/1, Except as Noted)					
	Pollutant	_Code	Typet	Source	Day 1	Day 2	Day 3	Average	
113.	toxaphene	116	7	ND	ND				
114.	antimony	116	7	<0.1	<0.1			<0.1	
115.	arsenic	116	7	<0.01	0.57			0.57	
117.	beryllium	116	7	<0.001	0.5	relative radio A		0.5	
118.	cadmium	116	7	<0.002	0.2	:	н — С.	0.2	
119.	chromium	116	7	<0.005	20			20	
120.	copper	116	7	<0.006	0.7	· ·		0.7	
121.	cyanide	116	7		0.002	0.002	0.002	0.002	
122.	lead	116	7	<0.02	10	* • •		10	
123.	mercury	116	.7	0.0001	0.0003			0.0003	
124.	nickel	116	7	<0.005	10			10	
125.	selenium	116	7	<0.01	0.02			0.02	
126.	silver	116	7	<0.02	0.05			0.05	
127.	thallium	116	7	<0.1	<0.1			<0.1	
128.	zinc	116	7	<0.06	1	·	•	1	

PRIMARY COLUMBIUM AND TANTALUM SECT -

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PRIMARY OOLUMBIUM-TANTALUM SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT B

	-		Concentrations (mg/l, Except as Noted)					
Pollutant	Stream Code	Sample Typet	Source	Day 1	Day 2	Day 3	Average	
Nonconventionals	9							
chemical oxygen demand (COD)	116	2		497		· .	497	
total organic carbon (TOC)	116 2		119			•	119	
Conventionals						·		
oil and grease	116	1		5	5	3	4.3	
total suspended solids (TSS)	116	2		436			436	
pH (standard units)	116	1		2.4	2.1	2.1		

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT C

		Stream	Sample	Concentrations (mg/l, Except as Noted)					
	Pollutant	Code	Typet	Source	Day 1	Day 2	Day 3	Average	
Toxic	e Pollutants			е — с - с - с			• •	•	
1.	acenaphthene	96	7			• * *	0.015	0.015	
4.	benzene	96	1	ND	<0.04	*	ND	<0.02	
6.	carbon tetrachloride		· · · · · · · · · · · · · · · · · · ·	ND	0.109	ND	0.035	0.072	
7.	chlorobenzene	96	. 1		ND	<0.029	0.065	0.033	
8.	1,2,4-trichlorobenzene	96	7		•	a.	0.018	0.018	,
10.	1,2-dichloroethane	96	1	ND	*	0.018	ND	0.009	
14.	1,1,2-trichloroethane	96	1	ND	ND	ND	*	*	j.
15.	1,1,2,2-tetrachloroethane	96	1	ND	<0.049	ND	ND	<0.049	
23.	chloroform	96	1	ND	0.059	0.018	*	0.026	
29.	1,1-dichloroethylene	96	1	ND	0.136	*	ND	0.068	
38.	ethylbenzene	96	1	ND	ND	ND	0.049	0.049	
48.	dichlorobromomethane	96	1	ND	0.016	ND	ND	0.016	
51.	chlorodibromomethane	96	. 1	ND	ND	*	ND	*	
66.	bis(2-ethylhexyl) phthalate	96	7	*			0.018	0.018	

PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT C

				Conce	ntrations	(mg/1, Exc	cept as No	oted)
	Pollutant	Stream Code	Sample <u>Typet</u>	Source	Day 1	Day 2	Day 3	Average
68.	di-n-butyl phthalate	96	7	*			*	*
71.	dimethyl phthalate	96	7	*			0.02	0.02
81.	phenanthrene	96	7				<0.012	<0.012
85.	tetrachloroethylene	96	1	ND	<0.046	0.189	0.137	0.109
86.	toluene	96	1	ND	0.092	*	0.012	0.035
87.	trichloroethylene	96	1	ND	<0.185	ND	0.020	0.010
108.	PCB-1221	96	7		-		**	**
111.	PCB-1060	96	7				**	**
113.	toxaphene	96	. 7	· .	ND		۶	
114.	antimony	96	, 7 -		· • •		<0.01	<0.01
115.	arsenic	96	7				0.18	0.18
117.	beryllium	96	7				0.02	0.02
118.	cadmium	-96	7 -	<0.002		-	. 0.008	0.008
119.	chromium	96	7				3	3

PRIMARY COLUMBIUM AND TANTALUM SECT

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT C

	Stream	Sample	Concentrations (mg/l, Except as Noted)					
Pollutant	Code	Typet	Source Day 1 Day 2 Day 3 Average					
120. copper	96	7	0.5 0.5					
121. cyanide	96	7	0.001 0.002 0.003 0.002					
122. lead	96	7	3 3					
123. mercury	96	7	0.0017 0.0017					
124. nickel	96	7	0.6 0.6					
125. selenium	96	7	<0.01 <0.01					
126. silver	96	7	<0.02 <0.02					
127. thallium	96	7	<0.1 <0.1					
128. zinc	96	7	6 6					
Nonconventionals								
ammonia	96	1	452 3,210 2,180 1,947					
chemical oxygen demand (COD)	96	7 · · ·	145 145					
chloride	96	7	118 118					
fluoride	96	7	2,200 2,200					
phenols (total; by 4-AAP method)	96	1	0.092 0.005 0.004 0.337					
total organic carbon (TOC)	96	7	45 45					

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PRIMARY COLUMBIUM-TANTALUM SAMPLING DATA MISCELLANEOUS - RAW WASTEWATER - PLANT C

		Sample Typet	Concentrations (mg/1, Except as Noted)						
Pollutant	Stream Code		Source	Day 1	Day 2	Day 3	Average		
Conventionals									
oil and grease	96	1		10	6	6	7.3		
total suspended solids (TSS)	96	1				566	566		

PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT A

נות	Stream	Sample	Concentrations (mg/1, Except as Noted)					
Pollutant	Code	Туре	Source	Day 1	Day 2	Day 3	Average	
Toxic Pollutants			•		e			
4. benzene	48 49	3 3	*	* 0 . 047	*	ND ND	* 0.0235	
6. carbon tetrachloride	48 49	3 3	ND ND	ND 0.043	ND ND	ND ND	0.043	
11. 1,1,1-trichloroethane	48 49	3 3	ND ND	ND 0.013	ND 0.046	ND ND	0.0295	
23. chloroform	48 49	3 3	0.032 0.032	0.069 0.03	ND *	0.027 0.038	0.048	
29. 1,1-dichloroethylene	48 49	3 3	ND ND	ND ND	ND 0.025	ND ND	0.025	
30. 1,2-transdichloro- ethylene	48 49	3 3	ND ND	ND ND	0.022 *	* ND	0.011 *	
48. dichlorobromomethane	48 49	3 3	ND ND	ND 0.089	ND ND	ND ND	0.089	
66. bis(2-ethylhexyl) phthalate	48 49	3 3	0.173 0.173	0.404 0.016	0.221 0.203	0.198 0.179	0.274 0.133	
67. butyl benzyl phthalate	48 49	3 3	ND ND	ND ND	* 0.047	* 0.043	* 0.045	

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PRIMARY COLUMBIUM AND TANTALUM

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PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT A

			Stream	Sample	Cor	ncentrations	(mg/l, Exc	ept as Noted)
		Pollutant	Code	Туре	Source	Day 1	Day 2	Day 3	Average
11	4.	antimony	48 49	3 3	<0.1 <0.1	<0.01 0.4	<0.1 <0.01	<0.01 <0.1	<0.04 0.4
11	15.	arsenic	48 49	3 3	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01	<0.01 <0.01
11	18.	cadmium	 48 49	3 3	ND ND	0.01	0.004	0.008 <0.02	0.007 <0.02
11	19.	chromium	48 49	3 3	ND ND	0.03 0.14	0.03	0.01 <0.05	0.023 0.14
12	20.	copper	48 49	3 3	ND ND	0.009 <0.06	0.03	0.007 <0.06	0.015 <0.06
13	21.	cyanide	48 49	3 3	-	<0.001 0.001	<0.001 0.002	<0.001 0.001	<0.001 0.0013
1:	22.	lead	48 49	3 3	ND ND	0.03 0.36	0.03	<0.02 <0.2	0.03 0.36
. 1	23.	mercury	48 49	3 3	ND ND	0.0001 0.0001	0.0001	0.0001 0.0002	0.0001 0.00015
1	24.	nickel	48 49	33	ND ND	0.04 <0.05	0.05	0.007 <0.05	0.032 <0.05

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PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT A

Pollutant	Stream Code	Sample Type	<u>Co</u> Source	ncentrations Day 1		pt as Note Day 3	ed) Average
126. silver	48 49	3 3	<0.02 <0.02	<0.02 0.05	<0.02 <0.02	<0.02 0.04	<0.02 0.045
128. zinc	48 49	3 3	ND ND	<0.06 <0.6	0.08	<0.06 <0.6	0.08 <0.6
<u>Conventionals</u>							
oil and grease	48 49	1	• • • • • • •	6 7	6 5	6 4	6.00 5.33

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PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT B

	Pollutant	Stream Code	Sample Type	Con Source	centrations Day 1	(mg/1, Exc Day 2	ept as Noted Day 3	d) Average
Toxic	Pollutants							
8.	1,2,4-trichlorobenzene	112 115	7 7	ND ND	0.011 ND			0.011
23.	chloroform	112 115	7 7	0.422 0.422	ND 0.048	0.022 0.03	0.02 0.032	0.02 0.037
30.	1,2-trans-dichloro- ethylene	112 115	7 7 7	ND ND	ND ND	0.21	ND .	0.21
51.	chlorodibromomethane	112 115	7 7	*	ND ND	0.019	ND	0.019
55.	naphthalene	112 115	7 7	ND ND	ND 0.084			0.084
66.	bis(2-ethylhexyl) phthalate	112 115	7 7	0.049 0.049	0.05 0.523			0.05 0.523
115.	arsenic	112 115	7 7	<0.01 <0.01	0.01 0.39			0.01 0.39
117.	beryllium	112 115	7 7	<0.001 <0.001	<0.001 0.09			<0.001 0.09
118.	cadmium	112 115	7	<0.002 <0.002	0.004		•	0.004
119.	chromium	112 115	7 7	<0.005 <0.005	0.008 3			0.008 3

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PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT B

	Stream Sample		Concentrations (mg/1, Except as Noted)						
Pollutant	Code	Туре	Source	Day 1	Day 2	Day 3	Average		
120. copper	112 115	7 7	<0.006 <0.006	0.01 0.4			0.01 0.4		
121. cyanide	112 115	7 7		0.002 0.007	0.002	<0.001 0.003	0.002 0.004		
122. lead	112 115	<u>7</u> 7	<0.02 <0.02	0.07 3	· · · · · · · · · · · · · · · · · · ·	nturtur ola zo zota. N	0.07 3		
123. mercury	112 115	7 7	0.0001 0.0001	<0.0001 0.0061			<0.0001 0.0061		
124. nickel	112 115	7 7	<0.005 <0.005	0.06 2	na series No series		0.06 2		
126. silver	112 115	7 7	<0.02 <0.02	0.02	1. A. A. A.		0.02		
128. zinc	112 115	7 7	<0.06 <0.06	<0.06 <0.6			<0.06 <0.6		
Nonconventionals	• • •								
ammonia	112 115	7 7	<0.1 <0.1	283 402	132 500	311 250	242 384		
chemical oxygen demand (COD)	112 115	7 7		44 408			44 408		

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PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT B

	Stream	Sample _Type_	Concentrations (mg/1, Except as Noted)						
Pollutant	Code		Source	Day 1	Day 2	Day 3	Average		
fluoride	112 115	7 7	1.6 1.6	16 3,525			16 3,525		
phenols (total; by 4-AAP method)	112 - 115	1 1		0.01 0.025	0.015	0.01	0.012 0.018		
total organic carbon (TOC)	112 115	7 7		9 118			9 118		
Conventionals									
oil and grease	112 115	1	-	- 4 5	3 6	5	3.5 5.3		
total suspended solids (TSS)	112 115	7		36 3,876			36 3,876		
pH (standard units)	112 115	1		9 3.6	8.8 4	8.8 3.4			

PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT D

Dollutant	Stream	Sample	Concentrations (mg/1, Except as Noted)						
Pollutant	Code	Туре	Source	Day 1	Day 2	Day 3	Average		
Toxic Pollutants					· · · · · · · · · · · ·				
10. 1,2-dichloroethane	24	2		0.019	0.024	ND	0.022		
15. 1,1,2,2-tetrachloro- ethane	24	2		ND	ND	ND			
23. chloroform	24	2		0.018	0.026	*	0.015		
30. 1,2-trans-dichloro- ethylene	24	2		ND	ND	ND			
38. ethylbenzene	24	2		*	*	ND	*		
44. methylene chloride	24	2		0.636	ND	ND	0.636		
47. bromoform	24	2		ND	ND	ND			
51. chlorodibromomethane	24	2		ND	ND	ND			
66. bis(2-ethylhexyl) phthalate	24	7		0.015			0.015		
67. butyl benzyl phthalate	24	7		0.08	*		0.08		
68. di-n-butyl phthalate	24	7	الاي والي الأنجار الجارية المالية إلى ال	0.02			0.02		
70. diethyl phthalate	24	7		ND					
71. dimethyl phthalate	24	7		ND	· · · · · · · · ·				

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PRIMARY COLUMBIUM AND

TANTALUM

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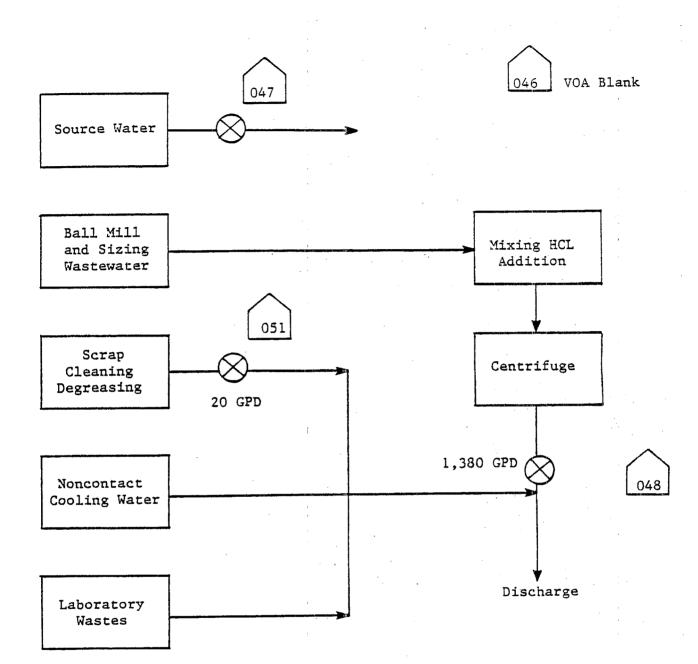
PRIMARY COLUMBIUM-TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT D

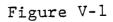
		cept as Note						
	Pollutant	Stream Code	Sample Type	Source	Day 1	Day 2	Day 3	Average
85.	tetrachloroethylene	24	2		0.024	ND	ND	0.024
87.	trichloroethylene	24	2		0.023	<0.024	ND	0.023
114.	antimony	24	7		0.2			0.2
115.	arsenic	··· · 24	7		0.45			0.45
117.	beryllium	24	7		<0.02			<0.02
118.	cadmium	24	7		<0.2			<0.2
119.	chromium	24	7		<0.24			<0.24
120.	copper	24	7		0.11			0.11
121.	cyanide	24	7	•	0.003	0.009	0.009	0.007
122.	lead	24	7		5			5
123.	mercury	24	7		0.0008			0.0008
124.	nickel	24	7		<0.5			<0.5
125.	selenium	24	7		0.045			0.045
128.	zinc	24	7		6			6

PRIMARY COLUMBIUM/TANTALUM - SAMPLING DATA MISCELLANEOUS - TREATMENT PLANT SAMPLES - PLANT D

Pollutant	Stream Code	Sample Type	<u>Co</u> Source	Distribution Day 1	s (mg/1, Ex Day 2	cept as Not Day 3	and the second
Nonconventionals					<u>Duy 1</u>	Day 5	Average
ammonia	24	2		21	16.5	27.8	21.77
chemical oxygen demand (OOD)	24	7		151	·	т. - ст ст	151
fluoride	24	7	•	6			6
phenols (total; by 4-AAP method)	24	2		0.031	0.03	0.03	0.031
total organic carbon (TOC)	24	7	•	27			27
Conventionals	,	· ·		• •	:		· · · · · · · · · · · · · · · · · · ·
oil and grease	24	1 .	· .	.4	4	4	4
total suspended solids (TSS)	24	7		89	,		89
pH (standard units)	24	1		12		ана салана 1973 — Салана 1973 — Салана 1974	

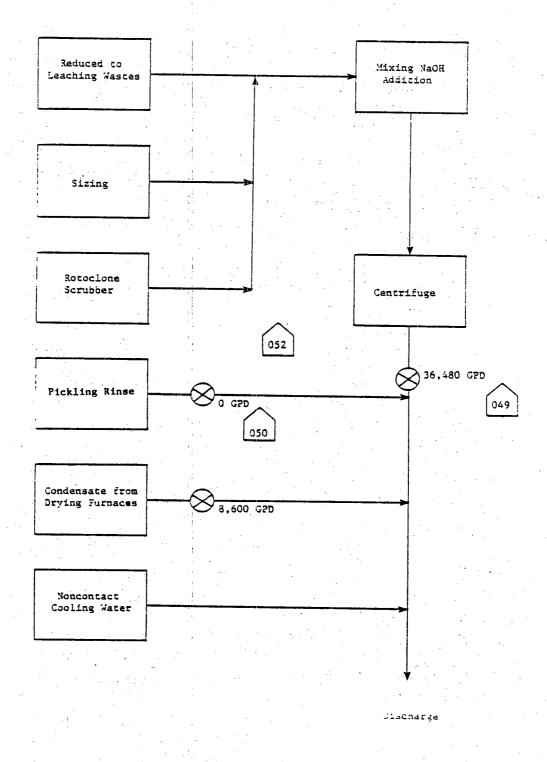
PRIMARY COLUMBIUM AND TANTALUM SECT -

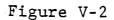




SAMPLING SITES AT COLUMBIUM-TANTALUM PLANT A

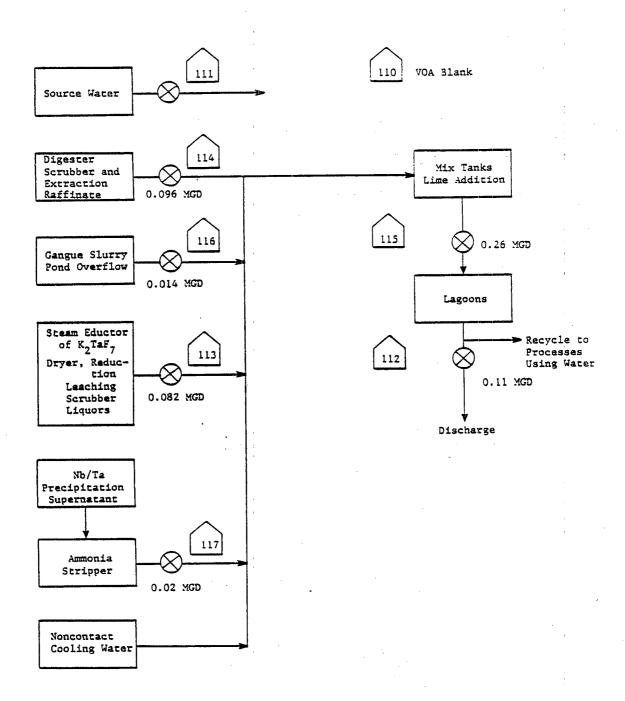
PRIMARY COLUMBIUM AND TANTALUM SECT - V

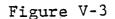




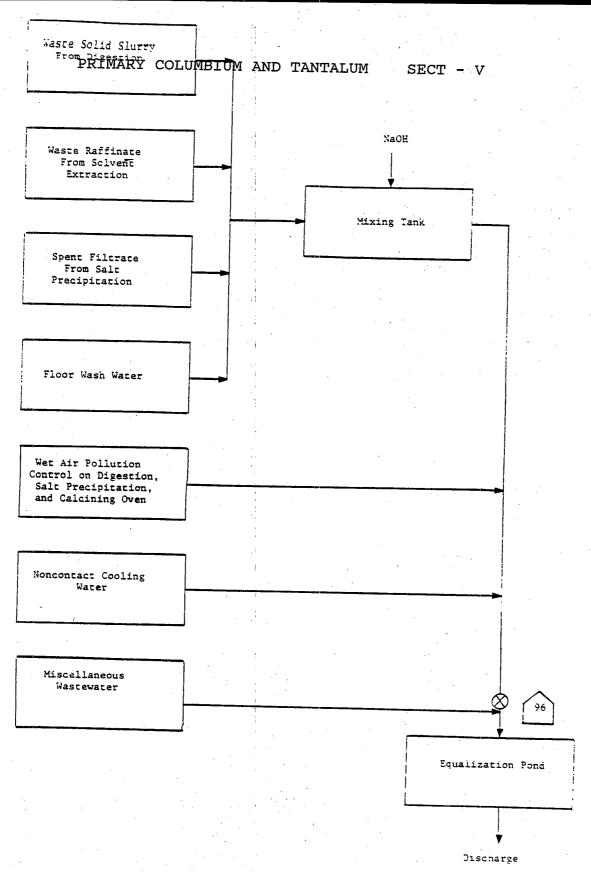
SAMPLING SITES AT COLUMBIUM-TANTALUM PLANT A (Continued)

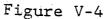
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SAMPLING SITES AT COLUMBIUM-TANTALUM PLANT B

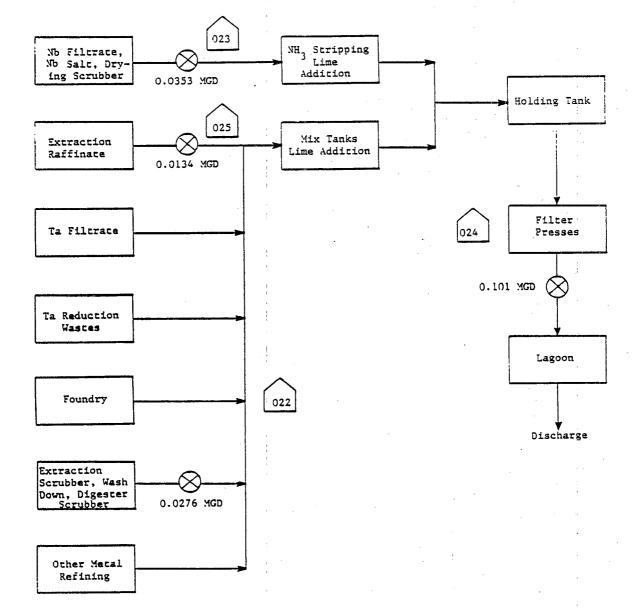


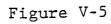


SAMPLING SITES AT COLUMBIUM-TANTALUM PLANT C

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SAMPLING SITES AT COLUMBIUM-TANTALUM PLANT D

SECT - VI

SECTION VI

SELECTION OF POLLUTANT PARAMETERS

This section examines the chemical analysis data presented in Section V and discusses the selection or exclusion of pollutants for potential limitation. The discussion that follows describes the analysis that was performed to select or exclude pollutants further consideration for limitations for and standards. Pollutants will be considered for limitations and standards if they are present in concentrations treatable by the technologies identified in this analysis. The treatable concentrations used for the toxic metals were the long-term performance values achievable by lime precipitation, sedimentation, and filtration. The treatable concentrations used for the toxic organics were the long-term performance values achievable by carbon adsorption.

After proposal, the Agency re-evaluated the treatment performance activated carbon adsorption to control toxic of organic pollutants. The treatment performance for the acid extractable, baseneutral extractable, and volatile organic pollutants has been equal to the analytical quantification limit of 0.010 set mq/l. analytical quantification limit for pesticides and total The phenols (by 4-AAP method) is 0.005 mg/l, which is below the 0.010 mg/l accepted for the other toxic organics. However, to be consistent, the treatment performance of 0.010 mg/l is used for pesticides and total phenols. The 0.010 mg/l concentration is achievable, assuming enough carbon is used in the column and a suitable contact time is allowed. The frequency of occurrence for 36 of the toxic pollutants has been redetermined based on the revised treatment performance value. As a result, the following pollutants, which were not selected at proposal, have been selected for further consideration for limitation:

- 4. benzene
- 6. carbon tetrachloride

85. tetrachloroethylene

CONVENTIONAL AND NONCONVENTIONAL POLLUTANT PARAMETERS SELECTED

This study examined samples from the primary columbium-tantalum subcategory for three conventional pollutant parameters (oil and grease, total suspended solids, and pH) and six nonconventional pollutant parameters (ammonia, chemical oxygen demand, chloride, fluoride, total organic carbon, and total phenols).

The following conventional and nonconventional pollutants or pollutant parameters are selected for consideration in establishing limitations for the columbium-tantalum subcategory:

ammonia total suspended solids (TSS) fluoride pH PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - VI

Five of eight samples analyzed for ammonia exhibited concentrations in excess of 40 mg/l (above the treatability concentration) with values reported as high as 3,210 mg/l. Since five of eight samples are above the 32 mg/l concentration attainable with steam stripping, ammonia is selected for further consideration.

The concentration of suspended solids in the ll samples for which it was analyzed ranged from 1 mg/l to 27,890 mg/l. Furthermore, most of the treatment used to remove toxic metals does so by precipitating the metals or their salts, and these toxic metal precipitates should not be discharged. A limitation on total suspended solids then, would help ensure that the toxic metals are removed. Thus, total suspended solids is selected for consideration for limitation.

Fluoride ions in low concentration (approximately 1.0 mg/l) are beneficial in drinking water supplies. However, higher concentrations (above 10 mg/l) can be harmful and even fatal to humans and animals. All four samples analyzed for fluoride contained very high concentrations of this pollutant (ranging from 2,800 to 24,000 mg/l). The identified treatment technology can reduce fluoride concentrations to 14.5 mg/l. Consequently, fluoride is selected for consideration for limitation.

The pH range measured was 1.87 to 11.0. Many deleterious effects are caused by either extreme pH values, or rapid changes in pH. Effective removal of toxic metals requires careful control of pH. Therefore, pH is considered for specific regulation in this subcategory.

TOXIC POLLUTANTS

The frequency of occurrence of toxic pollutants in the wastewater samples taken is presented in Table VI-1 (page 4435). These data provide the basis for the categorization of specific pollutants as discussed below. Table VI-1 is based on raw wastewater data from streams 22, 25, 113, 114, and 117 shown in Figures V-1 through V-5 and presented in Tables V-2, V-4, V-6, V-8, and V-12. Treatment plant samples were not considered in the frequency count. Streams 23, 48, 49, 50, 51, 52, 115, and 116 were not used because they contain either treated wastewater or wastewater from processes not considered for regulation in this rulemaking.

TOXIC POLLUTANTS NEVER DETECTED

The toxic pollutants listed in Table VI-2 (page 4439) were not detected in any wastewater samples from this subcategory; therefore, they are not selected for consideration in establishing limitations:

TOXIC POLLUTANTS NEVER FOUND ABOVE THEIR ANALYTICAL QUANTIFICATION LIMIT

The toxic pollutants listed below were never found above their analytical quantification concentration in any wastewater samples from this subcategory; therefore, they are not selected for consideration in establishing limitations.

14. 1,1,2-trichloroethane

- 15. 1,1,2,2-tetrachloroethylene
- 20. 2-chloronaphthalene
- 35. 2,4-dinitrotoluene
- 36. 2,6-dinitrotoluene
- 39. fluoranthene
- 67. butyl benzyl phthalate
- 73. benzo(a)pyrene
- 78. anthracene (a)
- 80. fluorene
- 81. phenanthrene (a)
- 113. toxaphene
- 121. cyanide

(a) Reported together as a combined value.

TOXIC POLLUTANTS PRESENT BELOW CONCENTRATIONS ACHIEVABLE BY TREATMENT

The pollutants listed below are not selected for consideration in establishing limitations because they were not found in any wastewater samples from this subcategory above concentrations considered achievable by existing or available treatment technologies. These pollutants are discussed individually following the list.

117. beryllium 126. silver

Beryllium was detected in five of six samples analyzed. However, it was found above its quantification limit in only two samples, both at concentrations below the treatable concentration of 0.20 mg/l for this pollutant. The concentrations of beryllium in the two samples were 0.18 and 0.02 mg/l. Therefore, beryllium is not selected for consideration for limitation.

Silver was detected in two of six samples analyzed, at values of 0.06 and 0.07 mg/l. However, treatment technology available cannot bring the silver concentration below 0.07 mg/l, so silver is not selected for consideration for limitation.

TOXIC POLLUTANTS DETECTED IN A SMALL NUMBER OF SOURCES

The following pollutants were not selected for regulation on the basis that they were detectable in the effluent from only a small number of sources within the subcategory and are uniquely related to only those sources.

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1. acenaphthene 12. hexachloroethane 23. chloroform 44. methylene chloride 47. bromoform 48. dichlorobromomethane 54. isophorone 56. nitrobenzene 66. bis(2-ethylhexyl) phthalate 68. di-n-butyl phthalate 71. dimethyl phthalate 106. PCB-1242 (a) PCB-1254 107. (a) 108. PCB-1221 (a) 109. PCB-1232 (a) 110. PCB-1248 (b) 111. PCB-1260 (b) 112. PCB-1016 (b)

123. mercury

(a), (b) Reported together as a combined value.

Acenaphthene was detected in one of seven samples, with the one detected value above the 0.010 mg/l concentration considered attainable with the identified treatment technology. The value detected in the sample was 0.017 mg/l. From the waste stream in which acenaphthene was detected, two other samples of this waste stream reported acenaphthene as a not detected. Therefore, acenaphthene is not considered characteristic of columbiumtantalum wastewaters and is not considered for limitation.

Hexachloroethane was present in only one out of seven samples taken, at 0.023 mg/l. Concentrations above 0.010 mg/l are considered treatable by the identified treatment technology. Also, in the dcp, all of the columbium-tantalum plants indicated that this pollutant was either known or believed to be absent. Therefore, hexachloroethane is not selected for consideration for limitation.

Chloroform, a common laboratory solvent, was detected in 10 of 14 samples, ranging from 0.017 to 0.24 mg/l. Concentrations above the analytical quantification limit in two of the three blanks (0.052 mg/l and 0.015 mg/l) analyzed raise the likelihood of sample contamination. Also, in the dcp, all of the columbiumtantalum plants indicated that this pollutant was either known or believed to be absent. Chloroform, therefore, is not selected for consideration for limitation.

One very high value of methylene chloride, 88.4 mg/l, was found in one of 14 samples; methylene chloride was not detected in the remaining 13 samples. But this solvent is so pervasive in laboratories that this one case of detection (out of 14) is probably due to sample contamination. The presence of methylene chloride in one of the blanks attests to this. Also, in the dcp,

all of the columbium-tantalum plants indicated that this pollutant was either known or believed to be absent. Therefore, methylene chloride is not selected for consideration for limitation.

Bromoform was found in two of the 14 raw waste samples. Only one of the concentrations found was above the 0.010 mg/l concentration considered attainable with identified treatment technology. The treatable value was 0.021 mg/l. The other concentration was below the analytical quantification limit. Therefore, bromoform is not selected for consideration for limitation.

Dichlorobromomethane was detected in one of the 14 samples, at a concentration above the 0.010 mg/l concentration considered attainable with identified treatment technology. The value detected in the sample was 0.038 mg/l. Two other samples from the waste stream in which dichlorobromomethane was reported were analyzed. The results were one "not detected" and one "detected below quantification concentration." Therefore, dichlorobromomethane is not considered characteristic of columbium-tantalum wastewaters and is not considered for limitation.

Isophorone was found in two of seven samples; only one was above the 0.010 mg/l concentration considered attainable with the identified treatment technology. The other sample had a concentration below the analytical quantification limit. The treatable value was 0.029 mg/l and was obtained from a sample of solvent extraction raffinate. Two other samples of solvent extraction raffinate were reported as not detected. Therefore, isophorone is not considered for limitation.

Nitrobenzene was detected in one of seven samples, and above the 0.010 mg/l concentration considered attainable with the identified treatment technology. The value detected was 0.1 mg/l. This value was obtained from a sample of solvent extraction raffinate in which two other samples were reported as not detected. Nitrobenzene, therefore, is not considered for limitation.

Bis(2-ethylhexyl) phthalate was reported present above its analytical quantification limit in five of seven samples; the reported concentrations ranged from 0.02 mg/l to 1.2 mg/l. This compound is a plasticizer found in many plastic materials used in manufacturing plants, thus it is not considered attributable to specific materials or processing in this subcategory. Also, in the dcp, all of the columbium-tantalum plants indicated that this pollutant was either known or believed to be absent. Therefore, bis(2-ethylhexyl) phthalate is not selected for consideration for limitation.

Di-n-butyl phthalate was measured above its concentration considered attainable with the identified treatment technology in three of seven samples; the measured concentrations ranged from

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0.012 mg/l to 0.08 mg/l. This substance is a plasticizer found in many products used in manufacturing plants; it is not considered a pollutant specific to this point source. Also, in the dcp, all of the columbium-tantalum plants indicated that this pollutant was either known or believed to be absent. Therefore, di-n-butyl phthalate is not selected for consideration for limitation.

Dimethyl phthalate was reported present above its analytical quantification limit in two of seven samples; the reported concentrations were 0.012 mg/l and 0.02 mg/l. This compound is a plasticizer found in many plastic materials used in manufacturing plants, and is not considered a point source specific pollutant. Also, in the dcp, all of the columbium-tantalum plants indicated that this pollutant was either known or believed to be absent. Therefore, dimethyl phthalate is not selected for consideration for limitation.

PCB-1242, PCB-1254, and PCB-1221 were measured above their analytical quantification limit in only one of seven samples. The observed concentration was 0.0516 mg/l. Since PCBs were found in just one plant, and since in the dcp, all of the columbium-tantalum plants indicated that this pollutant was either known or believed to be absent, they are not selected for consideration for limitation.

PCB-1232, PCB-1248, PCB-1260, and PCB-1016 were measured above their analytical quantification limit in one of seven samples. The observed concentration was 0.336 mg/l. Since PCB's were found in only one plant, and since in the dcp, all of the columbium-tantalum plants indicated that this pollutant was either known or believed to be absent, they are not selected for consideration for limitation.

Mercury was found above the concentration achievable by treatment in one of six samples. Only one sample at 0.063 mg/l was detected above the treatable concentration of 0.036 mg/l. Since the five other samples were below the treatable concentration, mercury is not selected for consideration for limitation.

TOXIC POLLUTANTS SELECTED FOR FURTHER CONSIDERATION FOR LIMITATIONS AND STANDARDS

The toxic pollutants listed below were selected for further consideration in establishing limitations and standards for this subcategory. The toxic pollutants selected are each discussed following the list.

- 4. benzene
- 6. carbon tetrachloride
- 7. chlorobenzene
- 8. 1,2,4-trichlorobenzene
- 10. 1,2-dichloroethane
- 30. 1,2-trans-dichloroethylene
- 38. ethylbenzene

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51. chlorodibromomethane 85. tetrachloroethylene 87. trichloroethylene 114. antimony 115. arsenic 116. asbestos 118. cadmium 119. chromium 120. copper 122. lead 124. nickel 125. selenium 127. thallium 128. zinc

Benzene was detected in 12 of 14 samples, with one of the concentrations above the 0.010 mg/l concentration considered attainable with the identified treatment technology. Eleven of these samples were below the quantification concentration. The value detected above the treatable concentration was 0.042 mg/l. Benzene was detected in four different process waste streams representing two plants. Therefore, benzene cannot be considered site-specific and is selected for further consideration for limitation.

Carbon tetrachloride was found in three of the 14 samples, with two of the concentrations above the 0.010 mg/l concentration considered attainable with identified treatment technology. The values found above the treatable concentration were 0.017 mg/l and 0.074 mg/l. Both of these values are from the same waste stream and represent two of the six samples from concentration digestion scrubber. Therefore, carbon tetrachloride is selected for further consideration for limitation.

Chlorobenzene was detected in three of 14 samples, with two of the concentrations above the 0.010 mg/l concentration considered attainable with the identified treatment technology. The values detected above the treatable concentration were 1.00 and 0.034 mg/l. Both of these values are from the same waste stream and represent two of the six samples analyzed from solvent extraction raffinate. Therefore, chlorobenzene is selected for further consideration for limitation.

1,2,4-Trichloroethylene was detected in two of seven samples, with one of the values above the 0.01 mg/l concentration considered attainable with the identified treatment technology. The value detected above its treatable concentration was 0.051 mg/l. Both samples in which 1,2,4-trichloroethylene was detected are from solvent extraction raffinate. Since the waste stream is from a solvent extraction process using an organic solvent, and 1,2,4-trichloroethylene was found above a treatable concentration, it is selected for further consideration for limitation.

1,2-Dichloroethane was detected in nine of 14 samples, with two

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of the concentrations above the 0.010 mg/l concentration considered attainable with the identified treatment technology. The values detected above the treatable concentration ranged from 0.016 mg/l to 0.156 mg/l. 1,2-Dichloroethane was detected above process waste quantification in five different streams representing two different plants. Therefore, 1,2-dichloroethane site-specific, and it is selected for is not further consideration for limitation.

1,2-trans-dichloroethylene was detected in two of 14 samples, both of the concentrations above the 0.010 with mq/1concentration considered attainable with the identified treatment technology. The values detected above the treatable concentration were 0.484 and 0.26 mg/l. These two values were taken from two different waste streams that were sampled three One of these streams is from solvent times each. extraction Therefore, where organic solvents are used. 1,2-transdichloroethylene is selected for consideration for limitation.

Ethylbenzene was detected in four of 14 samples, with two of the concentrations above the 0.010 mg/l concentration considered attainable with the identified treatment technology. The values detected above the treatable concentration were 0.04 mg/l and 0.057 mg/l. Ethylbenzene was detected in two different process waste streams representing two plants. Therefore, ethylbenzene is selected for further consideration for limitation.

Chlorodibromomethane was detected in four of 14 samples, with one of the concentrations above the 0.010 mg/l concentration considered attainable with the identified treatment technology. The values detected above its concentration considered attainable with identified treatment technology ranged from 0.02 to 7.08 mg/l. The detection of chlorodibromomethane was not sitespecific as it was detected in three different process wastewater streams representing two plants. Therefore, chlorodibromomethane is selected for further consideration for limitation.

Tetrachloroethylene was found in four of 14 samples, with two samples above the concentration considered attainable with identified treatment technology. The values detected above the treatable concentration were 0.157 mg/l and 0.235 mg/l. Tetrachloroethylene was found in three different process waste streams representing two plants. Therefore, this compound is selected for further consideration for limitation.

Trichloroethylene was detected in 12 of 14 samples, with one of the concentrations above the 0.010 mg/l concentration considered attainable with the identified treatment technology. Eleven of these samples were below the quantification concentration. The value detected above the treatable concentration was 0.235 mg/l. Trichloroethylene was detected in four different process waste streams representing two plants. Trichloroethylene cannot be considered site-specific and is therefore selected for further consideration for limitation.

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Antimony was found in five of seven samples analyzed; in all five of these, it was measured above its treatable concentration (0.47 mg/l) at concentrations ranging up to 30 mg/l. Therefore, antimony is selected for further consideration for limitation.

Arsenic was found in all six samples analyzed; three samples contained concentrations above its treatable concentration of 0.34 mg/l. Values were as high as 45 mg/l. Therefore, arsenic is selected for further consideration for limitation.

Analyses were made for asbestos at only one plant. The raw wastewater sample contained 980 million fibers per liter (MFL), while the plant influent contained less than 9 MFL. Since asbestos was detected and is above the treatable concentration of 10 MFL in the only sample analyzed, it is considered for further consideration for limitation.

Cadmium was detected in four of six samples, and was found above its treatable concentration of 0.049 mg/l. The concentration of cadmium in the sample was 40 mg/l. Cadmium, therefore, is selected for further consideration for limitation.

Five of six samples analyzed for chromium showed concentrations in excess of its treatable concentration (0.07 mg/l). Wastewater at one sampling site was found to contain 1,000 mg/l on each of three days sampled. Therefore, chromium is selected for further consideration for limitation.

Copper was found in all six samples analyzed, and occurred at concentrations above its treatable concentration of 0.39 mg/l in five of these. Values ranged from 0.8 to 300 mg/l. Therefore, copper is selected for further consideration for limitation.

Lead occurred far above its treatable concentration of 0.08 mg/l in five of six samples. Concentrations ranged from 1.0 to 1,000 mg/l. Lead, therefore, is selected for further consideration for limitation.

Two out of six samples analyzed for nickel yielded values above the treatable concentration of 0.22 mg/l. The reported concentrations were 5 and 1 mg/l. Therefore, nickel is selected for further consideration for limitation.

Selenium was found in three of six samples analyzed, all three above its treatable concentration (0.20 mg/l). Values were as high as 70 mg/l. Therefore, selenium is selected for further consideration for limitation.

Thallium was found above its treatable concentration of 0.34 mg/l in three of six samples, with concentrations of 0.83, 1.14, and 1.18 mg/l. Therefore, thallium is selected for further consideration for limitation.

Four of six samples analyzed contained zinc at concentrations above the treatability concentration of 0.23 mg/l. Values ranged

from less than 400 mg/l to 1,000 mg/l. Zinc is thus selected for further consideration for limitation.

Table VI-1

FREQUENCY OF OCCURRENCE OF TOXIC POLLUTANTS PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY RAW WASTEWATER

Pollutant		Analytica Quantificat Concentratio (mg/l)(a)	ion (on	Treatable Concentra- tion (mg/l)(b)	Number of Streams Analyzed	Numbe Samp Analy	les	ND	Quant	ted Below ification	Belc able	etected W Treat- e Concen- cation	Abov able	ected ve Treat- concen- ation	
1. acenaphthene		0.010		0.010	5	7		6	-					1	К
2. acrolein		0.010		0.010	5	14		14	1.1						Ω.
3. acrylonitrile	-	0.010	10 C 1	0.010	5	14		14	1.1						<u>o</u>
4. benzene		0'.010		0.010	5	. 14		2		11				1 .	2
5. benzidine		0.010	ς.	0.010	5	. 7		7	· ·	•••				• • • • •	M
6. carbon tetrachloric	е	0.010		0.010	5	14		11		1				2	μ
7. chlorobenzene		0.010		0.010	5	12		11		i				2	COLUMBIUM
8. 1,2,4-trichlorobenz	ene	0.010		0.010	5	7		5		1				1	R
9. hexachlorobenzene		0.010		0.010	5	7		7							
10. 1,2-dichloroethane	an ku si siya s	0.010		. 0.010 .	 5			- 5 -		. 1	· · ·		l in the second	- <u>8</u>	AND
11. 1,1,1-trichloroetha	ne	0.010		0.010	5	14		14	1.1.1		· .	•	2015 - 12 12	0	A
12. hexachloroethane		0.010	· · · · · ·	0.010	5	. 7		6					2 N.	1	1
13. 1,1-dichloroethane	1	0.010	*	0.010	5	14		- 14			1 - A - A - A - A - A - A - A - A - A -				۲.
14. 1,1,2-trichloroetha		0.010		0.010	5	14		12		2		-			TANTALUM
15. 1,1,2,2-tetrachloro	ethane	0.010		0.010	5	14		12	P1	2	•				H
16. chloroethane		0.010	÷	0.010	5	.14		14							$\overline{\mathbf{A}}$
17. bis(chloromethyl) e	ther	0.010		0.010	5	14	1.1.1	14				- 1 - E		÷ .	5
18. bis(2-chloroethyl)		0.010		0.010	 5	7		-7		-					. ₹
19. 2-chloroethyl vinyl	ether	0.010		0.010	5	14		14		•				•	
20. 2-chloronaphthalene		0.010		0.010	5	7		6		1.					
21. 2,4,6-trichlorophen	ol l	0.010		0.010	2	- 4		- 4		· · · ·					
22. parachlorometa cres	01	0.010		0.010	2	. 4		4				1 e			S
23. chloroform		0.010		0.010	5	14		4	1. 			100 C		10	E
24. 2-chlorophenol		0.010		0.010	2	- 4		-4							ECT
25. 1,2-dichlorobenzene		0.010		0.010	5	. 7		7.							
26. 1,3-dichlorobenzene	1	0.010		0.010	5 .			7							1
27. 1,4-dichlorobenzene	•	0.010		0.010	5			7				•		•	4
28. 3,3'-dichlorobenzid		0.010		0.010	5			. 7							ΤZ
29. 1,1-dichloroethylen	9 	0.010	1.1.1	0.010	5	14		14		· ·		·			• •
30. 1,2-trans-dichloroe	cnytene	0.010		0.010	5	14		12						2	
31. 2,4-dichlorophenol		0.010	*	0.010	- 2	. 4		4							
32. 1,2-dichloropropane		0.010	• •	0.010	5	- 14		14							
33. 1,3-dichloropropyle	ie .	0.010		0.010	5	- 14		14			1.111	· •		•	
34. 2,4-dimethylphenol		0.010	• •	0.010	2	- 4		. 4							
35. 2,4-dinitrotoluene	N	0.010		0.010	- 5	7	1971 - A.	6		1		· · ·		· · ·	. :
36. 2,6-dinitrotoluene		0.010		0.010	5	7		6		1				· · · ·	
37. 1,2-diphenylhydrazi	ie 	0.010		0.010	5	- 7		°7	· .						

Table VI-1 (Continued)

FREQUENCY OF OCCURRENCE OF TOXIC POLLUTANTS PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY RAW WASTEWATER

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Poll	<u>itant</u>	Analytical Quantification Concentration (mg/l)(a)	Treatable Concentra- tion (mg/1)(b)	Number of Streams Analyzed	Number of Samples Analyzed	ND	Detected Below Quantification Concentration	Detected Relow Treat- able Concen- tration	Detected Above Treat- able Concen- tration	
38. ethylbenzen	c	0.010	0.010	5	14	10	2		2	
39. fluoranthen		0.010	0.010	5	7	7				COLUMBIUM
	nyl phenyl ether	0.010	0.010	5	7	7				F
41. 4-bromophen		0.010	0.010	5	7	7				J.
	bisopropyl) ether	0.010	0.010	5	7	7				B
	vethoxy) methane	0.010	0.010	5	7	7				Н
44. methylene c		0.010	0.010	5	14	13			. 1	Ş
45. methyl chlo		0.010	0.010	5	14	14				
46. methyl brom		0.010	0.010	5	14	14				AND
47. bromoform		0.010	0.010	5	14	12	1			- Zi
48. dichlorobro	momethane	0.010	0.010	5	14	13			1	0
49. trichlorofl		0.010	0.010	5	14	14				H
	luoromethane	0.010	0.010	5	14	14				A
51. chlorodibro		0.010	0.010	5	14	10	3		I	E
52. hexachlorob		0.010	0.010	5	7	7				Ä
53. hexachloroc		0.010	0.010	5	· 7	7				브
54. isophorone)•	0.010	0.010	5	7	5	1		ł	TANTALUM
55. naphthalene		0.010	0.010	5	7	7			1	А
56. nitrobenzen		0.010	0.010	5	7	6			ł	
57. 2-nitropher		0.010	0.010	2	4	4				
58. 4-nitrophen		0.010	0.010	2	4	4				ດາ
59. 2.4-dinitro		0.010	0.010	2	4	4				SECT
60. 4.6-dinitro		0.010	0.010	2	4	4				G
61. N-nitrosodi		0.010	0.010	5	7	7				Ы
62. N-nitrosodi		0.010	0.010	5	7	7				1
63. N-nitrosodi		0.010	0.010	5	7	7		-		• •
64. pentachloro		0.010	0.010	2	4	4	. *			TΛ
65. phenol	•	0.010	0.010	2	-4	4			5	••
	hexyl) phthalate	0.010	0.010	5	7	1	1		C	
67. butyl benzy		0.010	0.010	5	7	7			3	
68. di-n-butyl		0.010	0.010	5	7	3	1		د	
69. di-n-octyl		0.010	0.010	5	7	7				
70. diethyl pht		0.010	0.010	5	7	7			1	
71. dimethyl pł		0.010	0.010	5	. 7 .	5	· · · · · ·		·····	
72. benzo(a)ant		0.010	0.010	· 5	7	7				
73. benzo(a)py		0.010	0.010	5	7	7				
74. 3,4-benzof		0.010	0.010	5	/	/				

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Table VI-1 (Continued)

FREQUENCY OF OCCURRENCE OF TOXIC POLLUTANTS PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY RAW WASTEWATER

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	Pollutant	. r	Analytical Quantification Concentration (mg/l)(a)	Treatable Concentra- tion (mg/l)(b)	Number of Streams Analyzed	Number of Samples Analyzed	<u>ND</u>	Detected Below Quantification Concentration	Detected Below Treat- able Concen- tration	Detected Above Treat- able Concen- tration	PRIMARY
76	 benzo(k)fluoranthene chrysene 		0.010 0.010	0.010 0.010	5 5	7 7.	7 7				COLUMBIUM
77 78	acenaphthylene anthracene (c)		0.010	0.010 0.010	5	7	7			-	LU1
- 79 80	<pre>. benzo(ghi)perylene</pre>		0.010	0.010	5	7	7	1			B
	fluorene (c)		0.010 0.010	0.010	· 5 ·	7	7			× 1	IUI
82	dibenzo(a,h)anthracene indeno(1,2,3-cd)pyrene		0.010 0.010	0.010 0.010	5	, 7 7	7	······································			
. 84.	pyrene tetrachloroethylene		0.010	0.010	5	7	7			· .	AND
86.	toluene		0.010 0.010	0.010 0.010	· 5	14 14	10 14	2		2	
	trichloroethylene vinyl chloride		0.010 0.010	0.010 0.010	5	14	2	- 11		1	TANTALUM
89.	aldrin		0.005	0.010	5	14	14 7	4 .			TA
	dieldrin chlordane	•	0.005 0.005	0.010 0.010	5	7	7		20		Ľ
	4,4'-DDT 4,4'-DDE		0.005	0.010	5	, 7	7		,	* :	Z,
<u>94</u> .	4,4'-DDD		0.005 0.005	0.010 0.010	5	7	7	н. -			
	alpha-endosulfan beta-endosulfan		0.005	0.010	. 5	ź	ź				Ŋ
97.	endosulfan sulfate		0.005 0.005	0.010 0.010	5	7.	-7	· .			SECT
. 98. 99.			0.005	0.010	5	7	2	•			Н
100.	heptachlor		0.005	0.010	5	7	.7				1
101.	heptachlor epoxide alpha-BHC		0.005	0.010 0.010	5	7	7				ΔI
103.	beta-BHC		0.005	0.010	5 5	. 7	1	р -			
104.	gamma-BHC delta-BHC		0.005	0.010	5 5	7	7			·	
106.	PCB-1242 (d) PCB-1254 (d)		0.005	0.010	5 5	7	, í	5	· ·	.1	
108.	PCB-1221 (d)	• •	0.005					•	1997 - L		
109.	PCB-1232 (e) PCB-1248 (e)		0.005	0.010	5	7	1	. j 5	·	1	
111.	(4)		0.005 0.005				-			· ·	•

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Table VI-1 (Continued)

FREQUENCY OF OCCURRENCE OF TOXIC POLLUTANTS PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY RAW WASTEWATER

<u>Pollutant</u>	Analytical Quantification Concentration (mg/l)(a)	Treatable Concentra- tion (mg/1)(b)	Number of Streams Analyzed	Number of Samples Analyzed	ND	Detected Below Quantification Concentration	Detected Below Treat- able Concen- tration	Detected Above Treat- able Concen- tration	RIMARY
<pre>113. toxaphene 114. antimony 115. arsenic 116. asbestos 117. beryllium 118. cadmium 119. chromium 120. copper 121. cyanide 122. lead 123. mercury 124. nickel 125. selenium 126. silver 127. thallium 128. zinc 129. 2,3,7,8-tetrachlorodibenzo- p-dioxin (TCDD)</pre>	0.005 0.100 0.010 10 MFL 0.010 0.002 0.005 0.009 0.100(g) 0.020 0.0001 0.005 0.01 0.02 0.100 0.050 Not Analyzed	0.010 0.47 0.34 10 MFL 0.20 0.049 0.07 0.39 0.047 0.08 0.036 0.22 0.20 0.07 0.34 0.23	5 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	7 6 6 1 6 6 6 6 6 6 6 6 6 6	2 2 1 2 1 1 4 3 4 3 2	5 3 11	3 2 1 4 2	4 3 1(f) 1 5 5 5 1 2 3 3 4	COLUMBIUM AND TANTALUM

(a) Analytical quantification concentration was reported with the data (see Section V).

(b) Treatable concentrations are based on performance of lime precipitation, sedimentation, and filtration for toxic metal pollutants and activated carbon adsorption for toxic organic pollutants.

(c),(d),(e) Reported together.

(f) 101+ MFL.

(g) Analytical quantification concentration for EPA Method 335.2, Total Cyanide Methods for Chemical Analysis of Water and Wastes, EPA-600/4-79-020, March 1979.

SECT -

4 H

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - VI

TABLE VI-2

TOXIC POLLUTANTS NEVER DETECTED

2.	acrolein
3.	acrylonitrile
5.	benzidine
9.	hexachlorobenzene
11.	1,1,1-trichloroethane
13.	1,1-dichloroethane
16.	chloroethane
17.	DELETED
18.	
	bis(2-chloroethyl) ether
19.	2-chloroethyl vinyl ether
21.	2,4,6-trichlorophenol
22.	parachlorometa cresol
24.	2-chlorophenol
25.	
	1,2-dichlorobenzene
26.	1,3-dichlorobenzene
27.	1 A-dichlorohonzono
	1,4-dichlorobenzene
28.	3,3'-dichlorobenzidine
29.	l,l-dichloroethylene
	2 4 dishistore ingrene
31.	2,4-dichlorophenol
32.	1,2-dichloropropane
33.	1 2-dichloropropulars
	1,3-dichloropropylene
34.	2,4-dimethylphenol
37.	l,2-diphenylhydrazine
	The arbuchymyarazine
10	
40.	4-chlorophenyl phenyl ether
40. 41.	4-chlorophenyl phenyl ether 4-bromophenyl phenyl ether
41.	4-bromophenyl phenyl ether
41. 42.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether
41. 42. 43.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane
41. 42.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane
41. 42. 43. 45.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride
41. 42. 43. 45. 46.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide
41. 42. 43. 45. 46. 49.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED
41. 42. 43. 45. 46.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide
41. 42. 43. 45. 46. 49. 50.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED
41. 42. 43. 45. 46. 49. 50. 52.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene
41. 42. 43. 45. 46. 49. 50. 52. 53.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene
41. 42. 43. 45. 46. 49. 50. 52.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene
41. 42. 43. 45. 46. 49. 50. 52. 53. 55.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol
41. 42. 45. 46. 49. 50. 52. 53. 55. 57. 58.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57.	4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol
41. 42. 45. 45. 50. 52. 53. 55. 57. 58. 59.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol</pre>
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57. 58. 59. 60.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol</pre>
41. 42. 45. 45. 50. 52. 53. 55. 57. 58. 59.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol</pre>
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57. 58. 59. 60. 61.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine</pre>
41. 42. 45. 45. 50. 52. 53. 55. 57. 58. 59. 60. 61. 62.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine</pre>
41. 42. 43. 45. 46. 50. 52. 53. 55. 57. 58. 59. 60. 61. 62. 63.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine N-nitrosodi-n-propylamine</pre>
41. 42. 45. 45. 50. 52. 53. 55. 57. 58. 59. 60. 61. 62.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine</pre>
41. 42. 43. 45. 46. 50. 52. 53. 55. 57. 58. 59. 60. 61. 62. 63. 64.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine N-nitrosodi-n-propylamine pentachlorophenol</pre>
41. 42. 43. 45. 46. 50. 52. 53. 55. 57. 58. 59. 61. 62. 63. 64. 65.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine N-nitrosodi-n-propylamine pentachlorophenol phenol</pre>
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57. 58. 59. 61. 62. 63. 64. 65. 69.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine N-nitrosodi-n-propylamine pentachlorophenol phenol di-n-octyl phthalate</pre>
41. 42. 43. 45. 46. 50. 52. 53. 55. 57. 58. 59. 61. 62. 63. 64. 65.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine N-nitrosodi-n-propylamine pentachlorophenol phenol di-n-octyl phthalate</pre>
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57. 58. 59. 61. 62. 63. 64. 65. 69. 70.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine N-nitrosodi-n-propylamine pentachlorophenol phenol di-n-octyl phthalate diethyl phthalate</pre>
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57. 58. 59. 61. 62. 63. 64. 65. 69. 70. 72.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodimethylamine N-nitrosodi-n-propylamine pentachlorophenol phenol di-n-octyl phthalate diethyl phthalate benzo(a)anthracene</pre>
41. 42. 43. 45. 46. 49. 50. 52. 53. 55. 57. 58. 59. 61. 62. 63. 64. 65. 69. 70.	<pre>4-bromophenyl phenyl ether bis(2-chloroisopropyl) ether bis(2-chloroethoxy) methane methyl chloride methyl bromide DELETED DELETED hexachlorobutadiene hexachlorocyclopentadiene naphthalene 2-nitrophenol 4-nitrophenol 2,4-dinitrophenol 4,6-dinitro-o-cresol N-nitrosodimethylamine N-nitrosodiphenylamine N-nitrosodi-n-propylamine pentachlorophenol phenol di-n-octyl phthalate diethyl phthalate</pre>

TABLE VI-2 (Continued)

TOXIC POLLUTANTS NEVER DETECTED

75. benzo(k)fluoranthene chrysene 76. 77. acenaphthylene benzo(ghi)perylene 79. dibenzo(a,h)anthracene 82. 83. indeno(1,2,3-cd)pyrene 84. pyrene 86. toluene vinyl chloride 88. aldrin 89. dieldrin 90. chlordane 91. 92. 4,4'-DDT 93. 4,4'-DDE 4, 4' - DDD94. alpha-endosulfan 95. beta-endosulfan 96. endosulfan sulfate 97. 98. endrin 99. endrin aldehyde 100. heptachlor 101. heptachlor epoxide 102. alpha-BHC 103. beta-BHC 104. gamma-BHC delta-BHC 105. 129. 2,3,7,8-tetrachlorodibenzo-p-dioxin (TCDD)

SECTION VII

CONTROL AND TREATMENT TECHNOLOGIES

The preceding sections of this supplement discussed the sources, flows, and characteristics of the wastewaters generated in the primary columbium-tantalum subcategory. This section summarizes the description of these wastewaters and indicates the level of treatment which is currently practiced for each waste stream.

CURRENT CONTROL AND TREATMENT PRACTICES

section presents a summary of the control and treatment This technologies that are currently applied to each of the sources generating wastewater in this subcategory. As discussed in Section V, wastewater associated with the primary columbiumtantalum subcategory is characterized by the presence of the toxic metal pollutants, ammonia, and suspended solids. This analysis is supported by the raw (untreated) wastewater data presented for specific sources as well as combined waste streams in Section V. Generally, these pollutants are present in each of the waste streams at concentrations above treatability, so these streams are commonly combined for treatment to reduce waste the concentrations of these pollutants. Construction of one wastewater treatment system for combined treatment allows plants to take advantage of economies of scale, and in some instances, combine streams of differing alkalinity to reduce treatment to chemical requirements. Four plants in this subcategory currently have lime and settle treatment systems. As such, three options were selected for consideration for BPT, BAT, BDT, and pretreatment in this subcategory after proposal, based on combined treatment of these compatible waste streams.

CONCENTRATE DIGESTION WET AIR POLLUTION CONTROL

All three plants which practice digestion use hydrofluoric acid leach the columbium and tantalum ore concentrates. The to leachate goes to solvent extraction. Wet scrubbers are used at all three plants, one with recycle (86 percent) and a bleed stream, and two with once-through water usage. Wet scrubbers are necessary due to the acidic nature of the emissions and the presence of gaseous fluoride. The scrubber liquor has treatable concentrations of suspended solids, fluoride and metals. The addition of alkali is used in all cases to reduce these high concentrations. Existing wastewater treatment schemes for this waste stream are lime precipitation and sedimentation, and neutralization with caustic.

SOLVENT EXTRACTION RAFFINATE

After methyl isobutyl ketone extraction the barren raffinate must be treated. One plant of the three plants with this wastewater recycles a portion of the raffinate to the leaching process to utilize the acidic nature of this waste stream. The raffinate has characteristics similar to the concentrate digestion scrubber liquor. This stream is treated as follows:

- 1. Lime addition and sedimentation (partial recycle);
- Lime addition, sedimentation, and filtration (no recycle); and
- 3. Neutralization and equalization pond (no recycle).

SOLVENT EXTRACTION WET AIR POLLUTION CONTROL

This waste stream is generated by wet air pollution control equipment located over the solvent extraction process. Two plants use wet scrubbers to control solvent extraction air emissions. One plant does not recycle the scrubber effluent; the other plant uses the same scrubber for solvent extraction and concentrate digestion, practicing 86 percent recycle. Waste characteristics are very similar to those found in the solvent extraction raffinate and concentrate digester scrubber waste streams; treatment similar to these two waste streams is Indeed, the established treatment techniques indicated. are identical:

- 1. Lime addition and sedimentation (partial recycle), and
- 2. Lime addition, sedimentation, (no recycle).

PRECIPITATION AND FILTRATION

The metal salts in the pregnant extraction solutions are precipitated either by oxide precipitation with ammonia or by potassium fluoride precipitation of potassium fluotantalate (K_2TaF_7) . The barren solutions must subsequently be treated. Three plants produce this wastewater. The wastewater contains treatable concentrations of ammonia, fluoride, metals, and suspended solids. The following wastewater treatment schemes are practiced for this stream:

- 1. Ammonia steam stripping, lime addition, and sedimentation (partial recycle); and
- 2. Neutralization and equalization pond (no recycle).

PRECIPITATION AND FILTRATION WET AIR POLLUTION CONTROL

This waste stream is generated by wet air pollution control equipment located over the precipitation process. Two plants use wet scrubbers to control precipitation air emissions. Neither plant recycles this wastewater. Waste characteristics are very similar to those found in the precipitation and filtration supernatant. One plant discharges this waste stream with no treatment, while the other uses ammonia steam stripping followed by lime and settle treatment.

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TANTALUM SALT DRYING

Only one of the five plants surveyed captures tantalum salt drying steam. This wastewater contains treatable concentrations of fluoride when potassium fluoride is used in precipitation. The treatment scheme used to treat tantalum salt drying wastewater by the one plant is lime and settle.

OXIDES CALCINING WET AIR POLLUTION CONTROL

Four of the five plants surveyed practice calcining of columbium pentoxide prior to shipment or further processing. Wet scrubbers are necessary to control ammonia and fluoride emissions during this process. Two plants practice recycle with recycle rates of 89 and 100 percent. Suspended solids and metals may be present in this wastewater in addition to ammonia and fluoride. The treatment schemes used to treat oxide calcining scrubber liquor by the four plants which practice calcining are as follows:

- Ammonia steam stripping and lime and settle (no recycle),
- Ammonia steam stripping and lime and settle (recycle), and
- 3. No treatment.

REDUCTION OF TANTALUM SALT TO METAL

Four plants reduce columbium or tantalum salts to the metal. One plant practices aluminothermic reduction, which produces no wastewater. The other three plants practice sodium reduction. Leaching after sodium reduction, a common practice for tantalum production, is a major source of wastewater. After completion of the reduction reaction and subsequent cooling, the tantalum exists as small particles of metal in a matrix of potassium and sodium salts. The salts are removed by successive leaches in water and acid to produce a pure metal powder. The resulting wastewater contains fluoride at treatable concentrations, as well as toxic metals and oil and grease. The wastewater treatment schemes used for this waste stream are as follows:

- 1. Lime addition and sedimentation, and
- 2. Caustic addition and centrifugation (no recycle).

REDUCTION OF TANTALUM SALT TO METAL WET AIR POLLUTION CONTROL

Wet scrubbers are used to control emissions during the reduction reaction. Two plants use wet scrubbers, neither practicing recycle of the scrubber liquor. This wastewater is similar in characteristic to the reduction wastewater. It contains toxic metals and fluoride and chloride in treatable concentrations. Treatment for the waste stream consists of:

Lime addition and sedimentation (no recycle), and
 Caustic addition and centrifugation (no recycle).

TANTALUM POWDER WASH

One plant washes tantalum powder after reduction and uses a scrubber in this process. No recycle was reported. This wastewater is expected to be acidic and contain little or no toxic pollutants. Currently this plant treats the wastewater using lime and settle treatment methods.

CONSOLIDATION AND CASTING CONTACT COOLING

Four plants reported consolidation and casting operations. One plant generates no wastewater. Two plants use noncontact cooling water. The fourth plant generates contact cooling water but recycles 100 percent through a cooling tower. Therefore, no wastewater is discharged for this waste stream.

CONTROL AND TREATMENT OPTIONS

The Agency examined three control and treatment technology alternatives since proposal that are applicable to the primary columbium-tantalum subcategory. The options selected for evaluation represent a combination of in-process flow reduction, pretreatment technology applicable to individual waste streams, and end-of-pipe treatment technologies.

OPTION A

Option A for the primary columbium-tantalum subcategory requires treatment technologies to reduce pollutant mass. The Option A treatment scheme consists of ammonia steam stripping preliminary treatment _applied to the combined streams of precipitation | and filtration of metal salts wastewater, precipitation and filtration wet air pollution control, and oxides calcining wet air pollution control. Preliminary treatment is followed by lime precipitation and sedimentation applied to the combined stream of steam stripper effluent and the remaining wastewater sources. Chemical precipitation is used to remove metals and fluoride by sedimentation. the addition of lime followed by gravity Suspended solids are also removed from the process.

OPTION B

Option B for the primary columbium-tantalum subcategory consists of all treatment requirements of Option A (ammonia steam stripping, lime precipitation, and sedimentation) plus control technologies to reduce the discharge of wastewater volume. Water recycle and reuse are the principal control mechanisms for 'flow reduction.

OPTION C

Option C for the primary columbium-tantalum subcategory consists of all control and treatment requirements of Option B (ammonia steam stripping, in-process flow reduction, lime precipitation, and sedimentation) plus multimedia filtration technology added at

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the end of the Option B treatment scheme. Multimedia filtration is used to remove suspended solids, including precipitates of metals. beyond the concentration attainable bv gravity The filter suggested is of the gravity, sedimentation. mixed media type, although other forms of filters such as rapid sand filters or pressure filters would perform as well. The addition of filters also provides consistent removal during periods in which there are rapid increases in flows or loadings of pollutants to the treatment system.

CONTROL AND TREATMENT OPTIONS REJECTED

Three additional treatment technologies were considered prior to proposing mass limitations for this subcategory as discussed below. Activated alumina and reverse osmosis were rejected because they were not demonstrated in the nonferrous metals manufacturing category, nor were they readily transferable. Activated carbon adsorption treatment did not receive further consideration because the levels of toxic organics present in the primary columbium-tantalum subcategory are present only in trace (deminimus quantities) and are neither causing nor likely to cause toxic effects.

OPTION D

Option D for the primary columbium-tantalum subcategory consisted of Option C (ammonia steam stripping, in-process flow reduction, lime precipitation, sedimentation, and multimedia filtration) with the addition of activated alumina technology at the end of the Option C treatment scheme. The activated alumina process is used to remove dissolved fluoride which remains after lime precipitation.

OPTION E

Option E for the primary columbium-tantalum subcategory consisted of Option C (ammonia steam stripping, in-process flow reduction, lime precipitation, sedimentation, and multimedia filtration) with the addition of granular activated carbon technology at the end of the Option C treatment scheme. The activated carbon process is utilized to control the discharge of toxic organics.

OPTION F

Option F for the primary columbium-tantalum subcategory consisted of Option C (ammonia steam stripping, in-process flow reduction, lime precipitation, sedimentation, and multimedia filtration) with the addition of reverse osmosis and multiple-effect evaporation technologies at the end of the Option C treatment scheme. Option F is used for complete recycle of the treated water by controlling the concentration of dissolved solids. THIS PAGE INTENTIONALLY LEFT BLANK

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - VIII

SECTION VIII

COSTS, ENERGY AND NONWATER QUALITY ASPECTS

This section describes the method used to develop the costs associated with the control and treatment technologies suggested in Section VII for wastewaters from primary columbium-tantalum plants. The energy consumption of each technology as well as solid waste and air pollution aspects are also discussed. Cost curves are presented in Section VIII of Vol. I showing the total annual cost of each treatment and control technology as a function of wastewater flow rate.

TREATMENT OPTIONS COSTED FOR EXISTING SOURCES

Three control and treatment options are considered for treating wastewater from the primary columbium-tantalum subcategory. Cost estimates have been developed for each of the control and treatment options. The options are summarized below and presented schematically in Figures X-1 through X-3 (pages 4494 - 4496).

OPTION A

Option A for the primary columbium-tantalum subcategory consists of lime precipitation and sedimentation end-of-pipe technology, with ammonia steam stripping preliminary treatment for waste streams containing treatable concentrations of ammonia. Streams with treatable concentrations of ammonia include precipitation and filtration of metal salts wastewater, precipitation and filtration scrubber water, and oxides calcining scrubber water.

OPTION B

Option B for the primary columbium-tantalum subcategory requires control and treatment technologies to reduce the discharge of wastewater volume and pollutant mass. The recycle of metal salt drying scrubber water, concentrate digestion scrubber, and solvent extraction scrubber water through holding tanks is the control mechanism for flow reduction. The Option B treatment scheme consists of ammonia steam stripping preliminary treatment for streams containing treatable concentrations of ammonia, and end-of-pipe treatment technology consists of lime precipitation and sedimentation.

OPTION C

Option C consists of all the control and treatment technologies of Option B (flow reduction, ammonia steam stripping, lime precipitation, and sedimentation) with the addition of multimedia filtration to the end-of-pipe treatment scheme.

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COST METHODOLOGY

detailed discussion of the methodology used to Α develop the compliance costs is presented in Section VIII of the General Development Document. Plant-by-plant compliance costs have been estimated for the nonferrous metal manufacturing category and are the administrative presented in record supporting this regulation. A comparison of the developed costs for proposal and the revised costs for the final regulation are presented in Tables VIII-1 and VIII-2 (pages 4452 and 4453) for the direct and indirect dischargers, respectively.

Each of the major assumptions used to develop compliance costs is presented in Section VIII of the General Development Document. However, each subcategory contains a unique set of waste streams requiring certain subcategory-specific assumptions to develop compliance costs. Seven major assumptions are discussed briefly below.

- (1) Several plants utilized sodium hydroxide addition for wastewater treatment. This type of treatment is not considered to be equivalent to lime addition due to the need to remove fluoride in the wastewater as calcium fluoride. The Agency therefore included compliance costs for treating with lime for these plants.
- (2) Ammonia steam stripping requirements may exceed the excess steam generation capacity at any given plant. Therefore, a steam generation unit is included in the steam stripping costs.
- (3) Due to the large volume of wastewater treatment sludge generated by some plants in this subcategory, the costs of developing and maintaining nonhazardous sludge disposal sites are used instead of the normal contract hauling.
- (4) EPA included the cost of segregation and treatment for one plant that currently commingles its wastewater and gangue. These costs eliminate any conceivable need for sludge disposal as a radioactive waste.
- (5) Recycle of air pollution scrubber liquor is based on recycle through a holding tank after lime and settle Annual costs associated with maintenance treatment. are included in the estimated compliance costs. If a plant currently recycles scrubber liquor, capital costs of the recycle equipment (holding tank, pumps, and piping) are not included in the compliance costs.
- (6) Subsequent to proposal, one columbium-tantalum plant commented that the selected lime and settle technology could not be installed on-site because of land limitations. Specifically, a clarifier of sufficient size could not fit within the available space at the plant. However, through Section 308 requests and telephone contacts, the Agency has determined that the clarifier could be installed at the

existing plant site. (This determination is documented in the administrative record supporting this regulation). Costs for the treatment technology were developed for this plant assuming a clarifier of sufficient size could be used.

(7) Annual costs for operation and maintenance of wastewater treatment systems are included in compliance costs for plants with treatment in place because there are no previous BPT or BAT regulations promulgated which account for these costs.

NONWATER QUALITY ASPECTS

A general discussion of the nonwater quality aspects of the control and treatment options considered for the nonferrous metals category is contained in Section VIII of the General Development Document. Nonwater quality impacts specific to the primary columbium-tantalum subcategory, including energy requirements, solid waste and air pollution are discussed below.

ENERGY REQUIREMENTS

Energy requirements for the three options considered are estimated at 5.22 mwh/yr, 5.22 mwh/yr, and 5.27 mwh/yr for Options A, B, and C respectively. Option C would increase energy requirements over Option A by approximately one percent. Option C represents roughly one percent of a typical plant's electrical usage. It is therefore concluded that the energy requirements of the treatment options considered will have no significant impact on total plant energy consumption.

SOLID WASTE

associated with the primary columbium-tantalum Sludges subcategory will necessarily contain additional quantities (and concentrations) of toxic metal pollutants. Wastes generated by smelters and refiners are currently primary exempt from regulation by Act of Congress (Resource Conservation and Recovery (RCRA)), Section 3001(b). Consequently, sludges generated Act from treating primary industries' wastewater are not presently subject to regulation as hazardous wastes.

Sludge generation in the primary columbium-tantalum subcategory is due to the precipitation of metal hydroxides and carbonates along with calcium fluoride using lime. If a small excess of lime is added during treatment, the Agency does not believe these sludges would be identified as hazardous under RCRA.

The Agency received comments stating that wastewater treatment sludges generated in the primary columbium-tantalum subcategory would have to be disposed of as low level radioactive waste. There are no RCRA regulations applicable to low level radioactive wastes, so the claim appears exaggerated. The Agency, therefore, requested specific data and information from the commenters so that the comments could be properly evaluated. However, no data or information were submitted to support this claim. In fact, one commenter submitted information and data showing the cost of disposal for gangue, the waste material remaining after the columbium-tantalum values are extracted from the raw material, rather than for wastewater treatment sludge. In any case, the Agency believes the disposal of gangue as a low level radioactive material is an expense of doing business and not attributable to the treatment of wastewaters.

Commenters in the secondary aluminum subcategory claim stripped ammonia will have to be disposed of as corrosive hazardous waste. The Agency does not agree with the commenters because ammonia has an intrinsic value. In the columbium-tantalum subcategory, ammonia is a process chemical and may be reused as a precipitating agent.

Although it is the Agency's view that solid wastes generated as a result of these guidelines are not expected to be hazardous, generators of these wastes must test the waste to determine if the wastes meet any of the characteristics of hazardous waste (see 40 CFR 262.11).

If these wastes should be identified or are listed as hazardous, they will come within the scope of RCRA's "cradle to grave" hazardous waste management program, requiring regulation from the point of generation to point of final disposition. EPA's of generator standards would require generators hazardous nonferrous metals manufacturing wastes to meet containerization, labeling, recordkeeping, and reporting requirements; if plants dispose of hazardous wastes off-site, they would have to prepare a manifest which would track the movement of the wastes from the generator's premises to a permitted off-site treatment, storage, or disposal facility. See 40 CFR 262.20 45 FR 33142 (May 19, 1980), as amended at 45 FR 86973 (December 31, 1980). The transporter regulations require transporters of hazardous wastes to comply with the manifest system to assure that the wastes are delivered to a permitted facility. See 40 CFR 263.20 45 FR 33151 (May 19, 1980), as amended at 45 FR 86973 (December 31, 1980). Finally, RCRA regulations establish standards for hazardous waste treatment, storage, and disposal facilities allowed to receive such wastes. See 40 CFR Part 464 46 FR 2802 (January 12, 1981), 47 FR 32274 (July 26, 1982).

Even if these wastes are not identified as hazardous, they still must be disposed of in compliance with the Subtitle D open dumping standards, implementing 4004 of RCRA. See 44 FR 53438 (September 13, 1979). The Agency has calculated as part of the costs for wastewater treatment the cost of hauling and disposing of these wastes. EPA estimates implementation of lime, settle, and filter technology will produce approximately 25,000 tons per year of sludge at 20 percent solids. Multimedia filtration technology will not result in any significant amount of sludge over that generated by lime precipitation.

AIR POLLUTION

There is no reason to believe that any substantial air pollution problems will result from implementation of chemical precipitation, sedimentation, and multimedia filtration. These technologies transfer pollutants to solid waste and do not involve air stripping or any other physical process likely to transfer pollutants to air.

Table VIII-1

COST OF COMPLIANCE FOR THE COLUMBIUM-TANTALUM SUBCATEGORY DIRECT DISCHARGERS

(March 1982 Dollars)

	Proposal	Costs	Promulgation Costs			
<u>Option</u>	Capital Cost	Annual Cost	Capital Cost	<u>Annual Cost</u>		
А	0	0	680,000	1,139,000		
B	117,000	18,000	736,000	1,157,000		
С	1,083,000	538,000	830,000	1,202,000		

Table VIII-2

COST OF COMPLIANCE FOR THE COLUMBIUM-TANTALUM SUBCATEGORY INDIRECT DISCHARGERS

(March 1982 Dollars)

	Proposa	l Costs	Promulgation Costs			
Option	Capital Cost	Annual Cost	Capital Cost	Annual Cost		
A	408,000	207,000	951,000	667,000		
В	627,000	238,000	979,000	674,000		
С	2,977,000	1,835,000	1,035,000	701,000		

PRIMARY COLUMBIUM AND TANTALUM

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - VIII

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SECTION IX

BEST PRACTICABLE CONTROL TECHNOLOGY CURRENTLY AVAILABLE

This section defines the effluent characteristics attainable through the application of best practicable control technology currently available (BPT), BPT reflects the existing performance by plants of various sizes, ages, and manufacturing processes within the primary columbium-tantalum subcategory, as well as the established performance of the recommended BPT systems. Particular consideration is given to the treatment already in place at plants within the data base.

The factors considered in identifying BPT include the total cost of applying the technology in relation to the effluent reduction benefits from such application, the age of equipment and facilities involved, the manufacturing processes used, nonwater quality environmental impacts (including energy requirements), and other factors the Administrator considers appropriate. In general, the BPT level represents the average of the existing performances of plants of various ages, sizes, processes, or common characteristics. Where existing performance is other uniformly inadequate, BPT may be transferred from a different subcategory or category. Limitations based on transfer of technology are supported by a rationale concluding that the technology is, indeed, transferable, and a reasonable prediction that it will be capable of achieving the prescribed effluent limits. BPT focuses on end-of-pipe treatment rather than process changes or internal controls, except where such practices are common industry practice.

TECHNICAL APPROACH TO BPT

The Agency studied the nonferrous metals category to identify the processes used, the wastewaters generated, and the treatment processes installed. Information was collected from the category using data collection portfolios, and specific plants were sampled and the wastewaters analyzed. Additional data used in the final rule were obtained through comments and Section 308 Some of the factors which must be considered in requests. establishing effluent limitations based on BPT have already been The age of equipment and facilities, processes used, discussed. and raw materials were taken into account in subcategorization and subdivision and are discussed fully in Section IV. Nonwater quality impacts and energy requirements are considered in Section VIII.

As explained in Section IV, the primary columbium-tantalum subcategory has been subdivided into 11 potential wastewater sources. Since the water use, discharge rates, and pollutant characteristics of each of these wastewaters is potentially unique, effluent limitations will be developed for each of the 11 subdivisions.

each of the subdivisions, a specific approach was For followed for the development of BPT mass limitations. To account for production and flow variability from plant to plant, a unit of production or production normalizing parameter (PNP) was determined for each waste stream which could then be related to flow from the process to determine a production normalized the flow. Selection of the PNP for each process element is discussed Section IV. Each process within the subcategory was in then analyzed to determine (1) whether or not operations included generated wastewater, (2) specific flow rates generated, and (3) the specific production normalized flows for each process. This analysis is discussed in detail in Section \overline{V} . Nonprocess wastewater such as rainfall runoff and noncontact cooling water is not considered in the analysis.

Normalized flows were analyzed to determine which flow was to be used as part of the basis for BPT mass limitations. The selected flow (sometimes referred to as a BPT regulatory flow or BPT discharge rate) reflects the water use controls which are common practices within the category. The BPT normalized flow is based on the average of all applicable data. Plants with normalized flows above the average may have to implement some method of flow reduction to achieve the BPT limitations.

For the development of effluent limitations, mass limitations were calculated for each wastewater source or subdivision. This calculation was made on a stream-by-stream basis, primarily because plants in this subcategory may perform one or more of the operations in various combinations. The mass limitations (milligrams of pollutant per kilogram of production unit - mg/kg) were calculated by multiplying the BPT normalized flow (l/kkg) by the concentration achievable using the BPT treatment system (mg/l) for each pollutant parameter to be limited under BPT.

The mass limitations which are allowed under BPT for each plant will be the sum of the individual mass loadings for the various wastewater sources which are found at particular plants. Accordingly, all the wastewater generated within a plant may be combined for treatment in a single or common treatment system, but the effluent limitations for these combined wastewaters are based on the various wastewater sources which actually contribute to the combined flow. This method accounts for the variety of combinations of wastewater sources and production processes which may be found at primary columbium-tantalum plants.

The Agency usually establishes wastewater limitations in terms of mass rather than concentration. This approach prevents the use of dilution as a treatment method (except for controlling pH). The production normalized wastewater flow (l/kkg) is a link between the production operations and the effluent limitations. The pollutant discharge attributable to each operation can be calculated from the normalized flow and effluent concentration achievable by the treatment technology and summed to derive an appropriate limitation for each subcategory. PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

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BPT effluent limitations are based on the average of the discharge flow rates for each source; consequently, the treatment technologies which are currently used by the lowest dischargers will be the treatment technologies most likely required to meet BPT effluent limitations. Section VII discusses the various treatment technologies which are currently in place for each wastewater source. In most cases, the current treatment technologies consist of chemical precipitation and sedimentation (lime and settle technology) and a combination of reuse and recycle to reduce flow. Ammonia steam stripping is added to streams with treatable concentrations of ammonia.

The overall effectiveness of end-of-pipe treatment for the removal of wastewater pollutants is improved by the application of water flow controls within the process to limit the volume of wastewater requiring treatment. The controls or in-process technologies recommended under BPT include only those measures which are commonly practiced within the subcategory and which reduce flows to meet the production normalized flow for each operation.

In making technical assessments of data, reviewing manufacturing processes, and assessing wastewater treatment technology options, both indirect and direct dischargers have been considered as a single group. An examination of plants and processes did not indicate any process differences based on the type of discharge, whether it be direct or indirect.

INDUSTRY COST AND POLLUTANT REMOVAL ESTIMATES

In balancing costs in relation to pollutant removal estimates, EPA considers the volume and nature of existing discharges, the volume and nature of discharges expected after application of BPT, the general environmental effects of the pollutants, and the cost and economic impacts of the required pollution control level. The Act does not require or permit consideration of water quality problems attributable to particular point sources or industries, or water quality improvements in particular water quality bodies. Accordingly, water quality considerations were not the basis for selecting the promulgated BPT.

The methodology for calculating pollutant removal estimates and plant compliance costs is discussed in Section X. Table X-2 (page 4485) shows the estimated pollutant removal estimates for each treatment option for direct dischargers. Compliance costs for direct dischargers are presented in Table VIII-1 (page 4452).

BPT OPTION SELECTION

The BPT selected consists of chemical precipitation and sedimentation (lime and settle technology) with ammonia steam stripping preliminary treatment of wastewaters containing treatable concentrations of ammonia. The best practicable

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technology is presented in Figure IX-1 (page 4472). The BPT treatment is equivalent to Option A described in Section VII and does not differ from that proposed. Lime and settle technology is currently demonstrated by all three primary columbium-tantalum direct dischargers.

Ammonia steam stripping is demonstrated in the nonferrous metals manufacturing category and at two primary columbium-tantalum facilities. EPA proposed treatment performance concentrations based on levels achieved in the iron and steel manufacturing category.

Chemical analysis data were collected of raw waste (treatment influent) and treated waste (treatment effluent) from one coke plant of the iron and steel manufacturing category. A contractor for EPA, using EPA sampling and chemical analysis protocols, collected six paired samples in a two-month period. These data are the data base for determining the effectiveness of ammonia stripping technology and are contained within the steam administrative record supporting this document. Ammonia treatment at this coke plant consisted of two steam stripping columns in series with steam injected countercurrently to the flow of the wastewater. A lime reactor for pH adjustment separated the two stripping columns.

The raw untreated wastewater samples from the coke facility contained ammonia concentrations of 599, 226, 819, 502, 984, and 797 mg/l. Raw untreated wastewater samples from the primary columbium-tantalum subcategory contained ammonia concentrations of 53.1, 496.1, 25,700, 18,500, and 16,900 mg/l. These latter three concentrations represent three days of sampling from a calciner scrubber.

The Agency has verified the proposed steam stripping performance values using steam stripping data collected at a zirconiumhafnium plant in the nonferrous metals manufacturing category. Data collected by the plant represent almost two years of daily operations and support the long-term mean and variability used to establish treatment effectiveness. Although the ammonia concentrations in columbium-tantalum wastewater are higher than those in iron and steel wastewater, the columbium-tantalum ammonia data are comparable to raw wastewater data from this zirconium-hafnium plant.

The Agency has chosen not to regulate toxic organic pollutant parameters on a subcategory-wide basis for the primary columbiumtantalum subcategory. Primary columbium-tantalum plants may use an organic solvent in a liquid-liquid ion exchange process to extract columbium-tantalum from digested concentrates. In the pollutant reduction removals prior to proposal, it was estimated subcategory generates 170 kg/yr toxic that the organic pollutants. The Agency believes the toxic organic pollutants in the primary columbium-tantalum subcategory are present only in trace amounts and thus are not regulated on a subcategory-wide basis. However, it is possible toxic organic pollutants may be

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present in larger concentrations at an individual plant than the Agency sampling data indicate. Therefore, the permitting or control authority should check for the presence of toxic organic pollutants on a case-by-case basis and determine if they require treatment.

Agency has re-evaluated lime The and settle technology performance for fluoride removal. The proposed treatment performance for fluoride was transferred from the electrical and electronic component manufacturing (phase I) lime and settle mean performance. Commenters urged the Agency to transfer treatability Agency values from the inorganic chemical industry instead. The The Agency believes the electronics data base disagrees. more closely reflects the treatability of fluoride in nonferrous metals manufacturing wastewaters because of the type of fluoride chemicals The fluoride present in inorganic present. manufacturing (hydrofluoric acid production) exists as a complex fluoride mineral containing silicates and other compounds that complicate removal by lime precipitation. In nonferrous metals manufacturing and electronics, the fluoride disassociates in water to fluoride ion, which can be readily removed from solution by lime as calcium fluoride.

However, examination of the electronics data has led the Agency to conclude that the raw concentrations of fluoride in nonferrous metals manufacturing wastewaters more closely resemble the higher concentrations found in electrical and electronics phase II rather than phase I (49 FR 55690, December 14, 1983). Therefore, the Agency believes it is appropriate to use the mean performance and daily maximum variability developed for electronics phase II to establish treatment effectiveness for fluoride removal by lime and settle treatment.

BPT will result in the removal of an estimated 61,093 kg/yr of toxic pollutants, 1,692,000 kg/yr of conventional pollutants, 973,000 kg/yr of fluoride, and 941,000 kg/yr of ammonia from raw The estimated capital investment cost of BPT discharge levels. is \$0.68 million (March, 1982 dollars) and the estimated annual cost is \$1.1 million. These costs represent wastewater treatment equipment not currently in place.

WASTEWATER DISCHARGE RATES

BPT discharge rate is calculated for each subdivision based on А the average of the flows of the existing plants, as determined from analysis of dcp. The discharge rate is used with the achievable treatment concentration to determine BPT effluent Since the discharge rate may be different for each limitations. wastewater source, separate production normalized discharge rates each of the 11 wastewater sources are discussed below for and summarized in Table IX-1 (page 4465). The discharge rates are generally normalized on a production basis by relating the amount of wastewater generated to the mass of the intermediate product which is produced by the process associated with the waste stream question. These production normalizing parameters, or PNPs, in

are listed in Table IX-1 (page 4465).

Section V of this document further describes the discharge flow rates and presents the water use and discharge flow rates for each plant by subdivision.

CONCENTRATE DIGESTION WET AIR POLLUTION CONTROL

The proposed BPT wastewater discharge rate for concentrate digestion wet air pollution control was 10,915 l/kkg (2,618 gal/ton) of columbium-tantalum salt produced from digestion. This rate was allocated only for plants practicing wet air pollution control for concentrate digestion. Three plants reported wastewater discharges from concentrate digestion wet air pollution control, but dcp information provided by one plant was insufficient to calculate a discharge rate. Therefore, the BPT discharge rate was based on the average of two plants which discharged 8,692.4 and 13,135.5 l/kkg (2,084.5 and 3,150 gal/ton).

The proposed concentrate digestion wet air pollution control flow allowance has been revised based on new data and information received through comments and special information requests. It has been demonstrated to the Agency that water usage for concentrate digestion scrubbers correlates better with the mass of concentrate or slag digested than the product recovered from digestion. This is due to the different columbium-tantalum values contained in ore concentrates and tin slags. A plant processing tin slags, which contain much less columbium-tantalum than ore concentrates, would digest much more raw material than a plant using ore concentrates to recover equal amounts of columbium-tantalum. Therefore, the Agency has changed the production normalizing parameter from columbium-tantalum salt recovered to the mass of raw material digested.

There are three values available from which to determine the discharge rate as shown in Table V-1 (page 4372). Plant 519 will not be used because it scrubs air emissions from digestion and solvent extraction. Water use in fume control is directly related to the volume of gas being scrubbed; consequently, plant 519 water usage may not be comparable to the other two plants. Water use at plant 509 will not be used because it discharges 16 more times water (on a production normalized basis) than plant 507. There is no engineering reason this discharge rate should be so high. The BAT discharge rate is thus set equal to the demonstrated rate at plant 507, or 6,219 1/kkg (1,491 gal/ton) of raw material digested.

SOLVENT EXTRACTION RAFFINATE

The proposal BPT wastewater discharge rate for solvent extraction raffinate was 26,916 1/kkg (6,470.4 gal/ton) of columbium or tantalum salt extracted. This rate was based on the average discharge rate of two plants, which discharged 19,268 and 34,694 1/kkg (4,620 and 8,320 gal/ton). A third plant reported

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insufficient data to calculate a discharge rate.

The proposal solvent extraction raffinate flow allowance has been revised based on new data and information received through and specific requests for information. comments As with concentrate digestion wet air pollution control, it has been demonstrated to the Agency water usage correlates better with the mass of concentrate or slag digested than the product recovered from solvent extraction. The production normalized discharge flows for plants 507 and 509 changed from proposal because of this change in production normalizing parameter. Plant 509 submitted data in response to a request, allowing a discharge flow to be calculated. Water usage rates for solvent extraction, production normalized with the mass of raw material digested, are presented in Table V-3. The promulgated discharge rate is based on the average of the three values, or 9,155 1/kkg (2,195 gal/ton) of raw material digested.

SOLVENT EXTRACTION WET AIR POLLUTION CONTROL

The proposal BPT discharge rate for solvent extraction wet air pollution control was 4,301 1/kkg (1,034 gal/ton) of columbium or tantalum salt extracted. This rate was allocated only for plants practicing wet air pollution control for solvent extraction. Two plants reported this wastewater, however, one plant uses the same scrubber for both solvent extraction and concentrate digestion wet air pollution control. The BPT discharge rate was based on the discharge of the single plant which only scrubbed emissions from solvent extraction.

As with concentrate digestion wet air pollution control, the discharge rate for solvent extraction raffinate wet air pollution control has been revised. Water use and discharge rates for this stream are presented in Table V-5. Plant 519 was not considered the determination of the BPT discharge rate because it in uses same scrubber to control emissions concentrate the from digestion. Water use for this scrubber is expected to be larger because of the added volume of gas scrubbed versus that for plant 507. The BPT discharge rate is set equal to the discharge rate 507 of 2,456 l/kkg (589 gal/ton) of raw material at plant Plant 519, however, should receive both digested. the concentrate digestion and solvent extraction wet air pollution control flow allowance.

PRECIPITATION AND FILTRATION OF METAL SALTS

The proposal BPT wastewater discharge rate for precipitation and filtration waste streams was 247,223 l/kkg (59,428 gal/ton) of columbium or tantalum salt precipitated. Three plants reported producing this waste stream. The BPT discharge rate was based on the discharge rate of one of the plants. The two other plants reported insufficient data to calculate a discharge rate.

Data and information were received through comments and specific requests for additional information so that production normalized

flows are available for three plants. Table V-7 (page 4387) presents production normalized discharges for the three plants. The production normalizing parameter for precipitation and filtration has also been changed to the mass of raw material digested rather than the columbium-tantalum recovered through precipitation. Water discharge and production were found to correlate much better with the mass of raw material used as the production normalizing parameter. The BPT discharge rate is determined from the average of the three reported values, or 13,689 l/kkg (3,283 gal/ton) of raw material digested.

PRECIPITATION AND FILTRATION WET AIR POLLUTION CONTROL

A discharge allowance for precipitation and filtration wet air pollution control has been added to cover wastewater generated from fume scrubbing during precipitation. This wastewater stream was not considered at proposal. Two plants reportedly operate this scrubber as shown in Table V-9 (page 4391). The BPT discharge is the average discharge rate of these two plants, or 63,513 1/kkg (15,231 gal/ ton) of raw material digested.

TANTALUM SALT DRYING WET AIR POLLUTION CONTROL

The proposal BPT wastewater discharge rate for metal salt drying wet air pollution control was 83,643 l/kkg (20,106 gal/ton) of columbium or tantalum salt dried. This rate was allocated only for plants practicing wet air pollution control for metal salt drying emissions. Four plants discharged a metal salt drying wet air pollution control waste stream. Two plants discharging this waste stream reported sufficient dcp information to calculate a discharge rate. The two plants generated 11,563 and 156,125 l/kkg (2,773 and 37,440 gal/ton) respectively, of metal salt drying wet air pollution wastewater. The BPT discharge rate was the average discharge rate of these two plants.

Based on the information obtained through comments and special requests for additional information, the proposed metal salt drying scrubber allowance has been divided into tantalum salt drying and oxides calcining wet air pollution control. Two plants reported capturing steam generated during tantalum salt drying. One of these plants reported its data in conjunction with oxides calcining wet air pollution control, and the data cannot be separated. Therefore, the discharge rate is set equal to one reported value of 60,542 l/kkg (14,518 gal/ton) of tantalum salt dried.

OXIDES CALCINING WET AIR POLLUTION CONTROL

A separate flow allowance is established for the calcining operations used to dry columbium-tantalum pentoxide. Four plants reported using a scrubber to control emissions from this operation as shown in Table V-11 (page 4393). One plant currently recycles at an 89 percent rate, one plant reported using treated effluent as scrubber liquor make-up (therefore 100 percent recycle), and two plants reported using once-through

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systems. The BPT discharge is based on the average water use at plants 4225, 509, and 507. This value is 38,422 l/kkg (9,214 gal/ton) of oxide production from calcining.

REDUCTION OF TANTALUM SALT TO METAL

The proposal BPT wastewater discharge rate for reduction of salt to metal was 352,663 l/kkg (84,775 gal/ton) of columbium or tantalum reduced. This rate was based on the average discharge rate of two plants, which discharged 170,740 and 536,282 l/kkg (40,945 and 128,605 gal/ton). A third plant reported insufficient dcp information to calculate a discharge rate.

Based on a re-evaluation of data from dcp and trip reports available to the Agency before proposal, discharge rates are now available for three plants as shown in Table V-13 (page 4400). The BPT discharge is thus chosen as the average discharge rate, or 166,071 1/kkg (39,825 gal/ton) of salt produced. Two of the plants currently meet this discharge rate.

REDUCTION OF TANTALUM SALT TO METAL WET AIR POLLUTION CONTROL

The proposal BPT wastewater discharge rate for reduction of salt to metal wet air pollution control was 21,521 l/kkg (5,173 gal/ ton) of columbium or tantalum reduced. This rate was allocated only for those plants practicing wet air pollution control for reduction emissions. The BPT discharge rate was based on the average discharge rate of the two plants reporting this wastewater. The two plants generated 2,168 and 40,978 l/kkg (520 and 9,827 gal/ton) respectively, of this wastewater.

No new information has been received on reduction of salt to metal wet air pollution control; however, this discharge allowance was re-evaluated to try to determine any reasons for the large variation in water usage. Both plants have similar reduction operations and both use rotoclone type scrubbers. Information available to the Agency does not suggest any reason for such a large variation in production normalized water usage. There fore, the BPT discharge is based on plant 519, which reports 19 times less water than plant 513. Thus, the BPT discharge rate is 2,043 l/kkg (490 gal/ton) of tantalum salt produced.

TANTALUM POWDER WASH AND SCRUBBER

A discharge allowance for tantalum powder wash and scrubber has been added to cover wastewater generated from tantalum powder washing after reduction. Only one plant reported this waste stream and the BPT discharge is set equal to the current discharge practices at this plant of 20,433 l/kkg (4,900 gal/ton) of tantalum powder washed. Flow data could not be separated from the scrubber and washing operation.

CONSOLIDATION AND CASTING CONTACT COOLING

No BPT wastewater discharge allowance was proposed for consolidation and casting contact cooling and no comments or data were received indicating a discharge allowance is warranted. Only one plant in this subcategory reported a consolidation and casting contact cooling waste stream. This plant does not discharge this wastewater. BPT is based on this plant.

REGULATED POLLUTANT PARAMETERS

The raw wastewater concentrations from individual operations and the subcategory as a whole were examined to select certain pollutant parameters for limitation. This examination and evaluation was presented in Section VI. A total of six pollutants or pollutant parameters were selected for limitation and are listed below:

122. lead
128. zinc
ammonia
fluoride
total suspended solids
pH

EFFLUENT LIMITATIONS

The treatment effectiveness achievable by application of the proposed BPT treatment are explained in Section VII of Vol. I and summarized there in Table VII-21 (page 248). The achievable treatment concentrations (both one day maximum and monthly average values) are multiplied by the BPT normalized discharge flows summarized in Table IX-1 (page 4465) to calculate the mass of pollutants allowed to be discharged per mass of product. The results of these calculations in milligrams of pollutant per kilogram of product represent the BPT effluent limitations and are presented in Table IX-2 (page 4467) for each individual waste stream.

Table IX-1

BPT WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

	Discha	rmalized rge Rate	
Wastewater Stream	1/kkg	gal/ton	Production Normalizing Parameter
Concentrate Digestion Wet Air Pollution Control	6,219	1,491	Concentrate Digested
Solvent Extraction Raffi- nate	9,155	2,195	Concentrate Digested
Solvent Extraction Wet Air Pollution Control	2,456	589	Concentrate Digested
Precipitation and Filtra- tion Wastewater	13,689	3,283	Concentrate Digested
Precipitation and Filtra- tion Wet Air Pollution Control	63,513	15,231	Concentrate Digested
Tantalum Salt Drying	60,542	14,518	Tantalum Salt Dried
Oxides Calcining Wet Air Pollution Control	38,422	9,214	Columbium or Tantalum Oxide Calcined
Reduction of Tantalum Salt to Metal	166,071	39,825	Tantalum Salt Reduced
Reduction of Tantalum Salt Metal Wet Air Pollution Control	2,043	490	Tantalum Salt Reduced

Table IX-1 (Continued)

BPT WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

		malized ge Rate	
Wastewater Stream	1/kkg	gal/ton	Production Normalizing Parameter
Tantalum Powder Wash and Scrubber	20,433	4,900	Tantalum Washed
Consolidation and Casting Contact Cooling	0	0	Columbium or Tantalum Cast or Consolidated

and a second second

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1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 - 1997 -

TABLE IX-2

BPT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(a) Concentrate Digestion Wet Air Pollution Control BPT

Pollutant or		Maximum for	Maximum for
Pollutant Property	7	Any One Day	Monthly Average
Mahada	TTo i har and i har		
		of concentrate die	
English Units	s = lbs/million	n lbs of concentrat	te digested
Antimony		17,850	7.960
Arsenic		13.000	5.784
Cadmium		2.114	0.933
Chromium		2.736	1.119
Copper		11.820	6.219
*Lead		2.612	1.244
Nickel		11.940	7.898
Selenium		7.649	3.420
Thallium		12.750	5.659
*Zinc		9.080	3.794
*Ammonia (as N)		829.000	364.500
*Fluoride		217.700	124.400
*TSS		255.000	121.300
*pH	Within the ra	ange of 7.0 to 10.0	

(b) Solvent Extraction Raffinate BPT

Pollutant or Pollutant Pro	operty	Maximum for Any One Day	Maximum for Monthly Average
iorracane ric	sperey	mig one bag	monenty mverage
Mc	etric Units - mg/kg of	annantrata di	roctod
	Units - lbs/million 1		
Engrish	OIIICS = IDS/MIIIIOIII	the of concentration	te digested
Antimony		26.280	11.720
Arsenic	:	19.130	8.524
Cadmium		3.113	1.373
Chromium		4.028	1.648
Copper		17.400	9.155
*Lead		3.845	1.831
Nickel	2 - C	11.580	11.630
Selenium		11.260	5.035
Thallium		18.770	8.331
*Zinc		13.370	5.585
*Ammonia (as	N)	1,221.000	536.500
*Fluoride		320.400	183.100
*TSS	•	375.400	178.500
*pH	Within the ran	nge of 7.0 to 10	.0 at all times

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TABLE IX-2 (Continued)

BPT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(c) <u>Solvent Extraction Wet Air Pollution</u> <u>Control</u> BPT

Pollutant or Pollutant Property		Maximum for nthly Average
	/kg of concentrate diges lion lbs of concentrate of	
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride *TSS *pH Within t	7.049 5.133 0.835 1.081 4.666 1.032 4.716 3.021 5.035 3.586 327.400 85.960 100.700 he range of 7.0 to 10.0 a	3.144 2.284 0.368 0.422 2.456 0.491 3.119 1.351 2.235 1.498 143.900 49.120 47.890 at all times

(d) Precipitation and Filtration BPT

Pollutant or	•		Maximum for	Maximum for
Pollutant Pr	operty		Any One Day	Monthly Average
	letric Units - n			
English	n Units - 1bs/mi	lllion ll	os or concentra	te digested
Antimony			39.290	17.530
Arsenic		1	28.610	12.730
Cadmium			4.654	2.053
Chromium			6.023	2.464
Copper			26.010	13.690
*Lead			5.570	2.733
Nickel			· 26.010	13.690
Selenium			16.840	7.529
Thallium			28.060	12.460
*Zinc			19.990	8.350
*Ammonia (as	; N)		1,825.000	802.200
*Fluoride			479.100	273.800
*TSS			561.300	267.000
*pH	Within	the rang	ge or 7.0 to 10	.0 at all times

BPT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(e) <u>Precipitation and Filtration Wet Air Pollution Control</u> BPT

Pollutant or Pollutant Property	nits - ma/ka	Maximum for Any One Day g of concentrate d	Maximum for Monthly Average
English Units	- lbs/millic	on lbs of concentrate d	ate digested
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride *TSS *pH	Within the	182.300 132.700 21.590 27.950 120.700 26.680 122.000 78.120 130.200 92.730 8,466.000 2,223.000 2,604.000 range of 7.0 to 10	81.300 59.070 9.527 11.430 63.510 12.700 80.660 34.930 57.800 38.740 3,722.000 1,270.000 1,239.000 0.0 at all times

(f) <u>Tantalum</u> <u>Salt</u> <u>Drying</u> BPT

Pollutant or	Maximum		Maximum for
Pollutant Property	Any One		Monthly Average
Metric Units - mg/kg of	tantalum	n salt o	dried
English Units - lbs/million :	lbs of ta	antalum	salt dried

Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride *TSS				173.800 126.500 29.580 26.640 115.000 25.430 116.200 74.470 124.100 88.390 8,070.000 2,119.000 2.482.000	77.490 56.300 9.081 10.900 60.540 12.110 76.890 33.300 55.090 36.930 3,548.000 1,211.000
*TSS *pH	Within	the	range	2,482.000	1,181.000

BPT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(g) Oxides Calcining Wet Air Pollution Control BPT

Pollutant or Pollutant Property		aximum for chly Average
Metric Units - mg/kg of colu English Units - lbs/million lbs of		
Antimony	110.300	49.180
Arsenic	89.300	35.730
Cadmium	13.060	5.763
Chromium	16.910	6.916
Copper	73.000	38.420
*Lead	16.140	7.685
Nickel	73,770	48.800
Selenium	47.260	21.130
Thallium	78.770	48.800
*Zinc	56.100	23.440

*Ammonia (as N)	5,122.000	2,252.000
*Fluoride	1,345.000	768.500
*TSS	1,576.000	749.200
*pH	Within the range of 7.0 to 10.0 at	all times
		1

(h) Reduction of Tantalum Salt to Metal BPT

Pollutant	or	Maximum	for	Maximum for
Pollutant	Property	Any One	Day	Monthly Average

Metric Units - mg/kg of tantalum salt reduced English Units - lbs/million lbs of tantalum salt reduced

										r
Antimony	*				476.	.600			21	L2.600
Arsenic					347.	.100			15	54.140
Cadmium					56.	460			2	24.910
Chromium					73.	.070				29.89
Copper					315.	500			16	56.100
*Lead					69.	.750			3	33.220
Nickel					315.	.500			16	56.100
Selenium					69.	.750			5	33,210
Thallium					318.	.900			21	L0.900
*Zinc					242.	.500			10	01.300
*Ammonia (as	N)				22,140.	.000			9,73	32,000
*Fluoride					5,813.	.000			3,32	22.000
*TSS					6,809.	.000			3,23	39.000
*pH		Within	the	range	of 7.0	to l	0.0	at	all	times

BPT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

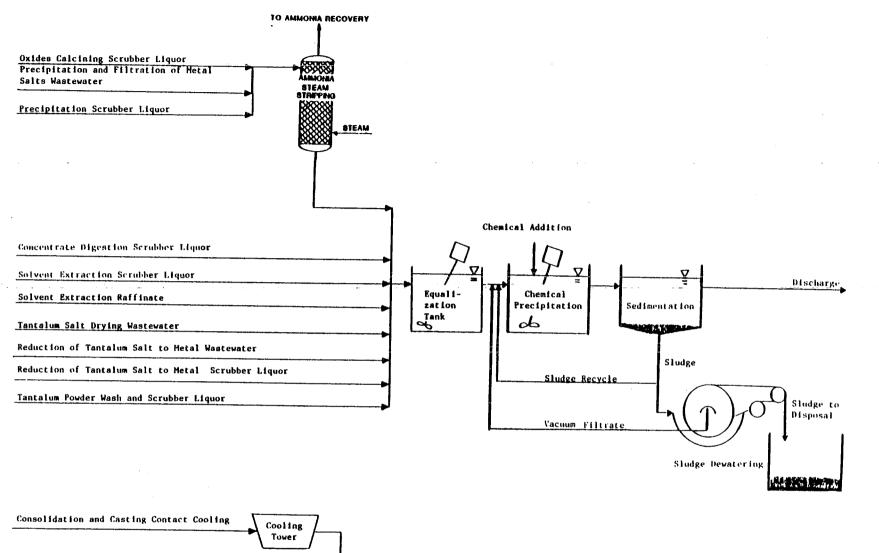
(i) <u>Reduction of Tantalum Salt to Metal Wet Air Pollution</u> <u>Control</u> BPT

	· · · ·		
Pollutant or		Maximum for	Maximum for
Pollutant Propert	y	Any One Day	
-	-		nonenity Average
Metric	Units - mg/k	g of tantalum sal	treduced
English Unit	s - lbs/mill	ion lbs of tantal	um salt reduced
- -	,		am bart reduced
Antimony	1	5.86	3 2.615
Arsenic		4.27	0 1.900
Cadmium	·	0.69	
Chromium		0.89	
Copper		3.88	
*Lead		0.85	
Nickel		3.95	
Selenium	;	2.51	
Thallium		4.18	
*Zinc		2.98	
*Ammonia (as N)		272.40	
*Fluoride	14	71.51	,
*TSS		83.770	
*pH	Within the		10.0 at all times
		· · · ·	
(j) <u>Tantalum</u> Powe	der Wash BP1	•	
	·		
Pollutant or			
Pollutant Property	•	Maximum for	
ioriucant Floperty	Ŷ	Any One Day	Monthly Average
Motria	Inita _ malle		
English Unite	= lbc/millio	of tantalum powe	ler washed
Engrish onics		on lbs of tantalum	l powder washed
Antimony		58.640	
Arsenic		42.710	
Cadmium		6.947	
Chromium		8.991	
Copper			
*Lead		38.820	
Nickel		8.852	
Selenium		39.230	
Thallium		25.130	
*Zinc		41.890	
		29.830	
*Ammonia (as N)		2,724.000	
*Fluoride	I.	715.200	408.700
*TSS		837.800	398.500
*pH			
F	within the	range of 7.0 to	10.0 at all times

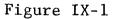
BPT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(k) Consolidation and Casting Contact Cooling BPT

Pollutant or Pollutant Proper	cty	Maximum for Any One Day	Maximum for Monthly Average
Metric Units - r	ng/kg of columbium	or tantalum cas	st or consolidated
English Units -	lbs/million lbs o consolidated	f columbium or t	cantalum cast or
Antimony		0.000	0.000
Arsenic		0.000	0.000
Cadmium		0.000	0.000
Chromium		0.000	0.000
Copper		0.000	0.000
*Lead		0.000	0.000
Nickel		0.000	0.000
Selenium		0.000	0.000
Thallium		0.000	0.000
*Zinc		0.000	0.000
*Ammonia (as N)		0.000	0.000
*Fluoride		0.000	0.000
*TSS		0.000	0.000
*155 *pH	Within the ra		0.0 at all times







BPT TREATMENT SCHEME PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

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SECTION X

BEST AVAILABLE TECHNOLOGY ECONOMICALLY ACHIEVABLE

These effluent limitations are based on the best control and treatment technology used by a specific point source within the industrial category or subcategory, or by another category where is readily transferable. Emphasis is placed on additional it treatment techniques applied at the end of the treatment systems currently used, as well as reduction of the amount of water used discharged, process control, and treatment and technology optimization.

The factors considered in assessing best available technology economically achievable (BAT) include the age of equipment and facilities involved, the process used, process changes, nonwater quality environmental impacts (including energy requirements), and the costs of application of such technology. At a minimum BAT technology represents the best available technology at plants of various ages, sizes, processes, or other characteristics. BAT may be transferred from a different subcategory or category. BAT may include feasible process changes or internal controls, even when not in common industry practice.

The statutory assessment of BAT considers costs, but does not require a balancing of costs against effluent reduction benefits However, in assessing BAT, the Agency has given substantial weight to the economic achievability of the selected technology.

TECHNICAL APPROACH TO BAT

The Agency reviewed a wide range of technology options and evaluated the available possibilities to ensure that the most effective and beneficial technologies were used as the basis of BAT. To accomplish this, the Agency elected to examine three technology options which could be applied to the primary columbium-tantalum subcategory as treatment options for the basis of BAT effluent limitations.

For the development of BAT effluent limitations, mass loadings were calculated for each wastewater source or subdivision in the subcategory using the same technical approach as described in Section IX for BPT limitations development. The differences in mass loadings for BPT and BAT are due to increased treatment the effectiveness achievable with the more sophisticated BAT technology and reductions in the effluent treatment flows allocated to various waste streams.

The treatment technologies considered for BAT are presented below:

Option A (Figure X-1, page 4494) is based on

o Preliminary treatment with ammonia steam stripping

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o Chemical precipitation and sedimentation

Option B (Figure X-2, page 4495) is based on

- o Preliminary treatment with ammonia steam stripping
- o Chemical precipitation and sedimentation
- In-process flow reduction 0

Option C (Figure X-3, page 4496) is based on

- Preliminary treatment with ammonia steam stripping 0
- o Chemical precipitation and sedimentation
- 0 In-process flow reduction
- Multimedia filtration 0

The three options examined for BAT are discussed in greater detail below. The first option considered is the same as the BPT treatment which was presented in the previous section. The last two options represent substantial progress toward the prevention of polluting the environment above and beyond the progress achievable by BPT.

OPTION A

Option A for the primary columbium-tantalum subcategory is equivalent to the control and treatment technologies which were selected for BPT in Section IX. The BPT end-of-pipe treatment scheme includes lime precipitation, sedimentation, with ammonia steam stripping preliminary treatment (see Figure X-1). The discharge rates for Option A are equal to the discharge rates The allocated to each stream as a BPT discharge flow.

OPTION B

Option B for the primary columbium-tantalum subcategory achieves lower pollutant discharge by building upon the Option A end-ofpipe treatment technology, which consists of ammonia steam stripping, lime precipitation, and sedimentation. Flow reduction measures are added to Option A treatment (see Figure X-2). These flow reduction measures result in the concentration of pollutants in scrubber liquor effluents. Treatment of a more concentrated effluent allows achievement of a greater net pollutant removal and introduces the possible economic benefits associated with treating a lower volume of wastewater.

Methods used in Option B to reduce process wastewater generation or discharge rates are presented below:

Recycle of Water Used in Wet Air Pollution Control

There are five wastewater sources associated with wet air pollution control which are regulated under these effluent limitations:

- o Concentrate digestion scrubber,
- o Solvent extraction scrubber,
- o Precipitation and filtration scrubber,
- o Oxides calcining drying scrubber, and
- o Reduction of salt to metal scrubber.

Table X-1 (page 4484) presents the number of plants reporting wastewater use with these sources, the number of plants practicing recycle of scrubber liquor, and the range of recycle values being used. Although some plants report total recycle of their scrubber water, some blowdown or periodic cleaning is likely to be needed to prevent the build-up of dissolved and suspended solids since the water picks up particulates and fumes from the air.

OPTION C

Option C for the primary columbium-tantalum subcategory consists of all control and treatment requirements of Option B (ammonia steam stripping, in-process flow reduction, lime precipitation, and sedimentation) plus multimedia filtration technology added at the end of the Option B treatment scheme (see Figure X-3). Multimedia filtration is used to remove suspended solids, including precipitates of toxic metals, beyond the concentrations attainable by gravity sedimentation. The filter suggested is of the gravity, mixed media type, although other filters, such as rapid sand filters or pressure filters, would perform as well.

As one means of evaluating each technology option, EPA developed estimates of the pollutant removal estimates and the compliance costs associated with each option. The methodologies are described below.

POLLUTANT REMOVAL ESTIMATES

A complete description of the methodology used to calculate the estimated pollutant reduction achieved by the application of the various treatment options is presented in Section X of the General Development Document. The pollutant removal estimates have been revised from proposal based on comments and new data. However, the methodology for calculating pollutant removals was not changed. The data used for estimating pollutant removals are the same as those used to revise the compliance costs.

Sampling data collected during the field sampling program were used to characterize the major waste streams considered for regulation. At each sampled facility, the sampling data were production normalized for each unit operation (i.e., mass of pollutant generated per mass of product manufactured). This value, referred to as the raw waste, was used to estimate the mass of toxic pollutants generated within the columbium-tantalum subcategory. By multiplying the total subcategory production for a unit operation by the corresponding raw waste value, the mass of pollutant generated for that unit operation was estimated. The volume of wastewater discharged after the application of each treatment option was estimated for each operation at each plant by comparing the actual discharge to the regulatory flow. The smaller of the two values was selected and summed with the other plant flows. The mass of pollutant discharged was then estimated by multiplying the achievable concentration values attainable by the option (mg/l) by the estimated volume of process wastewater discharged by the subcategory. The mass of pollutant removed is simply the difference between the estimated mass of pollutant generated within the subcategory and the mass of pollutant discharged after application of the treatment option.

The pollutant removal estimates for the direct dischargers in the primary columbium-tantalum subcategory are presented in Table X-2 (page 4485).

COMPLIANCE COST

Compliance costs presented at proposal were estimated using cost curves, which related the total costs associated with installation and operation of wastewater treatment technologies to plant process wastewater discharge. EPA applied these curves on a per plant basis, a plant's costs -- both capital, and operating and maintenance -- being determined by what treatment it has in place and by its individual process wastewater discharge (from dcp). The final step was to annualize the capital costs, and to sum the annualized capital costs, and the operating and maintenance costs, yielding the cost of compliance for the subcategory.

Since proposal, the cost estimation methodology has been changed as discussed in Section VIII of Vol. I. A design model and plant specific information were used to size a wastewater treatment system for each discharging facility. After completion of the design, capital and annual costs were estimated for each unit of the wastewater treatment system. Capital costs rely on vendor quotes, while annual costs were developed from the literature. The revised compliance costs for direct dischargers are presented in Table VIII-1 (page 4452).

BAT OPTION SELECTION

For BAT, EPA is promulgating mass limitations based on lime precipitation and sedimentation with ammonia steam stripping with additional reduction in pollutant discharge achieved through inprocess wastewater flow reduction and the use of filtration as an effluent polishing step. The end-of-pipe and pretreatment technology basis for BAT limitations being promulgated is the same as that for the proposed limitations. Ammonia steam stripping is currently demonstrated at two columbium-tantalum facilities. Filtration is not demonstrated within this subcategory, but is transferred from six nonferrous metals subcategories where it is demonstrated in 23 plants. With the limits for fluoride, the treatment performance exception of

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - X

concentrations upon which the mass limitations are based are equal to the values used to calculate the proposed mass limitations. The mass limitations for fluoride have been revised for the reasons discussed in Section IX - BPT Option Selection.

Revision of the proposed flow allowances is consistent with the changes made for the promulgated BPT limitations. The differences between the promulgated BPT and BAT flow allowances are due to flow reduction of scrubber liquors at BAT. The BAT flow allowances are discussed in detail below.

EPA estimates that application of BAT will remove 61,400 kg/yr of toxic metals and 1,694,000 kg/yr of nonconventional pollutants over raw discharge rates. BAT will result in the estimated 283 kg/yr of toxic pollutants and 1,980 kg/yr of removal of nonconventional pollutants over the estimated BPT discharge. The final BAT effluent mass limitations will remove 57 kg/yr of toxic metals over the intermediate BAT option considered, which lacks filtration. Both options are economically achievable. The Agency believes that the incremental removal justifies selection filtration as part of BAT model technology. of The estimated capital investment cost of BAT is \$0.83 million (March, 1982 dollars) and the estimated annual cost is \$1.2 million.

WASTEWATER DISCHARGE RATES

A BAT discharge rate was calculated for each subdivision based upon the flows of the existing plants, as determined from analysis of dcp. The discharge rate is used with the achievable treatment concentration to determine BAT effluent limitations. Since the discharge rate may be different for each wastewater source, separate production normalized discharge rates for each the ll wastewater sources were determined and are summarized of Table X-3 (page 4486). The discharge rates are generally in normalized on a production basis by relating the amount of wastewater generated to the mass of the intermediate product which is produced by the process associated with the waste stream in question. These production normalizing parameters (PNP) are also listed in Table X-3.

The BAT wastewater discharge rate equals the BPT wastewater discharge rate for seven of the ll waste streams in the primary columbium-tantalum subcategory. Based on the available data, the Agency did not find that further flow reduction would be feasible for these wastewater sources. The rationale for determining these regulatory flows is presented in Section IX. Wastewater streams for which BAT discharge rates differ from BPT are discussed below.

CONCENTRATE DIGESTION WET AIR POLLUTION CONTROL

The proposed BAT wastewater discharge rate for concentrate digestion wet air pollution control was 5,156 l/kkg (1,237 gal/ton) of columbium-tantalum salt produced from digestion. The BAT discharge rate was based on 90 percent recycle of the average

water use of two plants. A third plant reported insufficient dcp information to calculate a discharge rate.

The proposed BAT discharge rate has been revised based on the considerations presented in Section IX of this document. The promulgated BAT discharge rate is based on 90 percent recycle of the BPT discharge rate, or 622 l/kkg (149 gal/ton) of raw material digested.

SOLVENT EXTRACTION WET AIR POLLUTION CONTROL

The proposed BAT wastewater discharge rate for solvent extraction wet air pollution control was 430 l/kkg (103 gal/ton) of columbium or tantalum salt extracted. The BAT discharge rate was based on 90 percent recycle of the water use at one of the two plants which generate this waste stream.

The proposed BAT discharge rate has been revised based on the considerations presented in Section IX of this document. The promulgated discharge rate is 246 l/kkg (60 gal/ton) of raw material digested, and it is based on 90 percent recycle of the BPT discharge rate. One plant uses the same scrubber for both solvent extraction and concentrate digestion wet air pollution control. Both discharge allowances apply to this plant since water use in the scrubber is probably increased due to the added volume of air scrubbed.

PRECIPITATION AND FILTRATION WET AIR POLLUTION CONTROL

A discharge allowance for this waste stream was not proposed. Re-evaluation of the information supplied to the Agency has shown that a discharge allowance is necessary for plants operating scrubbers on precipitation and filtration processes. The promulgated BAT discharge rate is based on 90 percent recycle of the BPT discharge rate, or 6,351 1/kkg (1,523 gal/ton) of concentrate digested.

OXIDES CALCINING WET AIR POLLUTION CONTROL

The proposed BAT wastewater discharge rate for metal salt drying wet air pollution control was 16,479.4 l/kkg (3,961.4 gal/ton) of columbium or tantalum salt dried. The BAT discharge rate was based on 90 percent recycle of the water use at one of these plants. Two plants reported insufficient dcp information to calculate water usage, and the water usage of one plant was extremely high. These plants were not considered in calculating the BAT discharge rate.

As discussed in Section IX, the proposed metal salt drying wet air pollution control allowance has been divided into tantalum salt drying and oxides calcining wet air pollution control. The promulgated BAT discharge rate for oxides calcining wet air pollution control is based on 90 percent recycle of the BPT discharge rate, or 3,842 1/kkg (921 gal/ton) of columbium or tantalum oxide dried.

REDUCTION OF TANTALUM SALT TO METAL WET AIR POLLUTION CONTROL

The BAT promulgation discharge rate is equal to the BPT presented in Section IX of this document, which is 2,043 rate l/kkq gal/ton) of tantalum salt reduced. The proposal discharge (490 was based on 90 percent recycle; , recycle is rate not appropriate for the two plants that operate this scrubber. Both plants utilize rotoclone scrubbers which are much different than once through scrubbers such as packed towers or venturi Although there may be a discharge from a rotoclone scrubbers. scrubber, the scrubber acts as a sparge tank by drawing the gas stream through a body of water in a tank. Water droplets and mist created due to turbulence are captured and routed back to the tank.

REGULATED POLLUTANT PARAMETERS

In implementing the Clean Water Act, the Agency placed particular emphasis on the toxic pollutants. The raw wastewater concentrations from individual operations and the subcategory as a whole were examined to select certain pollutants and pollutant parameters for limitation. This examination and evaluation was presented in Section VI. The Agency, however, has chosen not to regulate all 21 toxic pollutants selected in this analysis.

The columbium-tantalum subcategory generates an estimated 80,000 kg/yr of toxic pollutants, of which only 170 kg/yr are toxic organic pollutants. The Agency believes that the toxic organic pollutants in the columbium-tantalum subcategory are present only in trace (deminimus quantities) and are neither causing nor likely to cause toxic effects. However, it is possible toxic organic pollutants may be present in larger concentrations at an individual plant than the Agency sampling data indicate. Therefore, the permitting or control authority should check for the presence of toxic organic pollutants on a case-by-case basis and determine if they require treatment. The following toxic organic pollutants are excluded from regulation:

- 4. benzene
- 6. carbon tetrachloride
- 7. chlorobenzene
- 8. 1,2,4-trichlorobenzene
- 10. 1,2-dichloroethane
- 30. 1,2-trans-dichloroethylene
- 38. ethylbenzene
- 51. chlorodibromomethane
- 85. tetrachloroethylene
- 87. trichloroethylene

The cost associated with analysis for toxic metal pollutants has prompted EPA to develop an alternative method for regulating and monitoring toxic pollutant discharges from the nonferrous metals manufacturing category. Rather than developing specific effluent

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mass limitations and standards for each of the toxic metals found in treatable concentrations in the raw wastewater from a given subcategory, the Agency is promulgating effluent mass limitations only for those pollutants generated in the greatest quantities as shown by the pollutant removal estimate analysis. The pollutants selected for specific limitation are listed below:

122. lead
128. zinc
ammonia (as N)
fluoride

By establishing limitations and standards for certain toxic metal pollutants, dischargers will attain the same degree of control over toxic metal pollutants as they would have been required to achieve had all the toxic metal pollutants been directly limited.

This approach is justified technically since the treatable concentrations used for lime precipitation and sedimentation technology are based on optimized treatment for concomitant multiple metals removal. Thus, even though metals have somewhat different theoretical solubilities, they will be removed at very nearly the same rate in a lime precipitation and sedimentation treatment system operated for multiple metals removal. Filtration as part of the technology basis is likewise justified because this technology removes metals non-preferentially.

The toxic metal pollutants selected for specific limitation in the columbium-tantalum subcategory to control the discharges of toxic metal pollutants are lead and zinc. Ammonia is also selected for limitation since the methods used to control lead and zinc are not effective in the control of ammonia. The following toxic pollutants are excluded from limitation on the basis that they are effectively controlled by the limitations developed for lead and zinc:

114. antimony 115. arsenic 116. asbestos 118. cadmium 119. chromium (Total) 120. copper 124. nickel 125. selenium 127. thallium

EFFLUENT LIMITATIONS

The concentrations achievable by application of BAT are discussed in Section VII of Vol. I and summarized there in Table VII-21 (page 248). The treatment effectiveness of both one day maximum and monthly average values are multiplied by the BAT normalized discharge flows summarized in Table X-3 (page 4486) to calculate the mass of pollutants allowed to be discharged per mass of

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product. The results of these calculations in milligrams of pollutant per kilogram of product represent the BAT effluent limitations and are presented in Table X-4 (page 4488) for each waste stream.

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TABLE X-1

CURRENT RECYCLE PRACTICE WITHIN THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

Process	Number of Plants With Wastewater	No. of Plants Practicing <u>Recycle</u>	Range of Recycle Values (%)
Concentrate Digestion Wet Air Pollution Control	3	1	0 - 86
Solvent Extraction We Air Pollution COntrol		1	0 - 86
Precipitation Wet Air Pollution Control	2	0	:
Oxides Calcining Wet Air Pollution Control	4	2	89 - 100
Reduction of Tantalum to Metal Wet Air Poll Control		0	• • •

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Table X-2

POLLUTANT REMOVAL ESTIMATES FOR PRIMARY COLUMBIUM-TANTALUM DIRECT DISCHARGERS

POLLUTANT	TOTAL RAW WASTE (kg/yr)	OPTION A DISCHARGED (kg/yr)	OPTION A REMOVED (kg/yr)	OPTION B DISCHARGED (kg/yr)	OPTION B REMOVED (kg/yr)	OPTION C DISCHARGED (kg/yr)	OPTION C REMOVED (kg/yr)
Ant imony	396.1	106.7	289.4	74.2	321.9	49.8	346.3
Arsenic	564.3	77.7	486.6	54.1	510.2	36.0	528.3
Cadmium	426.3	12.0	414.3	8.4	418.0	5,2	421.2
Chromium	20,888.1	12.8	20,875.3	8.9	20,879.2	7.4	20,880.6
Copper	3,503.7	88.4	3,415.3	61.5	3,442.2	41.3	3,462.3
Lead	18,149.9	18.3	18,131.6	12.7	18,137.2	8.5	18,141.4
Nickel	279.4	112.8	166.7	78.5	201.0	23.3	256.1
21nc	17,364.4	50.3	17,314.1	35.0	17,329.4	24.4	17,340.0
TOTAL TOXIC METALS	61,572.3	478 .9	61,093.4	333.2	61,239.1	196.0	61,376.3
Aluminum	7,435.6	341.3	7,094.2	237.5	7,198.1	158.0	7 177 6
Amnonia	945,842.4	4,684.8	941,157.6	3,372.8	942,469.6	3,372.8	7,277.6
Fluoride	974,798.6	2,209.5	972,589.0	1,537.3	973,261.3		942,469.6
Iron	352,794.3	62.5	352,731.8	43.5	352,750.8	1,537.3 29.7	973,261.3 352,764.6
TUTAL NONCONVENTIONALS	2,280,870.8	7,298.1	2,273,572.7	5,191.0	2,275,679.8	5,097.7	2,275,773.1
TSS	1,694,063.2	1,828.6	1,692,234.6	1,272.2	1,692,790.9	275.7	1,693,787.5
TOTAL CONVENTIONALS	1,694,063.2	1,828.6	1,692,234.6	1,272.2	1,692,790.9	275.7	1,693,787.5
TUTAL POLIJITANTS	4,036,506.3	9,605.6	4,026,900.7	•6,796.5	4,029,709.8	5,569.4	4,030,936.9
FLOW (1/yr)	· .	152,380,000		106,020,000		106,020,000	

NOTE: TOTAL TOXIC METALS = Antimony + Arsenic + Cadmium + Chromium + Copper + Lead + Nickel + Zinc TOTAL NONCONVENTIONALS = Aluminum + Ammonia + Fluoride + Iron TUTAL CONVENTIONALS = TSS TUTAL POLLUTANTS = Total Toxic Metals + Total Nonconventionals + Total Conventionals

OPTION A = Lime Preciptiation, Sedimentation, and Ammonia Steam Stripping. OPTION B = Option A, plus In-Process Flow Reduction.

OPTION C = Option B, plus Multimedia Filtration.

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Table X-3

BAT WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

		rmalized	
Wastewater Stream	1/kkg	gal/ton	Production Normalizing Parameter
Concentrate Digestion Wet Air Pollution Control	622	149	Concentrate Digested
Solvent Extraction Raffi- nate	9,155	2,195	Concentrate Digested
Solvent Extraction Wet Air Pollution Control	246	60	Concentrate Digested
Precipitation and Filtra- tion Wastewater	13,689	3,283	Concentrate Digested
Precipitation and Filtra- tion Wet Air Pollution Control	6,351	1,523	Concentrate Digested
Tantalum Salt Drying	60,542	14,518	Tantalum Salt Dried
Oxides Calcining Wet Air Pollution Control	3,842	921	Columbium or Tantalum Oxide Calcined
Reduction of Tantalum Salt to Metal	166,071	39,825	Tantalum Salt Reduced
Reduction of Tantalum Salt Metal Wet Air Pollution Control	2,043	490	Tantalum Salt Reduced

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Table X-3 (Continued)

BAT WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

		malized ge Rate	
Wastewater Stream	1/kkg	gal/ton	Production Normalizing Parameter
Tantalum Powder Wash and Scrubber	20,433	4,900	Tantalum Washed
Consolidation and Casting Contact Cooling	0	0	Columbium or Tantalum Cast or Consolidated

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TABLE X-4

BAT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(a) Concentrate Digestion Wet Air Pollution Control BAT

Pollutant or Pollutant Property		Maximum for onthly Average
Metric Units - mg/kg English Units - lbs/million		
Antimony	1.200	0.535
Arsenic	0.865	0.386
Cadmium	0.124	0.050
Chromium	0.230	0.093
Copper	0.796	0.379
*Lead	0.174	0.081
Nickel	0.342	0.230
Selenium	0.510	0.230
Thallium	0.871	0.379
*Zinc	0.635	0.261
*Ammonia (as N)	82.910	36.450
*Fluoride	21.770	12.440
(b) Solvent Extraction Raffinate	e BAT	1

Pollutant	or	Maximum	for	Maximum for
Pollutant	Property	Any One	Day	Monthly Average

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested

Antimony	17.670	7.873
Arsenic	12,730	5,676
Cadmium	1.831	0.732
Chromium	3.387	1.373
Copper	11.720	5.585
*Lead	2.563	1.190
Nickel	5.035	0.000
Selenium	7.507	3,387
Thallium	7.507	3.387
*Zinc	9.338	3.845
*Ammonia (as N)	1,221.000	536.500
*Fluoride	320.400	183.100
	,	

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TABLE X-4 (Continued)

BAT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(c) Solvent Extraction Wet Air Pollution Control BAT

Pollutant or		Maximum for	Maximum for
Pollutant Property		Any One Day	Monthly Average
Metric Units English Units - 1bs		of concentrate d lbs of concentr	
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride (d) Precipitation and F	hiltration	0.475 0.342 0.049 0.091 0.315 0.069 0.135 0.202 0.344 0.251 32.790 8.610	$\begin{array}{c} 0.212\\ 0.153\\ 0.020\\ 0.167\\ 0.000\\ 0.032\\ 0.091\\ 0.091\\ 0.150\\ 0.103\\ 14.420\\ 4.920\\ \end{array}$
Pollutant or		Maximum for	Maximum for
Pollutant Property		Any One Day	Monthly Average
Metric Units	- mg/kg c	f concentrate d	igested
English Units - lbs	/million	lbs of concentr	ate digested
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride		26.430 19.030 2.738 5.065 17.520 3.833 7.529 11.230 19.170 13.960 1,825.000 479.100	11.770 8.487 1.095 2.053 8.350 1.780 5.065 5.065 8.350 5.750 802.200 273.800

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(e) <u>Precipitation</u> and <u>Filtration</u>	Wet Air Pollutic	on <u>Control</u> BAT
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg o	f concentrate dig	gested
English Units - lbs/million	lbs of concentrat	ce digested
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride (f) <u>Tantalum Salt Drying</u> BAT	12.260 8.828 1.270 2.350 8.129 1.778 3.493 5.208 8.891 6.478 846.600 222.300	5.462 3.938 0.508 0.953 3.874 0.826 2.350 2.350 2.350 3.874 2.668 372.200 127.000
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg o	f tantalum salt	dried
English Units - lbs/million	lbs of tantalum	salt dried
Antimony	116.800	52.070
Arsenic	84.150	37.540
Cadmium	12.110	4.843
Chromium	22.400	9.081
Copper	77.490	36.930
*Lead	16.950	7.870
Nickel	33.300	22.400
Selenium	49.640	22.400
Thallium	84.760	36.930
Lead	16.950	7.871
Zinc	61.750	25.430
Ammonia (as N)	8,070.000	3,548.000
Fluoride	2,119.000	1,211.000

(g)	Oxides	Calcining	<u>Wet</u>	<u>Air</u>	Pollution	Control	BAT

Pollutant of Pollutant H		Maximum for Any One Day	Maximum for Monthly Average
Metri	ic Units - mg/kg o	f columbium-tantalum	oxide dried
		lbs of columbium-tant	
Antimony		7.415	3.304
Arsenic		5.340	2.382
Cadmium	·	0.768	0.307
Chromium		1.422	0.576
Copper		4.918	2.344
*Lead		1.076	0.500
Nickel	*	2.113	1.422
Selenium		3.150	1.422
Thallium	• •	5.379	2.344
*Zinc	· · · · ·	3.919	1.614
*Ammonia (a	N)	512.200	225.200
*Fluoride	~~ ~~ ,	134.500	76.840
(h) <u>Reduct</u>	<u>ion of Tantalum S</u>	alt to Metal BAT	
Pollutant c	Dr	<u>alt to Metal</u> BAT Maximum for Any One Day	Maximum for Monthly Average
Pollutant c Pollutant H	or Property Metric Units - mg/	Maximum for	Monthly Average
Pollutant c Pollutant f Englis	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum s	Monthly Average educed salt reduced
Pollutant o Pollutant I Englis Antimony	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum s 320.500	Monthly Average educed salt reduced 142.800
Pollutant c Pollutant H Englis Antimony Arsenic	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum 320.500 230.800	Monthly Average educed salt reduced 142.800 103.000
Pollutant o Pollutant H Englis Antimony Arsenic Cadmium	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum s 320.500 230.800 33.210	Monthly Average educed salt reduced 142.800 103.000 13.290
Pollutant o Pollutant H Englis Antimony Arsenic Cadmium Chromium	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum s 320.500 230.800 33.210 61.450	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910
Pollutant o Pollutant I Englis Antimony Arsenic Cadmium Chromium Copper	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum 320.500 230.800 33.210 61.450 212.600	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910 101.300
Pollutant o Pollutant I Englis Antimony Arsenic Cadmium Chromium Copper *Lead	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum 320.500 230.800 33.210 61.450 212.600 46.500	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910 101.300 21.590
Pollutant o Pollutant I Englis Antimony Arsenic Cadmium Chromium Copper *Lead Nickel	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt ro lion lbs of tantalum 320.500 230.800 33.210 61.450 212.600 46.500 91.340	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910 101.300 21.590 61.450
Pollutant o Pollutant I Englis Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt re lion lbs of tantalum 320.500 230.800 33.210 61.450 212.600 46.500 91.340 136.200	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910 101.300 21.590 61.450 61.450
Pollutant o Pollutant I Englis Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium	or Property Metric Units - mg/	Maximum for Any One Day kg of tantalum salt re lion lbs of tantalum s 320.500 230.800 33.210 61.450 212.600 46.500 91.340 136.200 232.500	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910 101.300 21.590 61.450 61.450 101.300
Pollutant of Pollutant I Englis Antimony Arsenic Cadmium Chromium Chromium Copper *Lead Nickel Selenium Thallium	or Property Metric Units - mg/ Sh Units - lbs/mil	Maximum for Any One Day kg of tantalum salt re lion lbs of tantalum s 320.500 230.800 33.210 61.450 212.600 46.500 91.340 136.200 232.500 169.400	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910 101.300 21.590 61.450 61.450 101.300 69.750
Pollutant o Pollutant I Englis Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium	or Property Metric Units - mg/ Sh Units - lbs/mil	Maximum for Any One Day kg of tantalum salt re lion lbs of tantalum s 320.500 230.800 33.210 61.450 212.600 46.500 91.340 136.200 232.500	Monthly Average educed salt reduced 142.800 103.000 13.290 24.910 101.300 21.590 61.450 61.450 101.300

BAT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(i) <u>Reduction of Tantalum Salt to Metal Wet Air Pollution</u> Control BAT

Pollutant Pollutant	Property	Maximum Any One	Day	Monthly	um for Average					
Metric Units - mg/kg of tantalum salt reduced English Units - lbs/million lbs of tantalum salt reduced										
Antimony			3.943		1.757					
Arsenic			2.840		1,267					
Cadmium			0.409		0.163					
Chromium			0.756		0.306					
Copper			2.615		1.246					
*Lead	· · · · · · · · · · · · · · · · · · ·		0.572		0.266					
Nickel			1.124	-	0.756					
Selenium			1.675	1	0.756					
Thallium			2.860		1.246					
*Zinc			2.084		0.858					
*Ammonia	(as N)		2.400		119.700					
*Fluoride		7	1.510		40.860					
(j) <u>Tant</u> <u>Pollutant</u>	alum Powder Wash BA	Maximum	for	Maxim	um for					
Pollutant		Any One		Monthly	Average					
FOTTUCANC		-	_	•						
Metric Units - mg/kg of tantalum powder washed English Units - lbs/million lbs of tantalum powder washed										
Antimony	1		9.440		17.570					
Arsenic			8.400		12.670					
Cadmium			4.087		1.635					
Chromium			7.560		3.065					
Copper	F		6.150		12.460					
*Lead	,		5.721		2.656					
Nickel			1.240		7.560					
Selenium	L		6.760		7.560					
Thallium			8.610	,	12.460					
*Zinc			0.840	-	8.582					
*Ammonia	(as N)		4.000		198.000					

715.200

408.700

*Fluoride

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY

SECT - X

TABLE X-4 (Continued)

BAT EFFLUENT LIMITATIONS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(k) Consolidation and Casting Contact Cooling BAT

Pollutant or Pollutant Prope	rty	Maximum for Any One Day	Maximum for Monthly Average
Metric Units - English Units -	mg/kg of columbium lbs/million lbs of consolidated	or tantalum ca columbium or	ast or consolidated tantalum cast or
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride		$\begin{array}{c} 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\end{array}$	$\begin{array}{c} 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\end{array}$

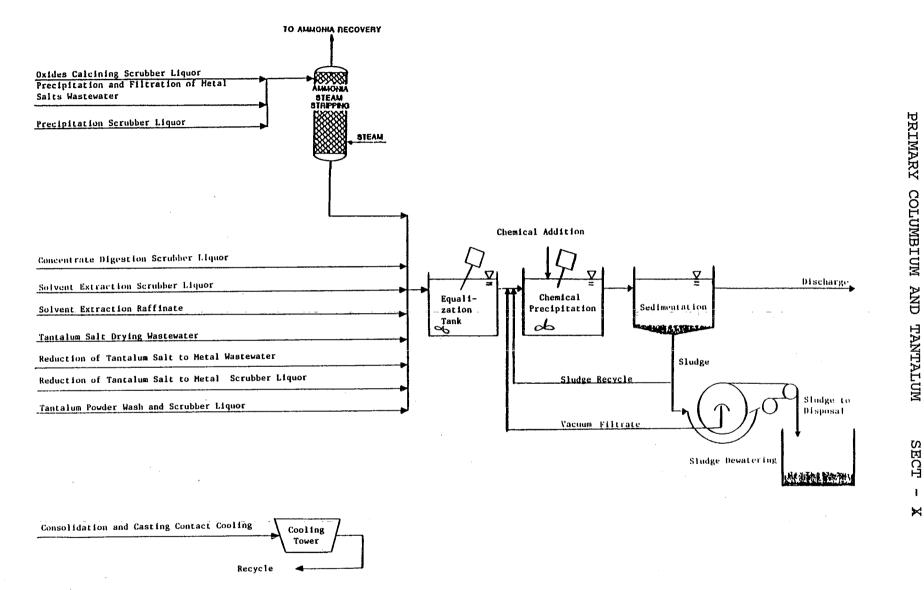
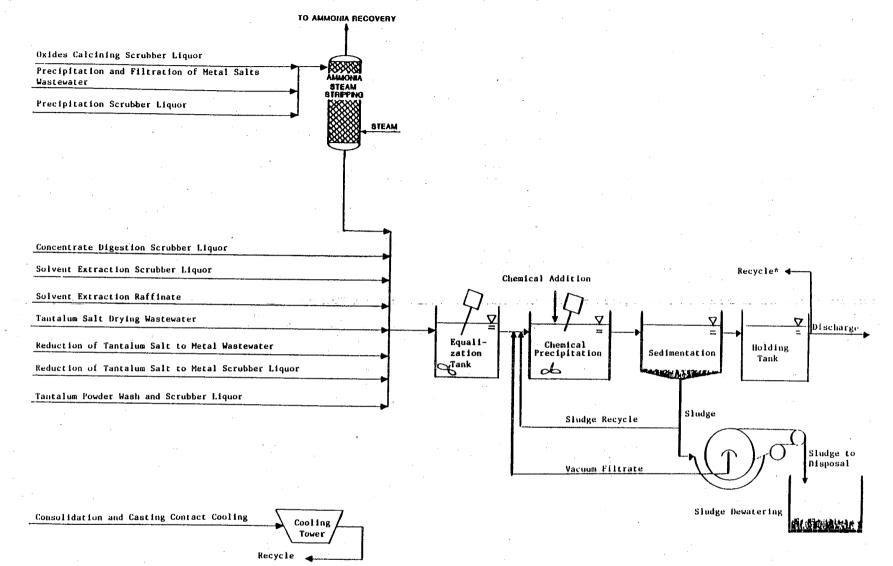


Figure X-1

BAT TREATMENT SCHEME OPTION A PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

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SECT



*90% Recycle of Concentrate Digestion, Solvent Extraction, Precipitation, and Oxides Calcining Scrubber Liquors.

Figure X-2

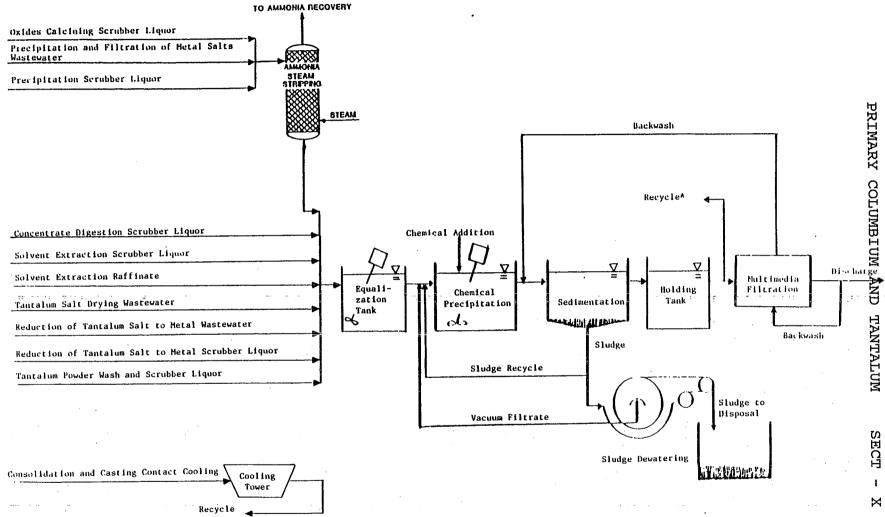
BAT TREATMENT SCHEME OPTION B PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

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SECT -

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PRIMARY COLUMBIUM AND TANTALUM



*90% Recycle of Concentrate Digestion, Solvent Extraction, Precipitation, and Oxides Calcining Scrubber Elquors.

Figure X-3

BAT TREATMENT SCHEME OPTION C PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

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PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - XI

SECTION XI

NEW SOURCE PERFORMANCE STANDARDS

This section describes the control technology for treatment of wastewater from new sources, and presents mass discharge limitations of regulated pollutants for NSPS in the primary columbium-tantalum subcategory, based on the described control technology. New plants have the opportunity to design the best and most efficient production processes and wastewater treatment technologies, without facing the added costs and restrictions encountered in retrofitting an existing plant. Therefore, EPA had considered the best demonstrated process changes, in-plant controls, and end-of-pipe treatment technologies which reduce pollution to the maximum extent feasible.

TECHNICAL APPROACH TO BDT

All of the treatment technology options applicable to a new source were previously considered for the BAT options. For this reason, three options were considered for BDT, all identical to BAT Options A, B, and C, which are discussed in Section X. Briefly, the treatment technologies used for the three options are as follows:

OPTION A

- o Chemical precipitation and sedimentation
- Ammonia steam stripping preliminary treatment of wastewaters containing treatable concentrations of ammonia

OPTION B

- o Chemical precipitation and sedimentation
- o Ammonia steam stripping preliminary treatment of
- wastewaters containing treatable concentrations of ammonia o In-process flow reduction

OPTION C

- o Chemical precipitation and sedimentation
- o Ammonia steam stripping preliminary treatment of
- wastewaters containing treatable concentrations of ammonia o In-process flow reduction
- o Multimedia filtration

Partial or complete recycle and reuse of wastewater is an essential part of the last two options. Recycle and reuse can precede or follow end-of-pipe treatment. A more detailed discussion of the treatment options is presented in Section X.

EPA proposed that the best available demonstrated technology for the primary columbium-tantalum subcategory be equal to BAT (Option C). Review of the subcategory indicated that no new demonstrated technologies that improve on BAT technology exist. Therefore, EPA is promulgating effluent mass limitations for NSPS equivalent to BAT.

Dry scrubbing is not demonstrated for controlling emissions from concentrate digestion, oxides calcining, precipitation and filtration, and salt to metal reduction. The nature of these emissions (acidic fumes, hot particulate matter) technically precludes the use of dry scrubbers. Therefore, EPA is including an allowance for these sources at NSPS equivalent to that promulgated for BAT. The Agency also does not believe that new plants could achieve any additional flow reduction beyond that promulgated for BAT.

Activated alumina (Option D) was considered; however, this technology was rejected because it too was not demonstrated in this category, nor was it clearly transferable to nonferrous wastewater. Activated carbon (Option E) was also considered; however, this technology was eliminated because it is not necessary, since toxic organic pollutants are not selected for limitation in this subcategory.

Reverse osmosis (Option F) was considered for the purpose of achieving zero discharge of process wastewater; however, the Agency ultimately rejected this technology because it was determined that its performance for this specific purpose was not adequately demonstrated in this category nor was it clearly transferable from another category.

REGULATED POLLUTANT PARAMETERS

The Agency has no reason to believe that the pollutants that will be found in treatable concentrations in processes within new sources will be any different than with existing sources. Accordingly, pollutants and pollutant parameters selected for limitation in Section X are also selected for limitation in NSPS.

NEW SOURCE PERFORMANCE STANDARDS

The NSPS discharge flows for each wastewater source are the same as the BAT discharge rates listed in Section X. The mass of pollutant allowed to be discharged per mass of product is calculated by multiplying the appropriate achievable treatment concentration by the production normalized wastewater discharge These achievable treatment concentrations are flows (1/kkq). discussed in Section VII. The results of these calculations are the production-based new source performance standards, and are presented in Table XI-2 (page 4501). The NSPS wastewater discharge rates are presented in Table XI-1 (page 4499). Since the discharge flows and the achievable treatment both concentrations are the same for new sources and BAT, the NSPS are identical to the BAT mass limitations.

Table XI-1

NSPS WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

Wastewater Stream		ormalized rge Rate gal/ton	Production Normalizing Parameter
Concentrate Digestion Wet Air Pollution Control	622	149	Concentrate Digested
Solvent Extraction Raffi- nate	9,155	2,195	Concentrate Digested
Solvent Extraction Wet Air Pollution Control	246	60	Concentrate Digested
Precipitation and Filtra- tion Wastewater	13,689	3,283	Concentrate Digested
Precipitation and Filtra- tion Wet Air Pollution Control	6,351	1,523	Concentrate Digested
Tantalum Salt Drying	60,542	14,518	Tantalum Salt Dried
Oxides Calcining Wet Air Pollution Control	3,842	921	Columbium or Tantalum Oxide Calcined
Reduction of Tantalum Salt to Metal	166,071	39,825	Tantalum Salt Reduced
Reduction of Tantalum Salt Metal Wet Air Pollution Control	2,043	490	Tantalum Salt Reduced

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PRIMARY COLUMBIUM AND TANTALUM SECT -XI

Table XI-1 (Continued)

NSPS WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

		ormalized rge Rate	
Wastewater Stream	1/kkg	gal/ton	Production Normalizing Parameter
Tantalum Powder Wash and Scrubber	20,433	4,900	Tantalum Washed
Consolidation and Casting Contact Cooling	0	0	Columbium or Tantalum Cast or Consolidated

ו דX PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - XI

TABLE XI-2

NSPS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(a) Concentrate Digestion Wet Air Pollution Control NSPS

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested Antimony 1.200 0.535 Arsenic 0.865 0.386 Cadmium 0.124 0.050 Chromium 0.230 0.093 Copper 0.796 0.379 *Lead 0.174 0.081 Nickel 0.342 0.230 Selenium 0.510 0.230 Thallium 0.871 0.379	Pollutant o Pollutant P		Maximum for Any One Day	Maximum for Monthly Average
Arsenic0.8650.386Cadmium0.1240.050Chromium0.2300.093Copper0.7960.379*Lead0.1740.081Nickel0.3420.230Selenium0.5100.230Thallium0.8710.379				
Arsenic0.8650.386Cadmium0.1240.050Chromium0.2300.093Copper0.7960.379*Lead0.1740.081Nickel0.3420.230Selenium0.5100.230Thallium0.8710.379	Antimony		1 200	0 535
Cadmium0.1240.050Chromium0.2300.093Copper0.7960.379*Lead0.1740.081Nickel0.3420.230Selenium0.5100.230Thallium0.8710.379	-			
Chromium0.2300.093Copper0.7960.379*Lead0.1740.081Nickel0.3420.230Selenium0.5100.230Thallium0.8710.379				
Copper0.7960.379*Lead0.1740.081Nickel0.3420.230Selenium0.5100.230Thallium0.8710.379				
*Lead0.1740.081Nickel0.3420.230Selenium0.5100.230Thallium0.8710.379				
Nickel0.3420.230Selenium0.5100.230Thallium0.8710.379				
Selenium 0.510 0.230 Thallium 0.871 0.379		a an		
Thallium 0.379				
*Ammonia 82.910 36.450				
*Fluoride 21.770 12.440				
*TSS 9.330 7.464				

Within the range of 7.0 to 10.0 at all times

(b) Solvent Extraction Raffinate NSPS

*pH

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
Metric Units - mg/kg of English Units - 1bs/million 1		
Antimony	17.670	7.873
Arsenic	12.730	5.676
Cadmium	1.831	0.732
Chromium	3.387	1.373
Copper	11.720	5.585
*Lead	2.563	1.190
Nickel	5.035	3.387
Selenium	7.507	3.387
Thallium	12.820	5.585
*Zinc	9.338	3.845
*Ammonia (as N)	1220.000	536.500
*Fluoride	320.400	183.100
*TSS	137.300	109.900
*pH Within the rar	ge of 7.0 to 10	

4501

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - XI

TABLE XI-2 (Continued)

NSPS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(c) Solvent Extraction Wet Air Pollution Control NSPS

Pollutant	or	Maximum	for	Maximu	um for
Pollutant	Property	Any One	Day	Monthly	Average

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested

Antimony Arsenic Cadmium	1	0.475 0.342 0.049	0.212 0.153 0.020
Chromium Copper		0.091 0.315	0.037 0.150
*Lead		0.069	0.032
Nickel		0.135	0.091
Selenium		0.202	0.091
Thallium		0.344	0.150
*Zinc	pro-	0.251	0.103
*Ammonia (as N)	÷	32.790	14.420
*Fluoride		8.610	4.920
*TSS		3.690	2.952
*pH	Within the	e range of 7.0 to 10	0.0 at all times

(d) Precipitation and Filtration NSPS

Pollutant or Pollutant Pr			ximum for y One Day	Maximu Monthly	
M	letric Units - mo	g/kg of com	ncentrate di	gested	
English	Units - 1bs/mi	llion lbs o	of concentra	te digest	ed
Antimony			26.420		11.770
Arsenic			19.030		8.487
Cadmium		•	2.738	•	1.095
Chromium			5.065		2.053
Copper			17.520		8.350
*Lead			3.833		1.780
Nickel			7.529		5.065
Selenium		6	11.230	·	5.065
Thallium			19.170		8.350
*Zinc			13.960		5.750
*Ammonia (as	5 N)		1825.000		02.200
*Fluoride			479.100		73.800
*TSS			205.400	. 1	64.300
*pH	Within	the range o	of 7.0 to 10	.0 at all	times

NSPS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(e) Precipitation and Filtration Wet Air Pollution Control NSPS

Pollutant or Pollutant Property			A	aximum ny One	Day M	onthly	um for Average
Metric U English Units	nits - n - lbs/m	mg/k illi	g of co on lbs	oncentr of con	ate dige centrate	sted digest	ced
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride *TSS *pH	Within	the	range	8 1 2 8 1 3 5 8 6 846 222 95	.260 .828 .270 .350 .129 .778 .493 .208 .891 .478 .600 .300 .270 to 10.0	3	5.462 3.938 0.508 0.953 3.874 0.826 2.350 2.350 3.874 2.668 72.200 27.000 76.210 times

(f) Tantalum Salt Drying NSPS

Pollutant or	. A	aximum for	Maximum for
Pollutant Property		ny One Day	Monthly Average
English Units -	s - mg/kg of ta lbs/million lb:	s of tantalum	salt dried
Antimony	hin the range	116.800	52.070
Arsenic		84.150	37.540
Cadmium		12.110	4.843
Chromium		22.400	9.081
Copper		77.490	36.930
*Lead		16.950	7.870
Nickel		33.300	22.400
Selenium		49.640	22.400
Thallium		84.760	36.930
*Zinc		61.750	25.430
*Ammonia (as N)		8070.000	3548.000
*Fluoride		2119.000	1211.000
*TSS		908.200	726.500
*pH Wit		of 7.0 to 10.	0 at all times

NSPS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(g) Oxides Calcining Wet Air Pollution Control NSPS

Pollutant	or	Maximum	for	Maximur	n for
Pollutant	Property	Any One	Day	Monthly A	lverage

Metric Units - mg/kg of columbium-tantalum oxide dried English Units - lbs/million lbs of columbium-tantalum oxide dried

Antimony						7.415		3.304
Arsenic						5.340		2.382
Cadmium						0.768		0.307
Chromium			L.			1.422		0.576
Copper			i.			4.918		2.344
*Lead						1.076		0.499
Nickel	·					2.113		1.422
Selenium						3.150		1.422
Thallium						5.379		2.344
*Zinc						3.919		1.614
*Ammonia (as	N)		1			512.200		225.200
*Fluoride						134.500		76.840
*TSS						57.630		46.110
*pH		Within	the	range	of	7.0 to 10.0	at	all times

(h) Reduction of Tantalum Salt to Metal NSPS

Pollutant Pollutant	Property			Aı	aximum ny One	Day	Мот	Maximum hthly Av	
	Metric Un								•
Englis	h Units -	lbs/mil	llion	lbs d	of tant	alum	salt	reduced	· .
Antimony					320	.500		142	.800
Arsenic		1			230	.800		103	.000
Cadmium				*	33	3.210		13	.290
Chromium					61	.450		24	.900
Copper					212	2.600		101	.300
*Lead			t		46	5.500		21	.590
Nickel					93	.340		61	.450
Selenium					136	5.200		61	.450
Thallium			÷.,		232	2.500		101	.300
*Zinc					169	9.400		69	.750
*Ammonia (as N)				22140	0.000		9732	.000
*Fluoride	•				5813	3.000		3322	.000
*TSS					249]	.000		1993	.000
*pH		Within	the	range	of 7.0) to 1	L0.0 a	at all t	imes

NSPS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(i) <u>Reduction of Tantalum Salt to Metal Wet Air Pollution</u> <u>Control</u> NSPS

Pollutant or		Maximum for	Maximum for
Pollutant Property		Any One Day	Monthly Average
Metric	Units - mg/k	g of tantalum	salt reduced
English Units - 1bs			
Antimony		3.943	1.757
Arsenic	1	2.840	1.267
Cadmium		0.409	0.163
Chromium		0.756	0.306
Copper		2.615	1.246
*Lead	· •	0.572	0.266
Nickel		1.124	0.756
Selenium		1.675	0.756
Thallium		2.860	1.246
*Zinc		2.084	0.858
*Ammonia (as N)		272.400	119.700
*Fluoride	1	71.510	40.860
*TSS	•	30.650	24.520
*pH	Within the ra	inge of 7.0 to 10	.0 at all times
(j) <u>Tantalum</u> Powde	er wasn NSPS		
	į		
Pollutant or		Maximum for	Maximum for
Pollutant Property		Any One Day	Monthlý Average

Metric Units - mg/kg of tantalum powder washed English Units - lbs/million lbs of tantalum powder washed

Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride *TSS		•		39.440 28.400 4.087 7.560 26.150 5.721 11.240 16.760 28.610 20.840 2724.000 715.200 306.500	17.570 12.670 1.635 3.065 12.460 2.656 7.560 12.460 8.582 198.000 408.700 245.220
*TSS *pH	Within	the	range	306.500 of 7.0 to 10.0	

NSPS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(k) Consolidation and Casting Contact Cooling NSPS

Pollutant or Pollutant Property		Maximum for Any One Day	Maximu Monthly	
Metric Units - mg/kg English Units - lbs/n consc Antimony Arsenic Cadmium	of columbium nillion lbs of olidated	or tantalum c columbium or 0.000 0.000 0.000))	0.000 0.000 0.000
Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N))))))	0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000
*Fluoride *TSS *pH V	Within the rar	0.00)	0.000

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - XII

SECTION XII

PRETREATMENT STANDARDS

This section describes the control and treatment technologies for pretreatment of process wastewaters from existing sources and new sources in the primary columbium-tantalum subcategory. PSES are designed to prevent the discharge of pollutants which pass through, interfere with, or are otherwise incompatible with the operation of publicly owned treatment works (POTW). The Clean Water Act of 1977 requires pretreatment for pollutants, such as toxic metals, that limit POTW sludge management alternatives. indirect discharge facilities, like new direct discharge ities, have the opportunity to incorporate the best New facilities, available demonstrated technologies, including process changes, in-plant controls, and end-of-pipe treatment technologies, and to use plant site selection to ensure adequate treatment system installation. Pretreatment standards are to be technology-based, analogous to the best available or best demonstrated technology for removal of toxic pollutants. Pretreatment standards for regulated pollutants are presented based on the selected control and treatment technology.

TECHNICAL APPROACH TO PRETREATMENT

Before proposing pretreatment standards, the Agency examines whether the pollutants discharged by the industry pass through POTW or interfere with the POTW operation or its chosen the sludge disposal practices. In determining whether pollutants pass through a well-operated POTW, achieving secondary treatment, the Agency compares the percentage of a pollutant removed by POTW with the percentage removed by direct dischargers applying the best available technology economically achievable. A pollutant is deemed to pass through the POTW when the average percentage nationwide by well-operated POTW meeting removed secondary requirements, is less than the percentage removed by treatment direct dischargers complying with BAT effluent limitations guidelines for that pollutant. (See generally, 46 FR at 9415-16 (January 28, 1981).)

This definition of pass through satisfies the two competing objectives set by Congress that standards for indirect dischargers be equivalent to standards for direct dischargers, while at the same time, the treatment capability and performance of the POTW be recognized and taken into account in regulating the discharge of pollutants from indirect dischargers.

The Agency compares percentage removal rather than the mass or concentration of pollutants discharged because the latter would not take into account the mass of pollutants discharged to the POTW from non-industrial sources nor the dilution of the pollutants in the POTW effluent to lower concentrations due to the addition of large amounts of non-industrial wastewater.

PRETREATMENT STANDARDS FOR EXISTING AND NEW SOURCES

Options for pretreatment of wastewaters are based on increasing the effectiveness of end-of-pipe treatment technologies. All inplant changes and applicable end-of-pipe treatment processes have been discussed previously in Sections X and XI. The options for PSES and PSNS, therefore, are the same as the BAT options discussed in Section X.

A description of each option is presented in Section X, while a more detailed discussion, including pollutants controlled by each treatment process and achievable treatment concentrations are presented in Section VII of the General Development Document.

The treatment technology options for the PSES and PSNS are:

Option A

- o Chemical precipitation and sedimentation
- o Ammonia steam stripping of wastewaters containing treatable concentrations of ammonia

Option B

- o Chemical precipitation and sedimentation
- o Ammonia steam stripping of wastewaters containing treatable concentrations of ammonia
- o In-process flow reduction

Option C

- o Chemical precipitation and sedimentation
- o Ammonia steam stripping of wastewaters containing
- treatable concentrations of ammonia
- o In-process flow reduction
- o Multimedia filtration

INDUSTRY COST AND POLLUTANT REMOVAL ESTIMATES

The industry cost and environmental benefits of each treatment option were used to determine incremental removals and costs. The methodology applied in calculating pollutant reduction benefits and plant compliance costs is discussed in Section X. Table XII-1 (page 4510) shows the estimated pollutant removals for indirect dischargers, while compliance costs are presented in Table VIII-2 (page 4453).

PSES AND PSNS OPTION SELECTION

EPA is promulgating PSES equal to BAT for this subcategory. It is necessary to promulgate PSES to prevent pass-through of lead, zinc, fluoride, and ammonia. The technology basis for PSES is lime precipitation and sedimentation, ammonia steam stripping, wastewater flow reduction and filtration. Flow reduction for the

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - XII

selected technology represents an 80 percent reduction in flow over current discharge rates.

Implementation of the promulgated PSES limitations would remove an estimated 18,590 kg/yr of toxic pollutants, 290,460 kg/yr of ammonia, and 400,175 kg/yr of fluoride over estimated raw discharge. The final PSES effluent mass limitations will remove 57 kg/yr of toxic metals over the intermediate PSES options considered, which lacks filtration. Both options are economically achievable and both prevent pass-through. The Agency is thus selecting PSES equal to BAT. The estimated capital cost for achieving PSES is \$1.0 million (March, 1982 dollars), and the annual cost is \$0.7 million.

The technology basis for promulgated PSNS is identical to NSPS, PSES, and BAT. The same pollutants pass through as at PSES, for the same reasons. EPA knows of no economically feasible, demonstrated technology that is better than PSES technology. The flow allowances are based on minimization of process PSES whenever possible through the of wastewater use lime precipitation and sedimentation to remove fluoride for scrubbing wastewater. Because PSNS does not include wet any additional costs compared to NSPS and PSES, the Agency does not believe it will prevent entry of new plants.

REGULATED POLLUTANT PARAMETERS

The pollutants and pollutant parameters selected for limitation, in accordance with the rationale of Section X, are identical to those selected for limitation for BAT. PSES and PSNS prevent the pass-through of lead, zinc, fluoride, and ammonia.

PRETREATMENT STANDARDS

The PSES and PSNS discharge flows are identical to the BAT discharge flows for all processes. These discharge flows are listed in Table XII-2 (page 4511). The mass of pollutant allowed to be discharged per mass of product is calculated by multiplying the achievable treatment concentration (mg/l) by the normalized wastewater discharge flow (1/kkg). Pretreatment standards for existing and new sources, as determined from the above procedure, are shown in Tables XII-3 and XII-4 (pages 4513 and 4519) for each waste stream.

Mass-based standards are promulgated for the columbium-tantalum subcategory to ensure that the standards are achieved by means of pollutant removal rather than by dilution. They are particularly important since the standards are based upon flow reduction; pollutant limitations associated with flow reduction cannot be measured any other way but as a reduction of mass discharged. The flow reduction over estimated current flow for the columbiumtantalum subcategory is 80 percent.

Table XII-1

POLLUTANT REMOVAL ESTIMATES FOR PRIMARY COLUMBIUM-TANTALUM INDIRECT DISCHARGERS

TANTALI	TOTAL RAW WASTE (kg/yr)	OPTION A DISCHARGED (kg/yr)	OPTION A REMOVED (kg/yr)	OPTION B DISCHARGED (kg/yr)	OPTION B REMOVED (kg/yr)	OPTION C DISCHARGED (kg/yr)	OPTION C REMOVED (kg/yr)
	119.4	47.4	72.0	30.9	88.5	20.7	198.7
Antimony Arsenic	171.8	34.5	137.3	22.5	149.3	15.0	156.8
Cadmium	137.7	5.3	132.3	3.5	134.2	2.2	135.5
Cadalium	6,326.4	5.7	6,320.7	3.7	6,322.7	3.1	6,323.3
	1,065.8	39.2	1,026.6	25.6	1,040.2	17.2	1,048.6
Copper Lead	5,501.0	8.1	5,492.9	5.3	5,495.7	3.5	5,497.5
Nickel	114.6	50.1	64.5	32.7	81.9	9.7	104.9
Zinc	5,234.0	22.3	5,211.6	14.6	5,219.4	10.1	5,223.8
TUTAL TUXIC METALS	18,670.7	212.7	18,458.0	138.7	18,532.0	81.6	18,589.1
· • • • • • • •	2 151 2	151.6	2,999.8	98.8	3,052.5	65.7	3,085.6
Aluminum	3,151.3	2,165.2	289,709.9	1,411.9	290,463.1	1,411.9	290,463.1
Ammonia	291,875.0	981.1	399,834.3	639.8	400,175.6	639.8	400,175.6
Fluoride Iron	400,815.4 107,248.8	27.7	107,221.1	18.1	107,230.7	12.4	107,236.5
TUTAL NONCONVENTIONALS	803,090.6	3,325.6	799,765.0	2,168.6	800,921.9	2,129.8	800,960.8
TSS	658,952.3	811.9	658,140.3	529.5	658,422.8	114.7	658,837.6
TUTAL CONVENTIONALS	658,952.3	811.9	658,140.3	529.5	658,422.8	114.7	658,837.6
TUTAL POLIJITANTS	1,480,713.5	4,350.2	1,476,363.4	2,836.8	1,477,876.7	2,326.1	1,478,387.4
FLOW (1/yr)		67,662,000		44,123,000		44,123,000	

NOTE: TOTAL TOXIC METALS = Antimony + Arsenic + Cadmium + Chromium + Copper + Lead + Nickel + Zinc TOTAL NUMCONVENTIONALS = Aluminum + Ammonia + Fluoride + Iron TOTAL CONVENTIONALS = TSS TUTAL POLIJIANTS = Total Toxic Metals + Total Nonconventionals + Total Conventionals

OPTION A = Lime Precipitation, Sedimentation, and Ammonia Steam Stripping OPTION B = Option A, plus in-Process Flow Reduction OPTION C = Option B, plus Multimedia Filtration

Table XII-2

PSES AND PSNS WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

	Norm	nd PSNS alized rge Rate			
Wastewater Stream	1/kkg	gal/ton	Production Normalizing Parameter		
Concentrate Digestion Wet Air Pollution Control	622	149	Concentrate Digested		
Solvent Extraction Raffi- nate	9,155	2,195	Concentrate Digested		
Solvent Extraction Wet Air Pollution Control	246	60	Concentrate Digested		
Precipitation and Filtra- tion Wastewater	13,689	3,283	Concentrate Digested		
Precipitation and Filtra- tion Wet Air Pollution Control	6,351	1,523	Concentrate Digested		
Tantalum Salt Drying	60,542°	14,518	Tantalum Salt Dried		
Oxides Calcining Wet Air Pollution Control	3,842	921	Columbium or Tantalum Oxide Calcined		
Reduction of Tantalum Salt to Metal	166,071	39,825	Tantalum Salt Reduced		
Reduction of Tantalum Salt Metal Wet Air Pollution Control	2,043	490	Tantalum Salt Reduced		

PRIMARY COLUMBIUM AND TANTALUM SECT I XII

Table XII-2

PSES AND PSNS WASTEWATER DISCHARGE RATES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

		nd PSNS alized ge Rate	
Wastewater Stream	1/kkg	gal/ton	Production Normalizing Parameter
Tantalum Powder Wash and Scrubber	20,433	4,900	Tantalum Washed
Consolidation and Casting Contact Cooling	0	0	Columbium or Tantalum Cast or Consolidated

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SECT

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XII

TABLE XII-3

PSES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(a) Concentrate Digestion Wet Air Pollution Control PSES

Pollutant or	Maximum for Ma	aximum for				
Pollutant Property	Any One Day Mon	thly Average				
Metric Units - mg/kg of concentrate digested						
English Units - 1bs/mi	llion lbs of concentrate d	igested				
Antimony	1.200	0.535				
Arsenic	0.865	0.386				
Cadmium	0.124	0.050				
Chromium	0.230	0.093				
Copper	0.796	0.379				
*Lead	0.174	0.081				
Nickel	0.342	0.230				
Selenium	0.510	0.230				
Thallium	0.871	0.379				
*Zinc	0.635	0.261				
*Ammonia (as N)	82.910	36.450				
*Fluoride	21.770	12.440				

(b) Solvent Extraction Raffinate PSES PSES

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average			
Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested					
5		5			
Antimony	17.670	7.873			
Arsenic	12.730	5.676			
Cadmium	1.831	0.732			
Chromium	3.387	1.373			
Copper	11.720	5.585			
*Lead	2.563	1.190			
Nickel	5.035	3.387			
Selenium	7.507	3.387			
Thallium	12.820	5.585			
*Zinc	9.338	3.845			
*Ammonia (as N)	1221.000	536.500			
*Fluoride	320.400	183.100			

PSES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(c) Solvent Extraction Wet Air Pollution Control PSES

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
	mg/kg of concentrate o million lbs of concentr	
Antimony	0.475	0.212

Arsenic	0.342	0.153
Cadmium	0.049	0.020
Chromium	0.091	0.037
Copper	0.315	0.150
*Lead	0.069	0.032
Nickel	0.135	0.091
Selenium	0.202	0.091
Thallium	0.344	0.150
*Zinc	0.251	0.103
*Ammonia (as N)	32.790	14.420
*Fluoride	8.610	4.920

(d) Precipitation and Filtration PSES

Pollutant or		Maximum	for	Maximum for
Pollutant Prop	erty	Any One	Day	Monthly Average

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested

Antimony	26.420	11.770
Arsenic	19.030	8.487
Cadmium	2.738	1.095
Chromium	5.065	2.053
Copper	17.520	8.350
*Lead	3.833	1.780
Nickel	7.529	5.065
Selenium	11.230	5.065
Thallium	19.170	8.350
*Zinc	13.960	5.750
*Ammonia (as N)	1825.000	802.200
*Fluoride	479.100	273.800
	The second s	

PSES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(e)Precipitation and Filtration Wet Air Pollution Control PSES

Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested 12.260 5.462 Antimony Arsenic 3.938 8.828 Cadmium 1.270 0.508 Chromium 2.350 0.953 8.129 3.874 Copper hea.T* 1.778 0.826

"Deau		T•//Q	0.020
Nickel	1	3.493	2.350
Selenium		5.208	2.350
Thallium	1	8.891	3.874
*Zinc		6.478	2.668
*Ammonia (as N)		846.600	372.200
*Fluoride		222.300	127.000

(f) Tantalum Salt Drying PSES PSES

Pollutant or Pollutant Property		Maximum for Any One Day	Maximum for Monthly Average
	Jnits - mg/kg of		
English Units	s - lbs/million	lbs of tantalu	m salt dried
Antimony		116.800	52.070
Arsenic		84.150	37.540
Cadmium		12.110	4.843
Chromium		22.400	9.081
Copper		77.490	36.930
*Lead		16.950	7.870
Nickel		33.300	22.400
Selenium		49.640	22.400
Thallium		84.760	36.930
*Zinc		61.750	25.430
*Ammonia (as N)		8070.000	3548.000
*Fluoride	۰ ۰	2119.000	1211.000

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - XII

TABLE XII-3 (Continued)

PSES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(g) Oxides Calcining Wet Air Pollution Control PSES

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Pollutant orMaximum forMaximum forPollutant PropertyAny One DayMonthly Average

Metric Units - mg/kg of columbium-tantalum oxide dried English Units - lbs/million lbs of columbium-tantalum oxide dried

Antimony	7.415	3,304
Arsenic	5.340	2.382
Cadmium	0.768	0.307
Chromium	1.442	0.576
Copper	4.918	2.344
*Lead	1.076	0.500
Nickel	2.113	1.422
Selenium	3.150	1.422
Thallium	5.379	2.344
*Zinc	3.919	1.614
*Ammonia (as N)	512.200	225.200
*Fluoride	134.500	76.840

(h) <u>Reduction of Tantalum Salt to Metal PSES</u>

Pollutant or Pollutant P		Maximum for Any One Day	Maximum for Monthly Average
M	etric Units - mg/kg o	f tantalum salt r	educed
Englis	h Units - lbs/million	lbs of tantalum	salt reduced
Antimony	•	320.500	142.800
Arsenic		230.800	103.000
Cadmium		33.210	13.290
Chromium		61.450	24.910
Copper		212.600	101.300
*Lead		46.500	21.590
Nickel		51.340	61.450
Selenium		136.200	61.450
Thallium		232.500	101.300
*Zinc		169.400	69.750
		22,140.000	9,732.000
*Ammonia (a *Fluoride	2 11 /	5,813.000	3,322.000

PSES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(i) <u>Reduction of</u> <u>Tantalum Salt to Metal Wet Air</u> <u>Pollution</u> <u>Control</u> <u>PSES</u>

Pollutant Pollutant		Maximum for Any One Day	Maximum for Monthly Average
	Metric Units - mg/kg of sh Units - lbs/million		
Antimony		3.943	1.757
Arsenic		2.840	1.267
Cadmium		0.409	0.163
Chromium		0.756	0.306
Copper		2.615	1.246
*Lead		0.572	0.266
Nickel		1.124	0.756
Selenium		1.675	0.756
Thallium		2.860	1.246
*Zinc		2.084	0.858
*Ammonia (as N)	272.400	119.700
*Fluoride		71.510	40.860

(j) <u>Tantalum</u> Powder Wash PSES

Pollutant Pollutant			Maximum Any One		Maximu Monthly	
		ts - mg/kg of				
Englis	h Units -	lbs/million l	bs of tant	alum p	powder was	shed
Antimony	н. 1	÷	30	9.440		17.570
Arsenic				3.400		12.670
Cadmium		1		1.087		1.635
Chromium				7.560		3.065
Copper		i	26	5.150		12.460
*Lead				5.721		2.656
Nickel			11	.240		7.560
Selenium			16	5.760		7.560
Thallium		8	28	8.610		12.460
*Zinc			20).840		8.582
*Ammonia (as N)		2,724	.000	1 ,]	.98.000
*Fluoride			715	5.200	4	108.700

PSES FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(k) Consolidation and Casting Contact Cooling PSES

Pollutant or Pollutant Pro	Maximum Any One	Maximu Monthly	

Metric Units - mg/kg of columbium or tantalum cast or consolidated English Units - lbs/million lbs of columbium or tantalum cast or consolidated

Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc	$\begin{array}{c} 0.000\\ 0.$	0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000

PRIMARY COLUMBIUM AND TANTALUM SUBCATEGORY SECT - XII

TABLE XII-4

PSNS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(a) Concentrate Digestion Wet Air Pollution Control PSNS

Pollutant	or	Maximum	for	Maximum for
Pollutant	Property	Any One	Day	Monthly Average

Metric Units - mg/kg of concentrate digested				
	nillion lbs of concentrate d			
		•		
Antimony	1.200	0.535		
Arsenic	0.865	0.386		
Cadmium	0.124	0.050		
Chromium	0.230	0.093		
Copper	0.796	0.379		
*Lead	0.174	0.081		
Nickel	0.342	0.230		
Selenium	0.510	0.230		
Thallium	0.871	0.379		
*Zinc	0.635	0.261		
*Ammonia (as N)	82.910	36.450		
*Fluoride	21.770	12.440		

(b) Solvent Extraction Raffinate PSNS

Pollutant or	1	Maximum	for	Maximum for
Pollutant Property		Any One	Day	Monthly Average

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested

Antimony		17.670	7.873
Arsenic		12.730	5.676
Cadmium		1.831	0.732
Chromium		3.387	1.373
Copper		11.720	5.585
*Lead		2.563	1.190
Nickel		5.035	3.387
Selenium		7.507	3.387
Thallium		12.820	5.585
*Zinc		9.338	3.845
*Ammonia (as N)		1,221.000	536.500
*Ammonia (as N) *Fluoride	• • •	1,221.000 320.400	536.500 183.100

PSNS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(c) Solvent Extraction Wet Air Pollution Control PSNS

Pollutant o	٦r	Maximum	for	Maximu	ım for
Pollutant H		Any One	Day	Monthly	Average
	11	_			,

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested

Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N)	0.475 0.342 0.049 0.091 0.315 0.069 0.1353 0.202 0.344 0.251 32.790 8.610	$\begin{array}{c} 0.212\\ 0.153\\ 0.020\\ 0.037\\ 0.150\\ 0.032\\ 0.091\\ 0.091\\ 0.150\\ 0.103\\ 14.420\\ 4.920\end{array}$
*Fluoride	8.610	4.920

(d) Precipitation and Filtration PSNS

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
Metric Units - mg/kg c	of concentrate di	gested
English Units - lbs/million	lbs of concentra	te digested
Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride	26.420 19.030 2.738 5.065 17.522 3.833 7.529 11.230 19.170 13.960 1,825.000 479.100	11.770 8.487 2.053 2.053 8.350 1.780 5.065 5.065 8.350 5.750 802.200 273.800

PSNS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(e) Precipitation and Filtration Wet Air Pollution Control PSNS

Pollutant or		Maximum	for	Maximum for
Pollutant Proper	=y	Any One	Day	Monthly Average

Metric Units - mg/kg of concentrate digested English Units - lbs/million lbs of concentrate digested

Antimony	12.260	5.462
Arsenic	8.828	3.938
Cadmium	1.270	0.508
Chromium	2.350	0.963
Copper	8.129	3.874
*Lead	1.788	0.826
Nickel	3.493	2.350
Selenium	5.208	2.350
Thallium	8.891	3.874
*Zinc	6.478	2.668
*Zinc	6.478	2.668
*Ammonia (as N)	846.600	372.200
*Fluoride	222.300	127.000

(f) Tantalum Salt Drying PSNS

Pollutant o Pollutant P		, , i	Maximum Any One		Maximu Monthly	
······································	Metric Units - m	g/kg of	tantalur	n salt o	dried	· · · · · · · · · · · · · · · · · · ·
	sh Units - 1bs/m					ied
	•	ì				
Antimony			116	5.800		52.070
Arsenic			84	4.150		37.540
Cadmium			12	2.110		4.843
Chromium			22	2.400		9.081
Copper			77	7.490		36.930
*Lead			16	5.950		7.871
Nickel			33	3.300		22.400
Selenium			49	0.640		22.400
Thallium			84	1.760		36.930
*Zinc		i.	61	.750		25.430
*Ammonia (a	SN)		8,070	0.000	3,5	548.000
*Fluoride			2,119	000.	1,2	211.000

PSNS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(g) Oxides Calcining Wet Air Pollution Control PSNS

Pollutant or Pollutant Property	Maximum for Any One Day	

Metric Units - mg/kg of columbium-tantalum oxide dried English Units - lbs/million lbs of columbium-tantalum oxide dried

Antimony Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia (as N) *Fluoride	7.415 5.340 0.768 1.422 4.918 1.076 2.113 3.150 5.379 3.919 512.200 134.500	2.382 0.307 0.576 2.344 0.500 1.422 1.422 2.344 1.614 225.200 76.840
--	--	--

(h) <u>Reduction of Tantalum Salt to Metal</u> PSNS

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
FOILdeane iloperoy	1	

Metric Units - mg/kg of tantalum salt reduced English Units - lbs/million lbs of tantalum salt reduced

PSNS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(i) <u>Reduction of Tantalum Salt to Metal Wet Air Pollution</u> Control PSNS

Pollutant Pollutant		Maximum for Any One Day	Maximum for Monthly Average
Engli	Metric Units - mg/kg of ish Units - lbs/million		
2.1.9 - 1			
Antimony		3.943	1.757
Arsenic		2.840	1.267
Cadmium	1	0.409	0.163
Chromium		0.756	0.306
Copper		2.615	1.246
*Lead		0.572	0.266
Nickel		1.124	0.756
Selenium		1.675	0.756
Thallium	:	2.860	1.246
*Zinc		2.084	0.858
*Ammonia ((as N)	272.400	119.700
*Fluoride		71.510	40.860

(j) Tantalum Powder Wash PSNS

Metric Units - mg/kg of tantalum powder washed English Units - lbs/million lbs of tantalum powder washed Antimony 39.440 17.570 Arsenic 28.400 12.670 Cadmium 4.087 3.065 Chromium 7.560 3.065 Copper 26.150 12.460 *Lead 5.721 2.656 Nickel 11.240 7.560 Selenium 16.760 7.560 Thallium 28.610 12.460 *Zinc 20.840 8.582	Pollutant Pollutant		Maximum for Any One Day	Maximum for Monthly Average		
Arsenic28.40012.670Cadmium4.0873.065Chromium7.5603.065Copper26.15012.460*Lead5.7212.656Nickel11.2407.560Selenium16.7607.560Thallium28.61012.460*Zinc20.8408.582						
*Fluoride 715.200 408.700	Arsenic Cadmium Chromium Copper *Lead Nickel Selenium Thallium *Zinc *Ammonia	(as N)	28.400 4.087 7.560 26.150 5.721 11.240 16.760 28.610 20.840 2,724.000	12.670 3.065 3.065 12.460 2.656 7.560 7.560 12.460 8.582 1,198.000		

PSNS FOR THE PRIMARY COLUMBIUM-TANTALUM SUBCATEGORY

(k) Consolidation and Casting Contact Cooling PSNS

.

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average

Metric Units - mg/kg of columbium or tantalum cast or consolidated English Units - lbs/million lbs of columbium or tantalum cast or consolidated

Antimony	0.000	0.000
Arsenic	0.000	0.000
Cadmium	0.000	0.000
	0.000	0.000
Chromium	0.000	0.000
Copper	0.000	0.000
*Lead	0.000	0.000
Nickel	0.000	0.000
Selenium	0.000	0.000
Thallium	0.000	0.000
*Zinc	0.000	0.000
*Ammonia (as N)	0.000	0.000
*Fluoride	0.000	51000

SECTION XIII

BEST CONVENTIONAL POLLUTANT CONTROL TECHNOLOGY

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EPA is not promulgating best conventional pollutant control technology (BCT) for the primary columbium-tantalum subcategory at this time.

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NONFERROUS METALS MANUFACTURING POINT SOURCE CATEGORY

DEVELOPMENT DOCUMENT SUPPLEMENT

for the

Secondary Tantalum Subcategory

William K. Reilly Administrator

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May 1989

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SECTION I

SUMMARY AND CONCLUSIONS

This document provides the technical basis for promulgating effluent limitations based on best practicable technology (BPT) and best available technology (BAT) for existing direct dischargers, pretreatment standards for new indirect dischargers (PSNS), and standards of performance for new source direct dischargers (NSPS).

The secondary tantalum subcategory consists of three plants. All three plants treat their process wastewater and discharge the effluent directly to rivers or streams.

EPA first studied the secondary tantalum subcategory to determine whether differences in raw materials, final products, manufacturing processes, equipment, age and size of plants, or water usage, required the development of separate effluent limitations and standards for different segments of the This involved a detailed analysis of wastewater subcategory. discharge and treated effluent characteristics, including the sources and volume of water used, the processes used, the sources of pollutants and wastewaters in the plant, and the constituents of wastewaters, including priority pollutants. As a result, five subdivisions have been identified for this subcategory that warrant separate effluent limitations. These are:

- o Tantalum alloy leach and rinse,
- o Capacitor leach and rinse,
- o Tantalum sludge leach and rinse,
- o Tantalum powder acid wash and rinse, and
- o Leaching wet air pollution control.

EPA also identified several distinct control and treatment technologies (both in-plant and end-of-pipe) applicable to the secondary tantalum subcategory. The Agency analyzed both historical and newly generated data on the performance of these technologies, including their nonwater quality environmental impacts and air quality, solid waste generation, and energy requirements. EPA also studied various flow reduction techniques reported in the data collection portfolios (dcp) and plant visits.

Engineering costs were prepared for each of the control and treatment options considered for the subcategory. These costs were then used by the Agency to estimate the impact of implementing the various options on the subcategory. For each control and treatment option that the Agency found to be most effective and technically feasible in controlling the discharge of pollutants, we estimated the number of potential closures, number of employees affected, and impact on price. These results are reported in a separate document entitled "The Economic Impact

SECT - I

Analysis of Effluent Limitations and Standards for the Nonferrous Metals Manufacturing Industry."

After examining the various treatment technologies, the Agency has identified BPT to represent the average of the best existing technology. Metals removal based on chemical precipitation and sedimentation technology (lime and settle) is the basis for the BPT limitations. To meet the BPT effluent limitations based on this technology, the secondary tantalum subcategory is expected to incur an estimated capital cost of \$6,462 and an annual cost of \$58,854.

For BAT, filtration is added as an effluent polishing step to the BPT end-of-pipe treatment scheme. To meet the BAT effluent limitations based on this technology, the secondary tantalum subcategory is estimated to incur a capital cost of \$13,474 and an annual cost of \$63,466.

source performance standards are equivalent to BAT. In New NSPS, EPA recognizes that new plants have the selecting implement the best efficient and most opportunity to manufacturing processes and treatment technology. As such, the technology basis of BAT has been determined as the best demonstrated technology.

PSES is not being promulgated for this subcategory because there are no existing indirect dischargers in the secondary tantalum subcategory. For PSNS, the Agency selected end-of-pipe treatment techniques equivalent to BAT.

The best conventional technology (BCT) replaces 8AT for the control of conventional pollutants. BCT is not being promulgated at this time because the methodology for BCT has not yet been finalized.

The mass limitations and standards for BPT, BAT. NSPS, and PSNS are presented in Section II.

SECTION II

CONCLUSIONS

EPA has divided the secondary tantalum subcategory into five subdivisions for the purpose of effluent limitations and standards. These subdivisions are:

(a) Tantalum alloy leach and rinse,

(b) Capacitor leach and rinse,

(c) Tantalum sludge leach and rinse,

(d) Tantalum powder acid wash and rinse, and

(e) Leaching wet air pollution control.

BPT is promulgated based on the performance achievable by the application of chemical precipitation and sedimentation technology. The following BPT effluent limitations are promulgated:

(a) Tantalum Alloy Leach and Rinse BPT

Pollutant orMaximum forMaximum forPollutant PropertyAny One DayMonthly Average

mg/kg (lb/million lbs) of tantalum powder produced

Copper	438.100	230.600
Lead	96.850	46.120
Nickel	442.800	292.900
Zinc	336.700	140.700
Tantalum	103.800	
TSS	9,455.000	4,497.000
pH	Within the range of 7.5	to 10.0 at all times

(b) Capacitor Leach and Rinse BPT

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average

mg/kg (lb/million lbs) of tantalum powder produced from leaching

Copper	38.380	20.200
Lead	8.484	4.040
Nickel	38.780	25.650
Zinc	29.490	12.320
Tantalum	9.090	
TSS	828.200	393.900
pH	Within the range of 7.5 t	to 10.0 at all times

SECONDARY TANTALUM SUBCATEGORY SECT - II

Tantalum Sludge Leach and Rinse BPT (C) Maximum for Pollutant or Maximum for Any One Day Pollutant Property Monthly Average mg/kg (lb/million lbs) of equivalent pure tantalum powder produced 390.100 205.300 Copper 86.230 41.060 Lead 394.200 260.700 Nickel Zinc 299.700 125.200 Tantalum 92.390 TSS 8,417.000 4,003.000 Within the range of 7.5 to 10.0 at all times рH

(d) Tantalum Powder Acid Wash and Rinse BPT

Pollutan Pollutant P		Maximum for Any One Day	Maximum for Monthly Average
mg/k	g (lb/milli	on lbs) of tant	alum powder produced
Copper Lead Nickel Zinc Tantalum TSS pH	Within the	0.665 0.147 0 672 0.511 0.158 14.350 range of 7.5 t	0 350 0.070 0.445 0.214 6.825 0 10.0 at all times

(e) Leaching Wet Air Pollution Control

Pollutan Pollutant P		Maximum for Any One Day	Maximum for Monthly Average	`
mg/kg (lb/m	illion lbs)	of equivalent	pure tantalum powder	produced
Copper Lead Nickel Zinc Tantalum TSS		9.272 2.050 9.370 7.125 2.196 200.100	4.880 0.976 6.198 2.977 95.160	
рН	Within the	range of 7.5 of	of 10.0 at all times	

BAT is promulgated based on the performance achievable by the application of chemical precipitation, sedimentation, and multimedia filtration technology. The following BAT effluent limitations are promulgated:

(a) Tantalum Alloy Leach and Rinse BAT

Pollutant or	Maximum	for	Maximum for
Pollutant Property	Any One	Day	Monthly Average

mg/kg (lb/million lbs) of tantalum powder produced

Copper	295.200	140.700
Lead	64.570	29.980
Nickel	126.800	85.320
Zinc	235.200	96.850
Tantalum	103.800	

(b) Capacitor Leach and Rinse BAT

Pollutant or	Maximum for	Maximum for	-
Pollutant Property	Any One Day	Monthly Average	

mg/kg (lb/million lbs) of tantalum powder produced from leaching

Copper	25.860	12.320
Lead	5.656	2.626
Nickel	11.110	7.474
Zinc	20.600	8.484
Tantalum	9.090	

(c) Tantalum Sludge Leach and Rinse BAT

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

Copper	262.800	125.200
Lead	57.480	26.690
Nickel	112.900	75.960
Zinc Tantalum	209.400 92.390	86.230

Pollutar Pollutant H		Maximum for Any One Day	Maximum for Monthly Average
mg/k	g (lb/mill	ion lbs) of tant	talum powder produced
Copper Lead Nickel Zinc Tantalum		0.448 0.098 0.193 0.357 0.158	0.214 0.046 0.130 0.147

(d) Tantalum Powder Acid Wash and Rinse BAT

(e) Leaching Wet Air Pollution Control BAT

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

NSPS are promulgated based on the performance achievable by the application of chemical precipitation, sedimentation, and multimedia filtration technology The following effluent standards are promulgated for new sources:

(a) <u>Tantalum Alloy Leach</u> and <u>Rinse</u> NSPS

Pollutant Pollutant Pr		Maximum for Any One Day	Maximum for Monthly Average
mg/kg	(lb/milli	on lbs) of tant	alum powder produced
Copper Lead Nickel Zinc Tantalum TSS pH	Within th	295.200 64.570 126.800 235.200 103.800 3,459.000 e range of 7.5	120.700 29.980 85.320 96.850 2,767.000 to 10.0 at all times

SECONDARY TANTALUM SUBCATEGORY SECT - II

(b) <u>Capacitor</u> <u>Leach</u> and <u>Rinse</u> NSPS

	t or	Maximum for	Maximum for
Pollutant P	roperty	Any One Day	Monthly Average
mg/kg (lb/m	illion lbs)	of tantalum	powder produced from lead
Copper	,	25.860	12.320
Lead		5.656	2.626
Nickel		11.110	7.474
Zinc		20.600	8.484
Tantalum		9.090	
TSS		303.000	242.400
рH	Within the	range of 7.9	5 to 10.0 at all times
(c) <u>Tantal</u>	um Sludge Lo	each and Rins	se NSPS
Pollutan	t or	Maximum for	Maximum for
Pollutant P	roperty	Any One Day	Monthly Average
		57.480	26.690
Lead Nickel Zinc Tantalum TSS pH		112.900 209.400 92.390 3,080.000 range of 7.5	75.960 86.230 2,464.000 5 to 10.0 at all times
	um Powder Ad	112.900 209.400 92.390 3,080.000 range of 7.9 cid Wash and	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse</u> NSPS
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan	um Powder Ad	112.900 209.400 92.390 3,080.000 range of 7.5 cid Wash and Maximum for	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse</u> NSPS <u>Maximum for</u>
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan	um Powder Ad	112.900 209.400 92.390 3,080.000 range of 7.9 cid Wash and	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse</u> NSPS <u>Maximum for</u>
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan Pollutant P	um Powder Ad t or roperty	112.900 209.400 92.390 3,080.000 range of 7.5 cid Wash and Maximum for Any One Day	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse</u> NSPS <u>Maximum for</u>
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan Pollutant P mg/k	um Powder Ad t or roperty	112.900 209.400 92.390 3,080.000 range of 7.5 cid Wash and Maximum for Any One Day	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse NSPS</u> <u>Maximum for</u> Monthly Average antalum powder produced 0.214
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan Pollutant P mg/k Copper	um Powder Ad t or roperty	112.900 209.400 92.390 3,080.000 range of 7.5 cid Wash and Maximum for Any One Day	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse NSPS</u> <u>Maximum for</u> Monthly Average antalum powder produced
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan Pollutant P mg/k Copper Lead	um Powder Ad t or roperty	112.900 209.400 92.390 3,080.000 range of 7.5 cid Wash and Maximum for Any One Day on 1bs) of ta 0.448	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse NSPS</u> <u>Maximum for</u> Monthly Average antalum powder produced 0.214
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan Pollutant P mg/k Copper Lead Nickel	um Powder Ad t or roperty	112.900 209.400 92.390 3,080.000 range of 7.5 cid Wash and Maximum for Any One Day on 1bs) of ta 0.448 0.098	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse NSPS</u> <u>Maximum for</u> Monthly Average antalum powder produced 0.214 0.046
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan Pollutant P	um Powder Ad t or roperty	112.900 209.400 92.390 3,080.000 range of 7.5 <u>cid Wash and</u> <u>Maximum for</u> Any One Day on 1bs) of ta 0.448 0.098 0.193 0.357 0.15	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse NSPS</u> <u>Maximum for</u> Monthly Average antalum powder produced 0.214 0.046 0.130
Lead Nickel Zinc Tantalum TSS pH (d) <u>Tantal</u> Pollutan Pollutant P mg/k Copper Lead Nickel Zinc	um Powder Ad t or roperty	112.900 209.400 92.390 3,080.000 range of 7.5 <u>cid Wash and</u> <u>Maximum for</u> Any One Day on 1bs) of ta 0.448 0.098 0.193 0.357	75.960 86.230 2,464.000 5 to 10.0 at all times <u>Rinse NSPS</u> <u>Maximum for</u> Monthly Average antalum powder produced 0.214 0.046 0.130

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(e) Leaching Wet Air Pollution Control NSPS

Pollutant of		Maximum for	Maximum for
Pollutant Prope		Any One Day	Monthly Average
mg/kg (lb/mill: powder produced		of equivalent	pure tantalum
Copper Lead Nickel Zinc Tantalum		6.246 1.366 2.684 4.978 2.196	2.977 0.634 1.806 2.050
TSS	thin the	73.200	58.560
pH Wit		range of 7.5	to 10.0 at all times

PSES are not being promulgated for the secondary tantalum subcategory at this time because there are no existing indirect dischargers in the secondary tantalum subcategory.

PSNS are promulgated based on the performance achievable by the application of chemical precipitation, sedimentation, and multimedia filtration technology. The following pretreatment standards are promulgated for new sources:

(a) Tantalum Alloy Leach and Rinse PSNS

or operty	Maximum for Any One Day	Maximum for Monthly Average
(lb/mil)	lion lbs) of ta	ntalum powder produced
	295.200	140.700
	64.570	29.980
	126.800	85.320
	235.200	96.850
	103.800	
	operty	operty Any One Day (lb/million lbs) of ta 295.200 64.570 126.800 235.200

(b) Capacitor Leach and Rinse PSNS

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
mg/kg (lb/million lbs)	of tantalum	powder produced from leaching
Copper	25.860	12.320
Lead	5.656	2.626
Nickel	11.110	7.474
Zinc	20.600	8.484
Tantalum	9.090	

(c) <u>Tantalum Sludge Leach and Rinse</u> PSNS

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

262.800	125.200
57.480	26.690
112.900	75.960
209.400	86.230
92.390	
	112.900 209.400

(d) Tantalum Powder Acid Wash and Rinse PSNS

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average
mg/kg (lb/million lbs) of tantalum	powder produced
Copper	0.448	0.214
Lead	0.098	0.046
Nickel	0.193	0.130
Zinc	0.357	0.147
Tantalum	0.158	

SECONDARY TANTALUM SUBCATEGORY SECT - II

(e) Leaching Wet Air Pollution Control PSNS

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average	
mg/kg (lb/million lbs) powder produced	of equivalent	pure tantalum	
Copper Lead Nickel Zinc Tantalum	6.246 1.366 2.684 4.978 2.196	2.977 0.634 1.806 2.050	

EPA is not promulgating BCT at this time for the secondary tantalum subcategory.

SECTION III

INDUSTRY PROFILE

This section of the secondary tantalum supplement describes the raw materials and processes used in producing secondary tantalum and presents a profile of the secondary tantalum plants identified in this study.

The present uses of tantalum stem from three important properties; a high melting point, resistance to corrosive agents, and dielectric properties. Tantalum mill products such as sheet into corrosion resistant chemical equipment. heat are made exchangers, reaction vessels and other equipment that can high temperatures and severe acid environments. withstand Tantalum has been used in surgical applications such as surgical implants and suture wire because it is inert to body fluids and tissue. The electronic industry has many applications for tantalum, primarily in capacitors. Tantalum capacitors provide higher volumetric capacitance efficiency than other capacitor materials, and function well at high and low temperatures. Alloying with tantalum produces alloys with good high temperature strength that have applications in aerospace products. Tantalum alloys also have favorable fabricating characteristics.

DESCRIPTION OF SECONDARY TANTALUM PRODUCTION

Secondary tantalum production methods vary from time to time and from plant to plant primarily because of the different raw materials that may be used in the process. Basically, acid leaching is used to dissolve metal impurities in the raw material leaving behind an upgraded tantalum product. In the following sections the variations to this fundamental operation will be described. Figure III-1 (page 4550) presents a flow diagram for the secondary tantalum production processes.

RAW MATERIALS

The plants presently producing secondary tantalum use three types of raw materials: alloy scrap, scrap tantalum-bearing electrical components, and tantalum-bearing sludge. Stamping operations are the primary source of raw materials for one plant which uses alloy scrap as a raw material.

ACID LEACHING

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Different types of acid leaching processes are used depending on the raw material being processed. Scrap alloy is immersed in a nitric acid bath and the spent acid is discharged in a batch flow. This waste stream contains high concentrations of dissolved metals, the constituents depending upon the make-up of the scrap alloy.

SECONDARY TANTALUM SUBCATEGORY SECT - III

Electrical components such as capacitors are acid leached in a batch process. The acid which may be a combination of hydrochloric, nitric and sulfuric, is poured into a rotating digestor filled with tantalum-bearing scrap. Leaching in the digestor continues until the acid is spent. usually about two days per batch. The spent acid is then discharged as a waste stream along with rinse water, and the digestor is filled with fresh acid. This procedure is repeated up to 15 times depending on the amount of materials to be leached from each batch of scrap.

Recovering tantalum from sludge involves a series of acid leaches, and may sometimes include caustic leaching depending on the materials present in the sludge. The recovery of other valuable materials from tantalum-bearing sludges may determine how the leaching steps are done. After each leaching step, the more pure tantalum is filtered to separate it from the dissolved impurities. If the filtrate does not contain recoverable material it is combined with rinse water and discharged as a wastewater stream.

One plant practicing acid leaching of tantalum-bearing sludges reported the use of a wet air pollution control device on the leaching vessel. A wastewater stream is discharged from this device.

WASHING AND RINSING

The washing and rinsing operations employed are dependent upon the form of the product recovered and on the desired purity of the end product. When processing scrap tantalum alloys, a water rinse is employed on a continuous once-through basis. Rinsing is performed after completing the acid leaching step for a batch of scrap alloy. Water use is determined by the amount of product to be washed. The tantalum alloy scrap wash water, combined with acid leachate constitutes a waste stream.

Tantalum metal product derived from electrical components is also rinsed and acid washed in batch operations. An acid wash is used to polish the purified metal powder by removing any residual surface oxides from the metal. A water rinse follows which washes away any residual acid. This waste stream may be pretreated to recover dissolved tantalum before routing to wastewater treatment.

Water is used to wash the upgraded tantalum solids to remove all acid prior to the next leaching operation. Filtering is used to retain the upgraded tantalum while separating the liquid fraction. If the filtrate contains recoverable materials, both the filtrate and the wash water are routed to the recovery operation. Otherwise, both flows are routed to wastewater treatment.

PROCESS WASTEWATER SOURCES

Although two fundamental processes are involved in secondary tantalum production, variations in raw materials and specific procedures require that the process wastewater sources be subdivided as follows:

- (a) Tantalum alloy leach and rinse,
- (b) Capacitor leach and rinse
- (c) Tantalum sludge leach and rinse,
- (d) Tantalum powder acid wash and rinse and
- (e) Leaching wet air pollution control.

OTHER WASTEWATER SOURCES

There may be other wastewater streams associated with the secondary tantalum subcategory. These wastewaters may include stormwater runoff, and maintenance and cleanup water. These waste streams are not considered as a part of this rulemaking. EPA believes that the flows and pollutant loadings associated with these waste streams are insignificant relative to the waste streams selected and are best handled by the appropriate permit authority on a case-by-case basis under authority of Section 403 of the Clean Water Act.

AGE, PRODUCTION AND PROCESS PROFILE

Figure III-2 (page 4551) shows the location of the three secondary tantalum plants currently operating in the United States. All three plants are located in the eastern part of the United States.

Table III-1 (page 4548) shows the relative age and discharge status of the tantalum plants. All three plants are direct dischargers and all were built prior to World War II. From Table III-2 (page 4548) it can be seen that secondary tantalum production is not done on a large scale, and production at each plant varies.

Table III-3 (page 4549) provides a summary of the number of plants generating wastewater for the waste streams associated with various processes and the number of plants with the process.

TABLE III-1

INITIAL OPERATING YEAR (RANGE) SUMMARY OF PLANTS IN THE SECONDARY TANTALUM SUBCATEGORY BY DISCHARGE TYPE

Initial Operating Year (Range) (Plant Age in Years)					
Type of Plant	1983- 1944 (1-40)	1943- 1934 (41-50)	1933- 1904 (51-80)	Before 1904 <u>(80+)</u>	Total
Direct	0	1	1	1	,3
Indirect	0	0	0	0	0
Zero	0	· O	0	0	0
Total	0	1	1	1	.3

TABLE III-2

PRODUCTION RANGES FOR THE SECONDARY TANTALUM SUBCATEGORY

Tantalum Production Ranges for 1982

Plant Type	0-8 tons/yr	9-17 tons/yr	18-26 tons/yr	Total Number of Plants
Direct	1	1	1	3
Indirect	0	0	0	0
Zero	0	0	0	0

Table III-3

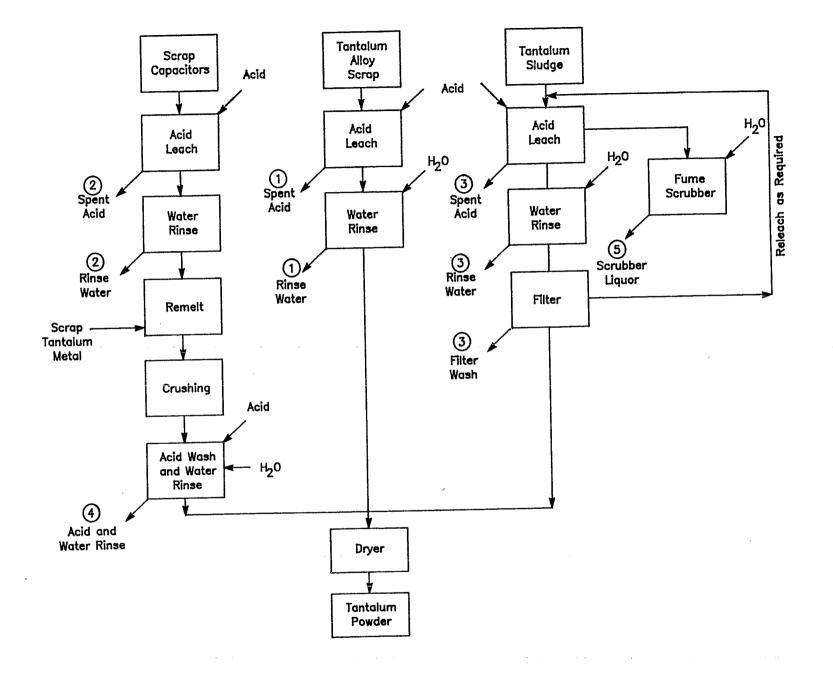
SUMMARY OF SECONDARY TANTALUM SUBCATEGORY PROCESSES AND ASSOCIATED WASTE STREAMS

Process	Number of Plants With Process	Number of Plants Reporting Generation of Wastewater*
Acid Leaching	3	
Tantalum Alloy Leach and Kinse	1	1
Capacitor Leach and Rinse	1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 -	1 1
Tantalum Sludge Leach and Rinse	1	1
Leaching Wet Air Pollution Control	- 1	1
Washing and Rinsing	1	
Tantalum Powder Acid Wash and Rinse	1	1

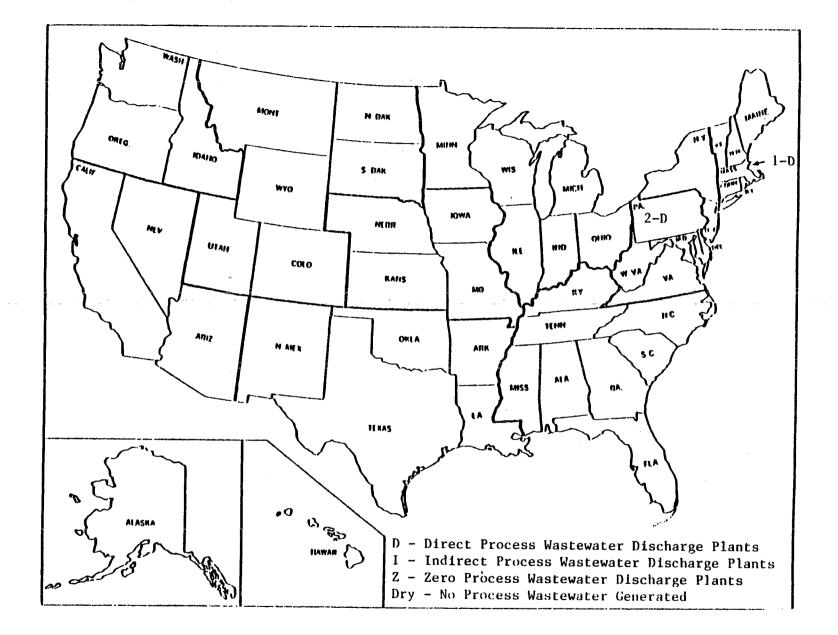
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^{*}Through reuse or evaporation practices, a plant may "generate" wastewater from a particular process but not discharge it.



SECONDARY TANATALUM SUBCATEGORY SECT I III



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Figure III-2

GEOGRAPHIC LOCATIONS OF THE SECONDARY TANTALUM SUBCATEGORY PLANTS

SECONDARY TANTALUM SUBCATEGORY SECT - III

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SECTION IV

SUBCATEGORIZATION

This section summarizes the factors considered during designation of the related subdivisions of the secondary tantalum subcategory. Production normalizing parameters for each subdivision are also discussed.

FACTORS CONSIDERED IN SUBDIVIDING THE SECONDARY TANTALUM SUBCATEGORY

The factors listed previously for general subcategorization were each evaluated when considering subdivision of the secondary tantalum subcategory. In the discussion that follows, the factors will be described as they pertain to this particular subcategory.

The rationale for considering further subdivision of the secondary tantalum subcategory is based primarily on differences in the production processes and raw materials used. Within this subcategory, different operations are performed which may or may have a water use or discharge, and which may require the not establishment of separate effluent limitations. While secondary tantalum is still considered a single subcategory, a more thorough examination of the production processes has illustrated the need for limitations and standards based on specific flow allowances for the following subdivisions:

- (a) Tantalum alloy leach and rinse,
- (b) Capacitor leach and rinse,

(c) Tantalum sludge leach and rinse,

- (d) Tantalum powder acid wash and rinse, and
- (e) Leaching wet air pollution control.

The following discussion is intended to clarify and support the reasons given above for subdividing the secondary tantalum subcategory.

Secondary tantalum production can be generally described as consisting of acid leaching of raw materials followed by water rinsing and drying of the final tantalum powder product. Variations of this process are due to differences in raw materials. Such factors account for the first three subdivisions listed above. A discussion of each subdivision follows.

Tantalum alloy scrap may be used as a raw material. This scrap is generated in forming operations in which a tantalum-containing alloy is rolled and stamped. The remaining metal skeleton is the raw material for the acid leaching process. Leaching is done in a batch mode by immersing the scrap in acid. Spent acid is discharged as a waste stream.

Scrap electrical components containing tantalum may be used as a

SECONDARY TANTALUM SUBCATEGORY SECT - IV

raw material. These components, predominantly capacitors, may have plastic parts and be diverse in composition. Successive acid leaching of batches of raw material is done in rotating digestors. The spent acid is discharged after each cycle. The process is complete when all impurities have been leached away, leaving only the tantalum product.

Tantalum-bearing sludges may used as a raw material for secondary tantalum recovery. The sludge is mixed with acid and acidsoluable impurities are leached away. The residual solids contain upgraded tantalum and are filtered to separate them from the spent acid prior to subsequent purification.

The fourth subdivision arises from an additional purification step that one plant includes in its production operations. After remelting leached tantalum powder and solid tantalum scrap to separate impurities, the tantalum product is crushed to a powder and washed with acid. The acid wash removes surface oxides from the tantalum powder resulting in a higher grade powder product.

The fifth subdivision accounts for wet scrubbers used to control emissions from acid leaching operations. Acid fume generation from the leaching of raw materials is a function of the type of processes used by individual plants. In this subcategory, only one plant uses a wet scrubber to control acid fumes from leaching operations.

OTHER FACTORS

The other factors considered in this evaluation were shown to be inappropriate bases for subdivision. Air pollution control methods, treatment costs, and total energy requirements are functions of the selected subcategorization factors -- metal product, raw materials, and production processes. Therefore, they are not independent factors and do not affect the subcategorization which has been applied. Certain other factors such as plant age, plant size, and the number of employees, were also evaluated and determined to be inappropriate for use as bases for subdivision of this subcategory.

PRODUCTION NORMALIZING PARAMETERS

As discussed previously, the effluent limitations and standards developed in this document establish mass limitations on the discharge of specific pollutant parameters. To allow these regulations to be applied to plants with various production capacities, the mass of pollutant discharged must be related to a unit of production. This factor is known as the production normalizing parameter (PNP).

In general, for each production process which has a wastewater associated with it, the actual mass of tantalum product or intermediate produced will be used as the PNP. Thus, the PNPs for the five subdivisions or building blocks are as follows:

SECONDARY TANTALUM SUBCATEGORY SECT - IV

leaching

produced

Building block

PNP

- Tantalum alloy leach and rinse
- tantalum powder produced

tantalum powder produced

tantalum powder produced from

equivalent pure tantalum powder

- 2. Capacitor leach and rinse
- 3. Tantalum sludge leach and rinse
- Tantalum powder acid wash and rinse
- 5. Leaching wet air pollution control

equivalent pure tantalum powder produced

Equivalent pure tantalum powder production was selected as the PNP for subdivisions three and five because the product of leaching tantalum-bearing sludge contains approximately 25 to 30 percent tantalum. Equivalent pure tantalum refers to the weight of tantalum contained in the product.

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SECONDARY TANTALUM SUBCATEGORY SECT - V

SECTION V

WATER USE AND WASTEWATER CHARACTERISTICS

This section describes the characteristics of the wastewaters associated with the secondary tantalum subcategory. Water use and discharge rates are explained and then summarized in tables at the end of this section. Data used to characterize the wastewaters are presented. Finally, the specific source, water use and discharge flows, and wastewater characteristics for each separate wastewater source are discussed.

The two principal data sources were used are data collection portfolios (dcp) and field sampling results. Data collection portfolios completed for each secondary tantalum plant contain information regarding wastewater flows and production levels.

In order to quantify the pollutant discharge from secondary tantalum plants, a field sampling program was conducted. Α complete list of the pollutants considered and a summary of the techniques used in sampling and laboratory analyses are included in Section V of Vol. I. Because the analytical standard for TCDD judged to be too hazardous to be made generally available, was samples were never analyzed for this pollutant. Samples were also not analyzed for asbestos. There is no reason to expect that or asbestos would be present in TCDD nonferrous metals manufacturing wastewater. One plant was selected for sampling in the secondary tantalum subcategory. In general, the samples were analyzed for the two classes of pollutants, priority metal pollutants and criteria pollutants (which include both conventional and nonconventional pollutants). Samples were not analyzed for priority organic pollutants because there is no reason to believe that organic pollutants would be present in wastewaters generated by the secondary tantalum subcategory.

After proposal, EPA gathered additional wastewater sampling data for two of the subdivisions in this subcategory. These data were acquired through a self-sampling program which was undertaken at the specific request of the Agency. The data include analyses for the toxic metals antimony, beryllium. cadmium, chromium, copper, lead, nickel, silver, thallium, and zinc. The data also include analyses for the nonconventional pollutant tantalum. These data support the assumptions which EPA had made concerning the presence and concentrations of pollutants in those subdivisions where we did not have analytical data for specific pollutants. For this reason, the selection of pollutant parameters for limitation in this subcategory (Section VI) has not been revised based on this new data.

As described in Section IV of this supplement, the secondary tantalum subcategory has been divided into five subdivisions or wastewater sources, so that the promulgated regulation contains mass discharge limitations and standards for five unit processes discharging process wastewater. Differences in the wastewater characteristics associated with these subdivisions are to be expected. For this reason, wastewater streams corresponding to each subdivision or building block are addressed separately in the discussions that follow. These wastewater sources are:

- (a) Tantalum alloy leach and rinse,
- (b) Capacitor leach and rinse,
- (c) Tantalum sludge leach and rinse,
- (d) Tantalum powder acid wash and rinse, and
- (e) Leaching wet air pollution control.

WASTEWATER FLOW RATES

Data supplied by dcp responses were evaluated, and two flow-toproduction ratios, water use and wastewater discharge flow, were calculated for each stream. The two ratios are differentiated by the flow value used in calculation. Water use is defined as the volume of water or other fluid required for a given process per mass of tantalum product and is therefore based on the sum of recycle and make-up flows to a given process. Wastewater flow discharged after pretreatment or recycle (if these are present) is used in calculating the production normalized flow -- the volume of wastewater discharged from a given process to further treatment, disposal, or discharge per mass of tantalum produced. Differences between the water use and wastewater fl ws associated with a given stream result from recycle, evaporation, and carry-over on the product. The production values used in calculation correspond to the production normalizing parameter, PNP, assigned to each stream, as outlined in Section IV. As an example, tantalum powder acid wash and rinse wastewater flow is related to the production of tantalum powder. As such, the discharge rate is expressed in liters of acid wash and rinse wastewater per metric ton of tantalum powder produced (gallons of acid wash and rinse water per ton of tantalum powder).

The production normalized discharge flows were compiled and statistically analyzed by stream type. These production normalized water use and discharge flows are presented by subdivision in Tables V-1 through V-5 (pages 4563 - 4564). Where appropriate, an attempt was made to identify factors that could account for variations in water use and discharge rates. These variations are discussed later in this section by subdivision. A similar analysis of factors affecting the wastewater flows is presented in Sections IX, X, XI, and XII where representative BPT, BAT, NSPS, and pretreatment flows are selected for use in calculating the effluent limitations.

The water use and discharge rates shown do not include nonprocess wastewater, such as rainfall runoff and noncontact cooling water.

WASTEWATER CHARACTERISTICS DATA

Data used to characterize the various wastewaters associated with secondary tantalum production come from two sources -- data collection portfolios and analytical data from field sampling

trips.

DATA COLLECTION PORTFOLIOS

In the data collection portfolios, the secondary tantalum plants that discharge wastewater were asked to specify the presence of priority pollutants in their wastewater. Of the three secondary tantalum plants, one did not respond to this portion of the questionnaire. None of the plants responding to the questionnaire reported the presence of priority organic pollutants. The responses for the priority metals and cyanide are summarized below:

Pollutant	Known Present	Believed Present (Based on Raw Materials and <u>Process Chemicals Used)</u>
Antimony	0	1
Arsenic	0	ō
Beryllium	0	Ő
Cadmium	0	1
Chromium	2	· 0
Copper	1	1
Cyanide	0	0
Lead	0	0
Mercury	1	0
Nickel	1	1
Selenium	0	0
Silver	0	1
Thallium	0	0
Zinc	1	1

FIELD SAMPLING DATA

In order to quantify the concentrations of pollutants present in wastewater from secondary tantalum plants, wastewater samples were collected at a single plant, which represents one-third of the secondary tantalum plants in the United States, and accounts for 44 percent of all secondary tantalum production. A diagram indicating the sampling sites and contributing production processes is shown in Figure V-1 (page 4586).

Raw wastewater data are summarized in Tables V-6 through V-9 (pages 4565 - 4577). Analytical results for capacitor leach and rinse and tantalum powder acid wash and rinse waste streams are given in Tables V-6 and V-7, respectively. Table V-8 shows analytical results from samples taken from a holding tank (sump) into which the spent acid stream flows, as well as other streams from unrelated plant processes. Table V-9 shows data from a similar type of holding tank (sump) into which the acid wash and water rinse stream flows, along with other unrelated waste streams. Finally, Table V-10 (page 4581) shows the analytical results of the samples taken of the final effluent, after having been treated and prior to discharge to a surface stream. Note that the stream numbers listed in the tables correspond to those

given in the plant sampling site diagram, Figure V-1 (page 4586). Where no data are listed for a specific day of sampling, the wastewater samples for the stream were not collected. Sampling was only done for two classes of pollutants: priority metal pollutants, and criteria pollutants which include both conventional and nonconventional pollutants.

The data tables include some samples measured at concentrations considered not quantifiable. The base-neutral extractable, acid extractable, and volatile organics generally are considered not quantifiable at concentrations equal to or less than 0.010 mg/l. Below this concentration, organic analytical results are not quantitatively accurate; however, the analyses are useful to indicate the presence of a particular pollutant. The pesticide fraction is considered not quantifiable at concentrations equal to or less than 0.005 mg/l.

It should be noted that the detection limits shown on the data tables for priority metals and conventional and nonconventional pollutants are not the same in all cases as the published detection limits for these pollutants by the same analytical methods. The detection limits used were reported with the analytical data and hence are the appropriate limits to apply to the data. Detection limit variation can occur as a result of a number of laboratory-specific, equipment-specific. and daily operator-specific factors. These factors can include day-to-day differences in machine calibration, variation in stock solutions, and variation in operators.

The statistical analysis of data includes some samples measured concentrations considered not quantifiable. For data at considered as detected but below quantifiable concentrations, a value of zero is used for averaging. Priority organic, nonconventional, and conventional pollutant data reported with a "less than" sign are considered as detected, but not further quantifiable. A value of zero is also used for averaging. If a pollutant is reported as not detected, it is assigned a value of zero in calculating the average. Finally, priority metal values reported as less than a certain value were considered as not quantifiable, and consequently were assigned a value of zero in the calculation of the average.

Finally, appropriate source water concentrations are presented with the summaries of the sampling data. The method by which each sample was collected is indicated by number, as follows:

1	one-time grab	4	8-hour automatic composite
2	manual composite during intermit-	5	24-hour manual composite
	tent process operation	6	24-hour automatic composite
3	8-hour manual composite		

WASTEWATER CHARACTERISTICS AND FLOWS BY SUBDIVISION

The secondary tantalum subcategory has been divided into five subdivisions. The wastewater characteristics and discharge rates

corresponding to each subdivision are described separately in this section.

TANTALUM ALLOY LEACH AND RINSE

Spent acid is generated when batches of scrap tantalum alloy from forming operations are acid leached to recover tantalum. After leaching, the residual tantalum metal is rinsed with water to remove excess acid. The rinse water is discharged to treatment along with the spent acid. Table V-1 (page 4563) shows the production normalized water use and discharge rates for the tantalum alloy leach and rinse waste stream in liters per metric ton of tantalum powder produced.

Although the two component waste streams in this subdivision were not directly sampled, it is expected that their respective waste characteristics will be similar to two waste streams from this subcategory that were sampled. Spent acid from tantalum alloy leaching is expected to have similar characteristics to capacitor leaching wastewater which was sampled. Consequently, treatable concentrations of toxic metals including copper, nickel, and zinc are expected, as well as low pH. Wastewater characteristics for capacitor leaching wastewater are shown in Table V-6 (page 4565).

The water rinse component of the tantalum alloy leach and rinse waste stream is expected to have similar pollutant concentrations to the tantalum powder acid wash and rinse waste stream. Table V-7 (page 4569) shows the analytical data for this waste stream. The water rinse portion of the waste stream is expected to be acidic and contain treatable concentrations of toxic metals including copper and nickel.

CAPACITOR LEACH AND RINSE

This waste stream is composed of spent acid generated by leaching scrap capacitors and other electrical components that contain tantalum. The acid leaches away all impurities leaving behind a residue of tantalum metal powder. The spent acid is discharged to treatment along with rinse water used to remove excess acid. Table V-2 shows production normalized flows in liters per metric ton of tantalum metal produced.

Table V-6 summarizes the field sampling data for this waste stream. From this data, it can be seen that capacitor leaching spent acid can be characterized by an acidic pH, treatable concentrations of toxic metals including copper, lead, and zinc, and treatable concentrations of suspended solids.

TANTALUM SLUDGE LEACH AND RINSE

This wastewater stream arises from the upgrading of tantalumbearing sludge. The sludge is leached with acid, rinsed with water and the residual solids separated from the liquid phase by filtration. Successive leaching operations are performed until the desired level of purity is attained. The production normalized flow is shown in Table V-3, (page 4563) in liters per metric ton of equivalent pure tantalum powder produced.

At proposal, specific wastewater characteristics data for this stream were not available. Following proposal, sampling data for this subdivision were acquired through a self-sampling effort initiated at the specific request of the Agency. These data are presented in Table V-11 (page 4585) and show show treatable concentrations of toxic and nonconventional metals.

TANTALUM POWDER ACID WASH AND RINSE

Acid washing is used to polish the powdered tantalum by removing surface oxides that may have formed in the previous stages of the production process. The subsequent water rinse is used to wash the acid from the powder prior to drying. Table V-4 (page 4564) shows the production normalized flows for this operation in liters per metric ton of tantalum powder produced.

A sample of this wastewater was taken after residual tantalum was recovered by ammonium hydroxide precipitation. This step is assumed not to affect constituents in the waste stream other than and ammonia. Because of the raw materials tantalum and production operations used by this plant, there is no reason to expect that treatable concentrations of ammonia are generated in the acid wash and water rinse process. Therefore, the concentrations of ammonia presented in Table V-7 (page 4569) are assumed to be caused by addition of ammonium hydroxide in the tantalum recovery opera-tion, and can be disregarded when characterizing the acid wash and water rinse waste stream. The pH may also be modified by the addition of ammonium hydroxide, but the data in Table V-7 show that the pH of the waste stream after tantalum recovery is still acidic. Accounting for these differences, the acid wash and water rinse waste stream is characterized by treatable concentrations of copper and nickel, and having an acidic pH.

LEACHING WET AIR POLLUTION CONTROL

One plant reported using a wet scrubber to control hydrochloric acid fumes generated in acid leaching operations. The scrubber liquor blowdown is discharged to treatment. Table V-5 (page 4564) shows the production normalized flows for the scrubbing operation in liters per metric ton of equivalent pure tantalum powder produced.

Following proposal, sampling data for this subdivision were acquired through a self-sampling effort at the specific request of the Agency. These data presented in table V-11 (page 4585) show treatable concentrations of toxic and nonconventional metals, thus corroborating the data used at proposal.

SECONDARY TANTALUM SUBCATEGORY SECT - V

TABLE V-1

WATER USE AND DISCHARGE RATES FOR TANTALUM ALLOY LEACH AND RINSE

(1000 1/kkg of tantalum powder produced)

Plant Code	Percent Recycle	Production Normalized <u>Water</u> <u>Use</u>	Production Normalized <u>Discharge</u> Flow	
1145	0	230.6	230.6	

TABLE V-2

WATER USE AND DISCHARGE RATES FOR CAPACITOR LEACH AND RINSE

(1000 1/kkg of tantalum powder produced)

<u>Plant</u> Code	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge Flow	
1089	0	20.2	20.2	
	· · · · · · · · · · · · · · · · · · ·			

TABLE V-3

WATER USE AND DISCHARGE RATES FOR TANTALUM SLUDGE LEACH AND RINSE

(1000 1/kkg of equivalent pure tantalum powder produced)

Percent	Normalized	Normalized
Plant Code Recycle	Water Use	Discharge Flow

205.3

1146

4563

SECONDARY TANTALUM SUBCATEGORY SECT - V

TABLE V-4

WATER USE AND DISCHARGE RATES FOR TANTALUM POWDER ACID WASH AND RINSE

(1000 1/kkg of tantalum powder produced)

<u>Plant</u> <u>Code</u>	Percent Recycle	Production Normalized Water Use	Production Normalized <u>Discharge</u> <u>Flow</u>
1089	0	0.350	0.350

TABLE V-5

WATER USE AND DISCHARGE RATES FOR LEACHING WET AIR POLLUTION CONTROL

(1000 1/kkg of equivalent pure tantalum powder produced)

		Production	Production
	Percent	Normalized	Normalized
<u>Plant</u> Code	Recycle	Water Use	Discharge Flow

1146

58.8

Table V-6

SECONDARY TANTALUM SAMPLING DATA WASTE ACID FROM TANTALUM LEACHING RAW WASTEWATER

	Pollutant	Stream Code	Sample Typet	<u>Conc</u>	entrations (mg/l) Day 1 Day 2	Day B
<u>Toxic</u>	Pollutants	2				Day Day
114.	antimony	464	1	<0.01	1.0	/ ¬
115.	arsenic	464	1	<0.01	<0.1	0.02 ANT
117.	beryllium	464	1	<0.005	<0.5	47 0.02 TANTALUM
118.	cadmium	464	1	<0.02	<2	
119.	chromium (total)	464	1	<0.02	<2	<2 BCAJ
120.	copper	464	1	<0.05	49,200 17,	<pre><2 SUBCATEGORY </pre>
122.	lead	464	1	<0.05	15,900 6,	100 ^R
123.	mercury	464	1	<0.0002	0.0037	0.001
124.	nickel	464	1	0.5	3,580 1,	890 H
125.	selenium	464	1	<0.01		<0.02
126.	silver	464	1	<0.01	30.0	50
127.	thallium	464	1	<0.01	0.075	0.02
128.	zinc	464	1	0.08	8,060 2,	810

4565

Table V-6 (Continued)

SECONDARY TANTALUM SAMPLING DATA WASTE ACID FROM TANTALUM LEACHING RAW WASTEWATER

Pollutant	Stream Code	Sample Typet	Conc Source	entrations (mg/l) Day 1 Day 2	Day 3 Day 3	
Nonconventional Pollutants					DARY	•
acidity	464	1	<1	390	10	
alkalinity	464	1 .	40	21	19 ТАМТАLUM 21 <10	
aluminum	464	1	0.2	<10	<10 ^L	
ammonia nitrogen	464	1	2.0	0.02	0.14 gg	5
barium	464	1	<0.05	<5	<5 CAT	;]
boron	464	1	<0.1	<10	0.14 SUBCATEGORY)) 1
calcium	464	1	25.7	20	30 ^R	i
chemical oxygen demand (COD)	464	1	110	2,000	760 ស្ព	2
chloride	464	1	11	<1	760 SE EC く1 日	Ē
cobalt	464	1	<0.05	<5	۱ 5 <	~ 7
fluoride	464	1	0.64	0.06	0.28	
	464	1	<0.5	60 5	,550	
iron magnesium	464	1	4.5	<10	<10	

Table V-6 (Continued)

SECONDARY TANTALUM SAMPLING DATA WASTE ACID FROM TANTALUM LEACHING RAW WASTEWATER

	Pollutant	Stream Code	Sample Typet	Conc Source		g/1) y 2
	Nonconventional Pollutants (Continued)					
	manganese	464	1	<0.05	2,380	1,060
	molybdenum	464	. 1	<0.05		<5
45		464	1	0.82	620,000	210,000
67	sodium	464	1	4.5	<10	<10
	sulfate	464	1	590	700	290
	tin	464	1	<0.05	65	8,960
	titanium	464	1	<0.05	<5	<5
	total organic carbon (TOC)	464	1	<1	12	27
	total solids (TS)	464	1	250	210,000	7,800
	vanadium	464	1	<0.05	<5	<5
	yttrium	464	1	<0.05	<5	<5

Table V-6 (Continued)

SECONDARY TANTALUM SAMPLING DATA WASTE ACID FROM TANTALUM LEACHING RAW WASTEWATER

	Stream	Sample	Conc	.) 전	
Pollutant	Code	<u>Typet</u>	Source	Day 1 Day	$\frac{2}{2}$ Day $\frac{3}{2}$
Conventional Pollutants					DARY
oil and grease	464	1	<1	<1	
total suspended solids (TSS)	464	- 1	- 19	80,000	1,200 ^M HA
pH (standard units)	464	1	7.60	2.26	3.00 g

tSample Type Code: 1 - One-time grab

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SUBCATEGORY

Table V-7

SECONDARY TANTALUM SAMPLING DATA FILTRATE FROM NH4OH PRECIPITATION OF TANTALUM RAW WASTEWATER

	Pollutant	Stream <u>Code</u>	Sample Typet	<u>Conc</u> Source	centrations (mg Day 1 Day	
<u>Toxic</u>	Pollutants					
114.	antimony	466	1	<0.01	<0.01	
115.	arsenic	466	1	<0.01	0.02	ANT
117.	beryllium	466	1	<0.005	<0.005	TANTALUM
118.	cadmium	466	1	<0.02	<0.02	
119.	chromium (total)	466	1	<0.02	<0.02	SUBCATEGORY
120.	copper	466	1	<0.05	4.65	EGO
122.	lead	466	1	<0.05	<0.5	RY
123.	mercury	466	1	<0.002	0.0004	Ŋ
124.	nickel	466	1	0.5	2.45	SECT
125.	selenium	466	1	<0.01	<0.01	ו ל
126.	silver	466	- 1	<0.01	<0.01	
127.	thallium	466	1	<0.01	0.02	
128.	zinc	466	1	0.08	0.12	

Table V-7 (Continued)

SECONDARY TANTALUM SAMPLING DATA FILTRATE FROM NH4OH PRECIPITATION OF TANTALUM RAW WASTEWATER

Pollutant	Stream <u>Code</u>	Sample Typet	Concentrations (mg/1) Source Day 1 Day 2 Day 3 Concentrations (mg/1)
Nonconventional Pollutants			AKY
acidity	466	1	
alkalinity	466	1	<1 <1 HAR
aluminum	466	1	
ammonia nitrogen	466	1	2.0 1,600
barium	466	1	<0.05 <0.05
boron	466	1	2.0 1,600 5 <0.05
calcium	466	1	25.7 0.7
chemical oxygen demand (COD)	466	1 .	110 230
chloride	466	1	11 410
cobalt	466	1	<0.05 <0.5
fluoride	466	1	0.64 530
iron	466	Î	<0.5 1.3
magnesium	466	1	4.5 1.6

Table V-7 (Continued)

SECONDARY TANTALUM SAMPLING DATA FILTRATE FROM NH4OH PRECIPITATION OF TANTALUM RAW WASTEWATER

	Pollutant	Stream	Sample		ncentrations		
	Forracant	Code	Typet	Source	Day 1	Day 2	Day 3
	Nonconventional Pollutants (Continued)						
	manganese	466	1	<0.05	<0.05		
	molybdenum	466	1	<5	0.1		
2	phosphate	466	1	0.82	59		
7	sodium	466	1	4.5	14		
	sulfate	466	1	590 1	,600		
	tin	466	1	<0.05	<0.5		
	titanium	466	1	<0.05	<0.05		
	total organic carbon (TOC)	466	1	<1	<1		
	total solids (TS)	466	1	250	880		
	vanadium	466	1	<0.05	<0.05		
	yttrium	466	1	<0.05	<0.05		

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Table V-7 (Continued)

SECONDARY TANTALUM SAMPLING DATA FILTRATE FROM NH4OH PRECIPITATION OF TANTALUM RAW WASTEWATER

	Stream	Sample	Con	centration	s (mg/l)	B
Pollutant	<u>Code</u>	Typet	Source	Day 1	<u>Day 2</u>	Day 30
Conventional Pollutants						DARY
oil and grease	466	1	<1	3		
total suspended solids (TSS)	466	1	19	528		TANTALUM
pH (standard units)	466	1	7.60	4.76		LUM

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Table V-8

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 2 EFFLUENT RAW WASTEWATER

Pollutant		Stream	Sample	Con	Concentrations (mg/l)			
	Pollucant	_Code_	Typet	Source	Day 1	Day 2	Day 3	- ON
Toxic	e Pollutants							SECONDARY
114.	antimony	479	3	<0.01	<0.01	<0.1	<0.05	
115.	arsenic	479	3	<0.01	<0.01	<0.09	<0.1	NTA
117.	beryllium	479	3	<0.005	<0.005	<0.05	<0.05	TANTALUM
118.	cadmium	479	3	<0.02	0.04	0.8	0.6	SUB
119.	chromium (total)	479	3	<0.02	0.32	6.2	6.4	CATI
120.	copper	479	3	<0.05	71.7	950	518	SUBCATEGORY
122.	lead	479	3	<0.05	2.35	73.5	<10	Ą
123.	mercury	479	3	<0.0002	<0.0002	<0.0002	0.0011	E
124.	nickel	479	3	0.5	145 2	,130	955	SECT
125.	selenium	479	3	<0.01	<0.01	<0.1	<0.01	י ל
126.	silver	479	3	<0.01	0.04	1	0.4	
127.	thallium	479	3	<0.01	<0.05		<0.5	
128.	zinc	479	3	0.08	15.4	89	25.4	

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RY TANTALUM SUBCATEGO

Table V-8 (Continued)

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 2 EFFLUENT RAW WASTEWATER

	Pollutant	Stream Code	Sample Typet	Con- Source	centrat: Day	lons (mg/l Day 2	
	Nonconventional Pollutants						
	acidity	479	3	<1	10	1,400	130
	alkalinity	479	3	40	<1	<1	<1
•	aluminum	479	3	0.2	0.4	12	6
7	ammonia nitrogen	479	3	2.0	0.45	2.3	1.7
	barium	479	3	<0.05	<0.05	<0.5	<0.5
	boron	479	3	<0.1	<1	<10	<2
	calcium	479	3	25.7	31.6	41	38
	chemical oxygen demand (COD)	479	3	110	56	4,300	6,140
	chloride	479	3	11	310	8,300	4,200
	cobalt	479	3	<0.05	5.75	75.5	37
	fluoride	479	3	0.64	0.21	0.72	0.52
	iron	* 479	3	<0.5	150	3,440	1,300
	magnesium	479	3	4.5	7.4	8	8
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Table V-8 (Continued)

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 2 EFFLUENT RAW WASTEWATER

Pollutant	Stream Code	Sample Typet	Cor Source	centratio Day 1	o <mark>ns (mg</mark> Day		SECONDARY
Nonconventional Pollutants (Continued)	,						NDAI
manganese	479	3	<0.05	7.5	59.5	28	
molybdenum	479	3	<0.05	<0.5	<5	<0.5	ANTI
phosphate	479	3	0.82	860 16	5,000	6,600	TANTALUM
sodium	479	3	4.5	9.3	33	610	
sulfate	479	3	590	470	93	91	SUBCATEGORY
tin	479	3	<0.05	6.15	9	<0.5	EGO
titanium	479	3	<0.05	<0.05	<0.5	<0.5	ЯY
total organic carbon (TOC)	479	3	<1	13	47	78	ល
total solids (TS)	479	3	250 2	,000 25	,000	13,000	SECT
vanadium	479	3	<0.05	<0.05	0.5	<0.5	י ע
yttrium	479	3	<0.05	<0.05	<0.5 [.]	<0.5	

Table V-8 (Continued)

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 2 EFFLUENT RAW WASTEWATER

	Stream	Sample	Concentrations (mg/1))
Pollutant	Code	Typet	Source	Day 1	Day 2	
Conventional Pollutants						UAKI
oil and grease	479	1	<1	110	3	14 1.920
total suspended solids (TSS)	479	3	19	630	160	1,920
pH (standard units)	479	3	7.60	3.83	1.84	2.26

1. Mune Cale. 1 One time and

tSample Type Code: 1 - One-time grab
3 - 8-hour manual composite

SECONDARY TANTALUM SUBCATEGORY SECT -

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Table V-9

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 3 EFFLUENT RAW WASTEWATER

Dollutant		Stream	Sample	<u>Concentrations (mg/l)</u>					
		Pollutant	Code	<u>Typet</u>	Source	Day 1	Day 2	Day 3	CON
	Toxic	Pollutants							SECONDARY
	114.	antimony	482	3	<0.01	<0.05	<0.01	<0.01	
	115.	arsenic	482	3	<0.01	<0.1	<0.1	<0.1	TANTALUM
	117.	beryllium	482	3	<0.005	<0.05	<0.005	<0.005	LUM
ר נ נ	118.	cadmium	482	3	<0.02	0.4	<0.02	<0.02	SUE
	119.	chromium (total)	482	3	<0.02	6.4	<0.02	<0.02	CAT
	120.	copper	482	3	<0.05	533	0.8	0.35	SUBCATEGORY
	122.	lead	482	3	<0.05	<10	<0.05	<0.05	ΥΥ.
	123.	mercury	482	3	<0.0002	<0.0002	<0.0002	<0.0002	IS
	124.	nickel	482	3	0.5	972	<0.05	<0.0002 0.1	CT
	125.	selenium	482	3	<0.01	<0.01	<0.01	<0.01	י ל
	126.	silver	482	3	<0.01	<0.01	<0.01	<0.01	
	127.	thallium	482	3	<0.01	<0.01	<0.01	<0.01	
	128.	zinc	482	3	0.08	26	0.06	0.06	

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Table V-9 (Continued)

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 3 EFFLUENT RAW WASTEWATER

Pollutant	Stream Code	Sample Typet	Conc Source	entratic Day 1	ons (mg/1) Day 2	Day 3	SECONDARY
Nonconventional Pollutants							DARS
acidity	482	3	<1	150	<1	<1	
alkalinity	482	3	40	<1	213	61	TANTALUM
aluminum	482	3	0.2	.6	0.2	0.2	LUM
ammonia nitrogen	482	3	2.0	0.08	0.43	0.14	SUE
barium	482	3	<0.05	<0.5	<0.05	<0.05	CAT
boron	482	3	<0.1	<2	<0.1	<0.1	SUBCATEGORY
calcium	482	3	25.7	38	22	28.5	Я
chemical oxygen demand (COD)	482	3	110 1,	,600	35	19.6	SH
chloride	482	3	11	25	46	20	SECT
cobalt	482	3	<0.05	37.5	<0.05	<0.05	י ל
fluoride	482	3	0.64	0.39	0.85	0.21	
iron	482	3	<0.5 1	,340	0.4	0.15	
magnesium	482	3	4.5	8	5.3	7.9	

Table V-9 (Continued)

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 3 EFFLUENT RAW WASTEWATER

Dollartont	Stream	Sample						
Pollutant	Code	<u>Typet</u>	Source	Day 1	<u>Day 2</u>	Day 3	ğ	
Nonconventional Pollutants (Continued)) .						SECONDARY	
manganese	482	3	<0.05	28.5	<0.05	<0.05	•	
molybdenum	482	3	<0.05	<0.5	<0.05	<0.05	TANTALUM	
phosphate	482	3	0.82	<0.732	<0.732	1.3	LUM	
sodium	482	3	4.5	620	247	7.4	SUB	
sulfate	482	3	590	520	1,300	590	CATE	
tin	482	3	<0.05	1.5	<0.1	<0.05	SUBCATEGORY	
titanium	482	3	<0.05	<0.5	<0.05	<0.05	К	
total organic carbon (TOC)	482	3	<1	<1	38	14	SECT	
total solids (TS)	482	3	250	280	920 1	,500	C I I	
vanadium	482	3	<0.05	<0.5	<0.05	<0.05	- ~	
yttrium	482	3	<0.05	<0.5	<0.05	<0.05		

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Table V-9 (Continued)

SECONDARY TANTALUM SAMPLING DATA SUMP NUMBER 3 EFFLUENT RAW WASTEWATER

Stream Code	Sample Typet	Cone Source	centration Day 1	ns (mg/1) Day 2	Day 3	SECONDARY
						IDAR
482	1	<1	<1	<1	<1	-
482	3	19	29	52	<1	TANTALUM
482	. 3	7.60	3.28	10.5	8.05	LUM
						SUBCATEGORY
	<u>Code</u> 482 482	<u>Code Typet</u> 482 1 482 3	<u>Code Typet Source</u> 482 1 <1 482 3 19	<u>Code Typet Source Day 1</u> 482 1 <1 <1 482 3 19 29	Code Typet Source Day 1 Day 2 482 1 <1	Code Typet Source Day 1 Day 2 Day 3 482 1 <1

tSample Type Code: 1 - One-time grab
3 - 8-hour manual composite

Table V-10

SECONDARY TANTALUM SAMPLING DATA TREATED EFFLUENT

	Pollutant	Stream <u>Code</u>	Sample Typet	Conc Source	centration Day 1	<u>ns (mg/1)</u> Day 2	Day 3	SE
<u>Toxic</u>	e Pollutants							CONI
114.	antimony	473	6	<0.01	0.13	<0.01	<0.01	SECONDARY
115.	arsenic	473	6	<0.01	<0.01	<0.01	<0.01	TAN
117.	beryllium	473	6	<0.005	<0.005	<0.005	<0.005	TANTALUM
118.	cadmium	473	6	<0.02	<0.02	<0.02	<0.02	
119.	chromium (total)	473	6	<0.02	<0.02	<0.02	<0.02	SUBC
120.	copper	473	6	<0.05	0.35	0.7	<0.05	SUBCATEGORY
122.	lead	473	6	<0.05	<0.5	<0.3	<0.5	30RY
123.	Mercury	473	6	<0.0002	<0.0002	<0.0002	<0.000	2
124.	nickel	473	6	0.5	5.35	4.6	1.85	SECT
125.	selenium	473	6	<0.01	<0.01	<0.01	<0.01	i I
126.	silver	473	6	<0.01	<0.02	0.02	<0.01	4
127.	thallium	473	6	<0.01	<0.02	<0.02	<0.02	
128.	zinc	473	6	0.08	1.46	0.74	0.16	

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Table V-10 (Continued)

SECONDARY TANTALUM SAMPLING DATA TREATED EFFLUENT

N 11 A 446	Stream Code	Sample Typet	Con Source	<u>centratio</u> Day 1	ns (mg/1 Day 2		IS
Pollutant		Typer	Dourod	<u> </u>			
Nonconventional Pollutants							CONDARY
acidity	473	6	<1	<1	<1	<1	
alkalinity	473	6	40	13	16	19	TAN
aluminum	473	6	0.2	<0.1	<0.1	<0.1	TANTALUM
ammonia nitrogen	473	6	2.0	1.2	1.3	1.4	
barium	473	6	<0.05	<0.05	<0.05	<0.05	UBC
boron	473	6	<0.1	0.2	0.1	0.2	SUBCATEGORY
calcium	473	6	25.7	745	762	808	; ORY
chemical oxygen demand (COD)	473	6	110	160	120	130	
chloride	473	6	11 1	,100	240	1,200	SECT
cobalt	473	6	<0.05	4.25	2.9	1.3	i i
	473	. 6	0.64	5.8	4.9	0.67	4
fluoride	473	6	<0.5	0.15	1.35	<0.15	
iron				2	1.9	1.8	
magnesium	473	6	4.5	L	1.7		

Table V-10 (Continued)

SECONDARY TANTALUM SAMPLING DATA TREATED EFFLUENT

Pollutant	Stream Code	Sample Typet	Source	oncentrat e Day		
Nonconventional Pollutants (Continued)						
manganese	473	6	<0.05	2.5	2.35	1.45
molybdenum	473	6	<0.05	<0.05	<0.05	<0.05
phosphate	473	6	0.82	15	8.4	39
sodium	473	6	4.5	14.7	15.7	21.2
sulfate	473	6	590	670	1,200	7,600
tin	473	6	<0.05	<0.05	<0.1	<0.1
titanium	473	6	<0.05	<0.05	<0.05	<0.05
total organic carbon (TOC)	473	6	<1	15	8.4	39
total solids (TS)	473	6	250	4,000	3,500	3,400
vanadium	473	6	<0.05	<0.05	<0.05	<0.05
yttrium	473	6	<0.05	<0.05	<0.05	<0.05

SECONDARY TANTALUM SUBCATEGORY SECT I 4

Table V-10 (Continued)

SECONDARY TANTALUM SAMPLING DATA TREATED EFFLUENT

	Stream	Sample	Concentrations (mg/1)				
Pollutant	Code	<u>Typet</u>	Source	Day 1	Day 2	Day	$\overline{}$
Conventional Pollutants							CONDARY
oil and grease	473	1	<1		<1	14	
total suspended solids (TSS)	473	6	19	16	5	<1	TANI
pH (standard units)	473	6	7.60	5.70	5.58	6.47	TANTALUM
							• •
							SUBCATEGORY
							EGO
							RY

tSample Type Code: 1 - One-time grab 6 - 24-hour automatic composite

SECT ı <

TABLE V-11

SECONDARY TANTALUM SAMPLING DATA RAW WASTEWATER DATA FROM SELF-SAMPLING PROGRAM

POLLUTANT	CONCENTRATION					
Sample No.	88143	88144				
Toxic Pollutants						
<pre>114. antimony 117. beryllium 118. cadmium</pre>	0.059 <0.050 0.120	<0.100 <0.050 0.600				
119. chromium 120. copper 122. lead	0.528 <0.100 <0.200	1.010 <0.100 <0.200				
124. nickel 126. silver 128. zinc	<0.200 1.600 <0.050	<0.200 <0.100 <0.050				
Nonconventional Pollutants						
aluminum cobalt iron	<0.500 <0.500 0.420	0.500 <0.500 1.200				
manganese molybdenum tantalum	<0.050 7.920 12.000	<0.050 <5.000				
tin titanium vanadium	5.000 <0.200 <1.000	<50.000 <20.000 10.300				

NOTES:

Sample No. 88143 = Tantalum Sludge Leach and Rinse Sample No. 88144 = Leaching Wet Air Pollution Control



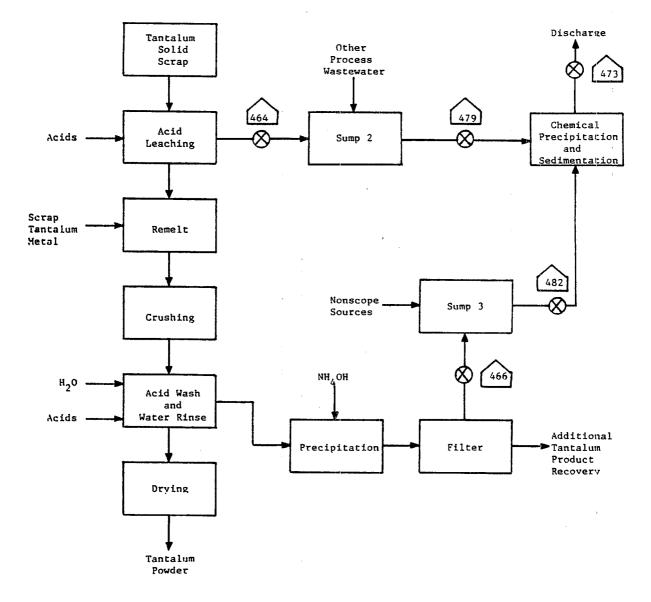


Figure V-1 SAMPLING SITES AT SECONDARY TANTALUM PLANT A

SECONDARY TANTALUM SUBCATEGORY SECT - VI

SECTION VI

SELECTION OF POLLUTANT PARAMETERS

This section examines the chemical analysis data presented in Section V and discusses the selection or exclusion of pollutants for potential limitation. The discussion that follows presents and briefly discusses selection of conventional and limitations. nonconventional pollutants for effluent Also described is the analysis that was performed to select or exclude priority pollutants for further consideration for limitations and standards. Pollutants will be considered for limitation if they are present in concentrations treatable by the technologies considered in this analysis. The treatable concentrations used the priority metals were the long-term performance values for by chemical precipitation. sedimentation, achievable and The treatable concentrations used for the priority filtration. organics were the long-term performance values achievable by carbon adsorption.

CONVENTIONAL AND NONCONVENTIONAL POLLUTANT PARAMETERS SELECTED

This study examined samples from the secondary tantalum subcategory for two conventional pollutant parameters (total suspended solids and pH) and several nonconventional pollutant parameters.

The conventional and nonconventional pollutants or pollutant parameters selected for limitation in this subcategory are:

tantalum total suspended solids (TSS) pH

Based on an examination of the production processes employed in the secondary tantalum subcategory, it is expected that concentrations of tantalum could be present in the wastewater generated in this subcategory. For this reason, tantalum is selected for limitation in this subcategory.

TSS concentrations ranging from 29 to 80,000 mg/l were observed in the raw waste samples analyzed for this study. All the concentrations are well above the 2.6 mg/l treatable concentration. Most of the specific methods used to remove toxic metals do so by converting these metals to precipitates, and toxic-metal-containing precipitates these should not be discharged. Meeting a limitation on total suspended solids helps ensure that removal of these precipitated toxic metals has been effective. For these reasons, total suspended solids are selected for limitation in this subcategory.

The pH values observed during this study ranged from 1.8 to 10.5. Seven of the values were equal to or less than 4.8, and one other was outside the 7.5 to 10.0 range considered desirable for discharge to receiving waters. Many deleterious effects are caused by extreme pH values or rapid changes in pH. Also, effective removal of toxic metals by precipitation requires careful control of pH. Since pH control within the desirable limits is readily attainable by available treatment, pH is selected for limitation in this subcategory.

TOXIC PRIORITY POLLUTANTS

The frequency of occurrence of the priority pollutants in the raw wastewater samples taken is presented in Table VI-1 (page 4591). Table VI-1 is based on the raw wastewater data from streams 464 and 466 (see Section V). These data provide the basis for the categorization of specific pollutants, as discussed below. Treatment plant and sump effluent samples were not considered in the frequency count. Note that sampling was not done for any priority organic pollutants.

TOXIC POLLUTANTS NEVER DETECTED

The toxic pollutants listed in Table VI-2 (page 4592) were never detected in any raw wastewater samples from this subcategory. Therefore, they are not selected for consideration in establishing limitations.

TOXIC POLLUTANTS NEVER FOUND ABOVE THEIR ANALYTICAL QUANTIFICATION CONCENTRATION

The toxic pollutants listed below were never found above their analytical quantification concentration in any raw wastewater samples from this subcategory; therefore, these pollutants are not selected for consideration in establishing limitations.

117. beryllium
118. cadmium
119. chromium
125. selenium
127. thallium

PRIORITY POLLUTANTS PRESENT BELOW CONCENTRATIONS ACHIEVABLE BY TREATMENT

The pollutants listed below are not selected for consideration in establishing limitations because they were not found in any raw wastewater samples from this subcategory above concentrations considered achievable by existing or available treatment technologies. These pollutants are discussed individually following the list.

```
115. arsenic
123. mercury
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Arsenic was detected above its quantification concentration of 0.010 mg/l in two of the samples analyzed. The detected values were both 0.02 mg/l. The treatable concentration for arsenic is

0.34 mg/l, much higher than any of the analyzed samples indicate. Therefore, arsenic is not selected for limitation.

Mercury was detected above its quantification concentration of 0.0001 mg/l in all three samples analyzed. The analysis showed a range of 0.0004 mg/l to 0.0037 mg/l, well below the treatable concentration for mercury of 0.036 mg/l. For this reason, mercury is not selected for limitation.

TOXIC POLLUTANTS SELECTED FOR FURTHER CONSIDERATION IN ESTABLISHING LIMITATIONS AND STANDARDS

The priority pollutants listed below are selected for further consideration in establishing limitations and standards for this subcategory. The toxic pollutants selected for further consideration for limitation are each discussed following the list.

114. antimony
120. copper
122. lead
124. nickel
126. silver
128. zinc

Antimony was detected below the quantification concentration of 0.100 mg/l in one sample (<0.01 mg/l). The other two samples indicated treatable concentrations of antimony of 1.0 mg/l and 47 mg/l. The treatable concentration for antimony is 0.47 mg/l. Therefore, antimony is selected for further consideration for limitation.

Copper was discovered above treatable concentrations in three samples analyzed. The treatable concentration for copper is 0.39 mg/l. The concentrations detected were 4.65 mg/l, 17,100 mg/l, and 49,200 mg/l. Since these waste streams contain substantial concentrations of treatable copper, this metal is selected for further consideration for limitation.

Lead was detected above treatable concentrations in two samples analyzed. The treatable concentration for lead is 0.08 mg/l. The sample concentrations showed 6,100 mg/l and 15,900 mg/l of lead. Because of such large lead concentrations in the waste streams, lead is selected for further consideration for limitation.

Nickel was detected above treatability (0.22 mg/l) in all three of the samples analyzed. Detected concentrations were found to be 2.45 mg/l, 1,890 mg/l, and 3,580 mg/l. Therefore, nickel is selected for further consideration for limitation.

Silver was detected below the quantification concentration of 0.02 mg/l in one of the three samples that were analyzed. The sample registered <0.01 mg/l. The remaining two samples were both above the treatable concentration of 0.07 mg/l. Concentrations of 30.0 and 50 mg/l of silver were detected, and

thus silver is selected for further consideration for limitation.

Zinc was detected above the treatable concentration of 0.23 mg/l in two samples showing 2,810 mg/l and 8,060 mg/l of zinc. The remaining sample was below treatability indicating only 0.12 mg/l zinc. However, because of the significant quantities found in two samples, zinc is selected for further consideration for limitation.

Table VI-1

FREQUENCY OF OCCURRENCE OF PRIORITY POLLUTANTS SECONDARY TANTALUM SUBCATEGORY RAW WASTEWATER

Pollutant	Analytical Quantification Concentration (mg/l)(a)	Treatable Concentra- tion (mg/1)(b)	Number of Streams Analyzed	Number of Samples Analyzed	ND	Detected Below Quantification Concentration	Detected Below Treat- able Concen- tration	Detected Above Treat- able Concen- tration
114. antimony	0.100	0.47	2	3		1	0	2
115. arsenic	0.010	0.34	2	3		1	ž	2
117. beryllium	0.010	0.20	2	3		à	ñ	0
118. cadmium	0.002	0.049	2	3		3	0	0
119. chromium	0.005	0.07	2	3		3	0	
120. copper	0.009	0.39	2	ĩ		ñ	0	0
121. cyanide (c)	0.02	0.047	ō			0	U	3
122. lead	0.020	0.08	2	3		1	٥	n
123. mercury	0.0001	0.036	- 2	ĩ		0	2	2
124. nickel	0.005	0.22	2	3		0	5	U
125. selenium	0.01	0.20	2	2		2	0	. 0
126. silver	0.02	0.07	$\overline{2}$	จิ		1	0	0
127. thallium	0.100	0.34	2	3		3	0	4
128, zinc	0.050	0.23	2	3		õ	1	2

(a) Analytical quantification concentration was reported with the data (see Section V).

(b) Treatable concentrations are based on performance of chemical precipitation, sedimentation, and filtration.

(c) Analytical quantification concentration for EPA Method 335.2, Total Cyanide Methods for Chemical Analysis of Water and Wastes, EPA 600/4-79-020, March 1979.

TABLE VI-2

TOXIC POLLUTANTS NEVER DETECTED

acenaphthene* 1. 2. acrolein* acrylonitrile* 3. 4 benzene* 5. benzidine* carbon tetrachloride (tetrachloromethane)* 6. chlorobenzene* 7. 8. 1,2,4-trichlorobenzene* 9. hexachlorobenzene* 1,2-dichloroethane* 10. 1,1,1-trichloroethane* 11. 12. hexachloroethane* 1,1-dichloroethane* 13. 1,1,2-trichloroethane* 14. 1,1,2,2-tetrachloroethane* 15. 16. chloroethane* bis (chloromethyl) ether (DELETED)* 17. bis (2-chloroethyl) ether* 18. 2-chloroethyl vinyl ether (mixed)* 19. 2-chloronaphthalene* 20. 2,4,6-trichlorophenol* 21. parachlorometa cresol* 22. chloroform (trichloromethane)* 23. 24. 2-chlorophenol* 25. 1,2-dichlorobenzene* 26. 1,3-dichlorobenzene* 1,4-dichlorobenzene* 27. 3,3'-dichlorobenzidine* 28. 1,1-dichloroethylene* 29. 1,2-trans-dichloroerhylene* 30. 2,4-dichlorophenol* 31. 1,2-dichloropropane* 32. 1,2-dichloropropylene (1,3-dichloropropene)* 33. 2,4-dimethylphenol* 34. 2,4-dinitrotoluene* 35. 2,6-dinitrotoluene* 36. 1,2-diphenylhydrazine* 37. ethylbenzene* 38. 39. fluoranthene* 4-chlorophenyl phenyl ether* 40. 4-bromophenyl phenyl ether* 41. bis(2-chloroisopropyl) ether* 42. 43. bis(2-choroethoxy) methane* 44. methylene chloride (dichloromethane)* methyl chloride (chloromethane)* 45. methyl bromide (bromomethane)* 46. 47. bromoform (tribromomethane)* dichlorobromomethane* 48. 49. trichlorofluoromethane (DELETED)*

TABLE VI-2 (Continued)

TOXIC POLLUTANTS NEVER DETECTED

50. dichlorodifluoromethane (DELETED)*

51. chlorodibromomethane*

52. hexachlorobutadiene*

53. hexachlorocyclopentadiene*

54. isophorone*

55. naphthalene*

56. nitrobenzene*

57. 2-nitrophenol*

58. 4-nitrophenol*

59. 2,4-dinitrophenol*

60. 4,6-dinitro-o-cresol*

61. N-nitrosodimethylamine*

62. N-nitrosodiphenylamine*

63. N-nitrosodi-n-propylamine*

64. pentachlorophenol*

65. phenol*

66. bis(2-ethylhexyl) phthalate*

67. butyl benzyl phthalate*

68. di-n-butyl phthalate*

69. di-n-octyl phthalate*

70. diethyl phthalate*

71. dimethyl phthalate*

72. benzo (a)anthracene (1,2-benzanthracene)*

73. benzo (a)pyrene (3,4-benzopyrene)*

74. 3,4-benzofluoranthene*

75. benzo(k)fluoranthane (11,12-benzofluoranthene)*

76. chrysene*

77. acenaphthylene*

78. anthracene*

79. benzo(ghi)perylene (1,11-benzoperylene)*

80. fluorene*

81. phenanthrene*

82. dibenzo (a,h)anthracene (1,2,5,6-dibenzanthracene)*

83. indeno (1,2,3-cd)pyrene (w,e,-o-phenylenepyrene)*

84. pyrene*

85. tetrachloroethylene*

86. toluene*

87. trichloroethylene*

88. vinyl chloride (chloroethylene)*

89. aldrin*

90. dieldrin*

91. chlordane (technical mixture and metabolites)*

92. 4,4'-DDT*

93. 4,4'-DDE(p,p'DDX)*

94. 4,4'-DDD(p,p TDE)*

95. Alpha-endosulfan*

96. Beta-endosulfan*

97. endosulfan sulfate*

99. endrin aldehyde*

TABLE VI-2 (Continued)

TOXIC POLLUTANTS NEVER DETECTED

*We did not analyze for these pollutants in samples of raw wastewater from this subcategory. These pollutants are not believed to be present based on the Agency's best engineering judgement which includes consideration of raw materials and process operations.

SECONDARY TANTALUM SUBCATEGORY SECT - VII

SECTION VII

CONTROL AND TREATMENT TECHNOLOGIES

The preceding sections of this supplement discussed the sources, flows, and characteristics of the wastewaters from secondary tantalum plants. This section summarizes the description of these wastewaters and indicates the treatment technologies which are currently practiced in the secondary tantalum subcategory for each wastewater stream. Secondly, this section presents the control and treatment technology options which were examined by the Agency for possible application to the secondary tantalum subcategory.

CURRENT CONTROL AND TREATMENT PRACTICES

Control and treatment technologies are discussed in Section VII of Vol. I and the pollutant concentrations achievable with these treatment technologies are presented in table VII-21 (page 248) that volume. The basic principles of these technologies and of the applicability to wastewaters similar to those found in this subcategory are presented there. This section presents a summary of the control and treatment technologies that are currently being applied to each of the sources generating wastewater in subcategory. As discussed in Section V, this wastewater associated with the secondary tantalum subcategory is characterized by the presence of the toxic metal pollutants and suspended solids. This analysis is supported by the raw (untreated) wastewater data presented for specific sources as well as combined waste streams in Section V. Generally, these pollutants are present in each of the wastewater streams at concentrations above the levels achievable by treatment, and these wastewaters are commonly combined for treatment. Construction of one wastewater treatment system for combined treatment allows plants to take advantage of economies of scale and in some instances to combine streams of different alkalinity to reduce treatment chemical requirements. All three plants in this subcategory currently have combined wastewater treatment systems including chemical precipitation and sedimentation. The options selected for consideration for BPT, BAT, NSPS, and on combined pretreatment based treatment of compatible wastewaters are summarized toward the end of this section.

TANTALUM ALLOY LEACH AND RINSE

Tantalum recovery from alloy scrap is accomplished by immersing the scrap into an acid bath and leaching away all impurities. Water rinsing of the tantalum powder residue follows the leaching operation and is designed to remove residual acid from the tantalum powder before drying. The spent acid, along with the once-through rinse water is discharged to lime and settle treatment. Polymer addition is used to aid flocculation and settling. The final effluent is discharged directly.

CAPACITOR LEACH AND RINSE

Tantalum is recovered from scrap capacitors and other electrical components by successive batch leaching. The spent acid contains high concentrations of dissolved metals and also some suspended solids. The wastewater from this operation and other wastewater streams is treated using chemical precipitation and sedimentation. The treated effluent is discharged to a surface water.

TANTALUM SLUDGE LEACH AND RINSE

Tantalum recovery from sludge requires successive leaching filtering and washing operations. The filtrate and wash water may be sent to a metal by-product recovery process prior to being discharged to the wastewater treatment facility. After treatment consisting of chemical precipitation and sedimentation, the effluent is discharged.

TANTALUM POWDER ACID WASH AND RINSE

One plant washes tantalum powder with acid and subsequently rinses it with water prior to the final drying of the product. The acid wash is designed to remove surface oxides from the tantalum powder, and the water rinse removes residual acid before drying. The acid and water stream are combined and pretreated with ammonium hydroxide to precipitate dissolved tantalum. After filtering the precipitate, the filtrate is routed to the treatment system for treatment consisting of chemical precipitation and sedimentation. It is then discharged to a surface water.

LEACHING WET AIR POLLUTION CONTROL

A wet scrubber may be used to control emissions of acid fumes generated by acid leaching operations. A caustic solution is circulated in the scrubber, and a 92 percent recycle rate is presently practiced. The scrubber discharge is combined with other wastewater streams and treated by chemical precipitation and sedimentation. The final effluent is discharged to a surface stream.

CONTROL AND TREATMENT OPTIONS

The Agency examined two control and treatment technology options that are applicable to the secondary tantalum subcategory. The options selected for evaluation represent applicable end-of-pipe treatment technologies.

Examination of the waste streams in this subcategory shows that no in-process flow reduction, beyond that presently being practiced, is achievable. Therefore, options including flow reduction were not considered.

OPTION A

Option A for the secondary tantalum subcategory requires control and treatment technologies to reduce the discharge of wastewater pollutant mass.

The Option A treatment scheme consists of chemical precipitation and sedimentation technology. Specifically, lime or some other alkaline compound is used to precipitate metal ions as metal hydroxides. The metal hydroxides and suspended solids settle out and the sludge is collected. Vacuum filtration is used to dewater sludge.

OPTION C

Option C for the secondary tantalum subcategory consists of all control and treatment requirements of Option A (chemical precipitation and sedimentation) plus multimedia filtration technology added at the end of the Option A treatment scheme. Multimedia filtration is used to remove suspended solids, including precipitates of metals, beyond the concentration attainable by gravity sedimentation. The filter suggested is of gravity, mixed-media type, although other forms of filters, the rapid sand filters or pressure filters would perform such as satisfactorily. The addition of filters also provides consistent removal during periods of time in which there are rapid increases in flows or loadings of pollutants to the treatment system.

SECONDARY TANTALUM SUBCATEGORY SECT - VII

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SECONDARY TANTALUM SUBCATEGORY SECT - VIII

SECTION VIII

COSTS, ENERGY, AND NONWATER QUALITY ASPECTS

This section presents a summary of compliance costs for the secondary tantalum subcategory and a description of the treatment options and subcategory-specific assumptions used to develop these estimates. Together with the estimated pollutant removal performance presented in Section X of this supplement, these cost estimates provide a basis for evaluating each regulatory option. These cost estimates are also used in determining the probable economic impact of regulation on the subcategory at different pollutant discharge levels. In addition, this section addresses nonwater quality environmental impacts of wastewater treatment and control alternatives, including air pollution, solid wastes, and energy requirements, which are specific to the secondary tantalum subcategory.

TREATMENT OPTIONS FOR EXISTING SOURCES

As discussed in Section VII, two treatment options have been developed and considered in promulgating limitations and standards for the secondary tantalum subcategory. These options are summarized below and schematically presented in Figures X-1 and X-2 (pages 4624 and 4625).

OPTION A

The Option A treatment scheme consists of chemical precipitation and sedimentation technology.

OPTION C

Option C for the secondary tantalum subcategory consists of all control and treatment requirements of Option A (chemical precipitation and sedimentation) plus multimedia filtration technology added at the end of the Option A treatment scheme.

COST METHODOLOGY

A detailed discussion of the methodology used to develop the compliance costs is presented in Section VIII of the General Development Document. Plant-by-plant compliance costs have been estimated for the nonferrous metals manufacturing category and are presented in the administrative record supporting this regulation. Compliance cost estimates developed for the promulgated regulation are presented in Table VIII-1 (page 4602) for the direct dischargers in this subcategory. These cost estimates are equivalent to those developed for the proposed regulation.

Each of the general assumptions used to develop compliance costs has been previously discussed. No subcategory-specific

assumptions were used in developing compliance costs for the secondary tantalum subcategory.

ENERGY REQUIREMENTS

Energy requirements for Option A are estimated at 37,000 kwh/yr, and for Option C the estimated requirement is 39,000 kwh/yr. Option C energy requirements increase over those for Option A because filtration is being added as an end-of-pipe treatment technology. Since recycle of scrubber liquor is already in place in this subcategory, energy requirement savings resulting from flow reduction measures are not reflected in this analysis. Both options represent about two percent of a typical plant's energy usage. It is therefore concluded that the energy requirements of the treatment options considered will not have a significant impact on total plant energy consumption.

SOLID WASTE

Sludge generated in the secondary tantalum subcategory is due to the precipitation of metal hydroxides and carbonates using lime or other chemicals. Sludges associated with the secondary tantalum subcategory will necessarily contain quantities of toxic metal pollutants. Wastes generated by secondary metal industries can be regulated as hazardous. However, the Agency examined the solid waste that would be generated at secondary nonferrous metals manufacturing plants by the suggested treatment technologies and believes they are not hazardous wastes under the Agency's regulations implementing Section 3001 of the Resource Conservation and Recovery Act. The one exception to this is solid wastes generated by cyanide precipitation. These sludges are expected to be hazardous and this judgment was included in None of the non-cyanide wastes are this study. listed specifically as hazardous. Nor are they likely to exhibit а characteristic of hazardous waste. This judgment is made based on the recommended technology of chemical precipitation and filtration. By the addition of a small excess of lime during treatment, similar sludges, specifically toxic metal bearing sludges, generated by other industries such as the iron and steel industry passed the Extraction Procedure (EP) toxicity test. See 40 CFR \$261.24. Thus, the Agency believes that the wastewater sludges will similarly not be EP toxic if the recommended technology is applied.

Although it is the Agency's view that solid wastes generated as a result of these guidelines are not expected to be hazardous, generators of these wastes must test the waste to determine if the wastes meet any of the characteristics of hazardous waste (see 40 CFR 262.11).

If these wastes should be identified or are listed as hazardous, they will come within the scope of RCRA's "cradle to grave" hazardous waste management program, requiring regulation, from the point of generation to point of final disposition. EPA's generator standards would require generators of hazardous nonferrous metals manufacturing wastes to meet containerization, labeling, recordkeeping, and reporting requirements; if plants dispose of hazardous wastes off-site, they would have to prepare a manifest, which would track the movement of the wastes from the generator's premise to a permitted off-site treatment, storage, or disposal facility. See 40 CFR 262.20 45 FR 33142 (May 19, 1980), as amended at 45 FR 86973 (December 31, 1980). The transporter regulations require transporters of hazardous wastes to comply with the manifest system to assure that the wastes are delivered to a permitted facility. See 40 CFR 263.20 45 FR 33151 (May 19, 1980), as amended at 45 FR 86973 (December 31, 1980). Finally, RCRA regulations establish standards for hazardous waste treatment, storage, and disposal facilities allowed to receive See 40 CFR Part 464 46 FR 2802 (January 12, 1981), such wastes. 47 FR 32274 (July 26, 1982).

Even if these wastes are not identified as hazardous, they still must be disposed of in compliance with the Subtitle D open dumping standards, implementing 4004 of RCRA. See 44 FR 53438 (September 13, 1979). It is estimated that the secondary tantalum subcategory will generate 386 metric tons of sludge per year when implementing the promulgated BPT treatment technology. The Agency has calculated as part of the costs for wastewater treatment the cost of hauling and disposing of these wastes.

AIR POLLUTION

There is no reason to believe that any substantial air pollution problems will result from implementation of chemical precipitation, sedimentation, and multimedia filtration. These technologies transfer pollutants to solid waste and are not likely to transfer pollutants to air.

TABLE VIII-1

COST OF COMPLIANCE FOR THE SECONDARY TANTALUM SUBCATEGORY DIRECT DISCHARGERS

(March, 1982 Dollars)

Option	Total Required <u>Capital</u> Cost	Total <u>Annual</u> Cost
A	6,462	58,854
С	13,474	63,466

4602

SECTION IX

BEST PRACTICABLE CONTROL TECHNOLOGY CURRENTLY AVAILABLE

This section defines the effluent characteristics attainable through the application of best practicable control technology currently available (BPT). BPT reflects the existing performance by plants of various sizes, ages, and manufacturing processes within the secondary tantalum subcategory, as well as the established performance of the recommended BPT systems. Particular consideration is given to the treatment already in place at plants within the data base.

The factors considered in identifying BPT include the total cost of applying the technology in relation to the effluent reduction benefits from such application, the age of equipment and facilities involved, the manufacturing processes used, nonwater quality environmental impacts (including energy requirements), and other factors the Administrator considers appropriate. In general, the BPT level represents the average of the existing performances of plants of various ages, sizes, processes, or common characteristics. Where existing performance other is uniformly inadequate, BPT may be transferred from a different subcategory or category. Limitations based on transfer technology are supported by a rationale concluding that of the technology is, indeed, transfera-ble, and a reasonable prediction that it will be capable of achieving the prescribed effluent limits. BPT focuses on end-of-pipe treatment rather than process changes or internal controls, except where such practices are common industry practice.

TECHNICAL APPROACH TO BPT

The Agency studied this subcategory to identify the processes used, the wastewaters generated, and the treatment processes Information was collected from industry using data installed. collection portfolios, and specific plants were sampled and the wastewaters analyzed. In making technical assessments of data, reviewing manufacturing processes, and assessing wastewater options, both indirect direct freatment technology and dischargers have been considered as a single group. An examination of plants and processes did not indicate any process differences based on the type of discharge, whether it be direct or indirect.

As explained in Section IV the secondary tantalum subcategory has been subdivided into five potential wastewater sources. Since the water use, discharge rates, and pollutant characteristics of each of these wastewaters is potentially unique, effluent limitations are developed for each of the five subdivisions.

For each of the subdivisions, a specific approach was followed for the development of BPT mass limitations. The first

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requirement to calculate these limitations is to account for production and flow variability from plant to plant. Therefore, a unit of production or production normalizing parameter (PNP) was determined for each waste stream which could then be related to the flow from the process to determine a production normalized flow. Selection of the PNP for each process element is discussed Each plant within the subcategory was Section IV. then in analyzed to determine which subdivisions were present, the specific flow rates generated for each subdivision, and the specific production normalized flows for each subdivision. This analysis is discussed in detail in Section V. Nonprocess wastewaters such as rainfall runoff and noncontact cooling water are not considered in the analysis.

Production normalized flows for each subdivision were then analyzed to determine the flow to be used as part of the basis for BPT mass limitations. The selected flow (sometimes referred to as the BPT regulatory flow or BPT discharge rate) reflects the water use controls which are common practices within the category. The BPT regulatory flow is based on the average of all applicable data. Plants with normalized flows above the average may have to implement some method of flow reduction to achieve the BPT limitations.

The second requirement to calculate mass limitations is the set of concentrations that are achievable by application of the BPT level of treatment technology. Section VII discusses the various control and treatment technologies which are currently in place for each wastewater source. In most cases, the current control and treatment technologies consist of chemical precipitation and sedimentation (lime and settle technology) and a combination of reuse and recycle to reduce flow.

Using these regulatory flows and the achievable concentrations, the next step is to calculate mass loadings for each wastewater source or subdivision. This calculation was made on a stream-bystream basis, primarily because plants in this subcategory may perform one or more of the operations in various combinations. The mass loadings (milligrams of pollutant per kilogram of production - mg/kg) were calculated by multiplying the BPT regulatory flow (l/kkg) by the concentration achievable by the BPT level of treatment technology (mg/l) for each pollutant parameter to be limited under BPT. These mass loadings are published in the Federal Register and in 40 CFR Part 421 as the effluent limitations.

The mass loadings which are allowed under BPT for each plant will be the sum of the individual mass loadings for the various wastewater sources which are found at particular plants. Accordingly, all the wastewater generated within a plant may be combined for treatment in a single or common treatment system, but the effluent limitations for these combined wastewaters are based on the various wastewater sources which actually contribute to the combined flow. This method accounts for the variety of combinations of wastewater sources and production processes which may be found at secondary tantalum plants.

The Agency usually establishes wastewater limitations in terms of mass rather than concentration. This approach prevents the use of dilution as a treatment method (except for controlling pH). The production normalized wastewater flow (l/kkg) is a link between the production operations and the effluent limitations. The pollutant discharge attributable to each operation can be calculated from the normalized flow and effluent concentration achievable by the treatment technology and summed to derive an appropriate limitation for each plant.

INDUSTRY COST AND POLLUTANT REMOVAL ESTIMATES

In balancing costs in relation to pollutant removal estimates, EPA considers the volume and nature of existing discharges, the volume and nature of discharges expected after application of BPT, the general environmental effects of the pollutants, and the cost and economic impacts of the required pollution control level. The Act does not require or permit consideration of water quality problems attributable to particular point sources or industries, or water quality improvements in particular water quality bodies. Accordingly, water quality considerations were not the basis for selecting the proposed or promulgated BPT.

The methodology for calculating pollutant removal estimates and plant compliance costs is discussed in Section X. Table X-1 (page 4618) shows the pollutant removal estimates for each treatment option for direct dischargers. Compliance costs for direct dischargers are presented in Table X-2 (page 4619).

BPT OPTION SELECTION

The technology basis for the proposed and promulgated BPT limitations is Option A, chemical precipitation and sedimentation technology to remove metals and solids from combined wastewaters and to control pH. These technologies are demonstrated and economically achievable since they are already in place at all of the direct dischargers in this subcategory. The BPT treatment scheme is presented in Figure IX-1 (page 4612).

Implementation of the promulgated BPT limitations will remove annually an estimated 26,268 kilograms of toxic metals and 20,079 kilograms of TSS from raw wastewater generated by the secondary tantalum industry. Projected capital and annual costs are \$6,462 and \$58,854 (1982 dollars), respectively, to achieve the promulgated BPT limitations.

WASTEWATER DISCHARGE RATES

A BPT discharge rate is calculated for each subdivision based on the average of the flows of the existing plants as determined from analysis of data collection portfolios. The discharge rate is used with the achievable treatment concentrations to determine BPT effluent limitations. Since the discharge rate may be

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source, for each wastewater different separate production normalized discharge rates for each of the five wastewater sources are discussed below and summarized in Table IX-1. The discharge rates are normalized on a production basis by relating the amount of wastewater generated to the mass of the intermediate product which is produced by the process associated with the waste scream in question. These production normalizing parameters, or PNPs. are also listed in Table IX-1.

Section V of this document further describes the discharge flow rates and presents the water use and discharge flow rates for each plant by subdivision in Tables V-1 through V-5 (pages 4564 - 4565).

TANTALUM ALLOY LEACH AND RINSE

The BPT wastewater discharge rate for tantalum alloy leach and rinse is 230,600 l/kkg (55,261 gal/ton) of tantalum powder produced based on the only water use rate reported. This rate is allocated only for those plants which leach tantalum alloy scrap material by immersion into an acid bath and use water to rinse the tantalum powder product before it is dried. Water use and wastewater discharge rates are presented in Table V-1 (page 4564).

CAPACITOR LEACH AND RINSE

The BPT wastewater discharge rate for capacitor leach and rinse is 20200 l/kkg (4841 gal/ton) of tantalum powder produced. This rate is allocated only for those plants whose raw material is scrap electrical components containing tantalum. Recovery of tantalum powder is performed by successive leachings of the raw material. The production normalized flows for this subdivision are presented in Table V-2 (page 4564).

TANTALUM SLUDGE LEACH AND RINSE

The proposed and promulgated BPT wastewater discharge rate for tantalum sludge leach and rinse is 205300 l/kkg (49198 gal/ton) of equivalent pure tantalum powder produced, based on the one water use rate reported. This rate is allocated only for those plants which use tantalum-bearing sludge as their raw material. The upgrading of tantalum-bearing sludge involves filtration for solids and spent acid separation and rinsing of the residual solids with water prior to the next leaching step. Water use and wastewater discharge rates are presented in Table V-3 (page 4564).

TANTALUM POWDER ACID WASH AND RINSE

The BPT wastewater discharge rate for tantalum powder acid wash and rinse is 350 l/kkg (84 gal/ton) of tantalum powder produced by the plant, based on the only reported water use rate. This rate is allocated only for those plants that incorporate a final acid wash of the tantalum powder to remove surface oxides,

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followed by a water rinse which cleans the powder prior to drying. The one plant that reported using such a system uses a tantalum recovery operation consisting of pH adjustment by ammonia addition and recovery of the precipitated tantalum solids. After treatment for tantalum recovery, the wastewater is further treated and discharged. The production normalized water use and discharge rates are presented in Table V-4 (page 4565).

LEACHING WET AIR POLLUTION CONTROL

The BPT wastewater discharge rate for acid leach wet air pollution control is 4880 l/kkg (ll69 gal/ ton) of equivalent pure tantalum powder produced. This rate is allocated for those plants which use a wet air pollution control system to control acid fumes which arise from the leaching operations. The available data indicate that this scrubber operates at 92 percent recycle. The BPT flow is based on this demonstrated recycle performance of the acid fume scrubber. Production normalized water use and discharge rates for this subdivision are presented in Table V-5 (page 4565).

REGULATED POLLUTANT PARAMETERS

The raw wastewater concentrations from individual operations and the subcategory as a whole were examined to select certain pollutant parameters for limitation. This examination and evaluation was presented in Section VI and also in Section X. A total of seven pollutants or pollutant parameters are selected for limitation under BPT and are listed below:

120.	copper
122.	lead
124.	nickel
128.	zinc
	tantalum
	TSS
	рH

EFFLUENT LIMITATIONS

The pollutant concentrations achievable by application of the promulgated BPT (both one-day maximum and monthly average values) are multiplied by the BPT normalized discharge flows summarized in Table IX-1 (page 4608) to calculate the mass of pollutants allowed to be discharged per mass of product. The results of these calculations in milligrams of pollutant per kilogram of product represent the BPT effluent limitations and are presented in Table IX-2 (page 4609) for each individual waste stream.

Table IX-1

BPT WASTEWATER DISCHARGE RATES FOR THE SECONDARY TANTALUM SUBCATEGORY

	BPT Normalized Discharge Rate		Production Normalizing
Wastewater Stream	l/kkg	gal/ton	Parameter
Tantalum Alloy Leach and Rinse	230,600	55,261	Tantalum powder produced
Capacitor Leach and Rinse	20,200	4,841	Tantalum powder produced from leaching
Tantalum Sludge Leach and Rinse	205,300	49,198	Equivalent pure tantalum powder produced
Tantalum Powder Acid Wash and Rinse	350	84	Tantalum powder produced
Leaching Wet Air Pollution Control	4,880	1,169	Equivalent pure tantalum powder produced

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TABLE IX-2

BPT MASS LIMITATIONS FOR THE SECONDARY TANTALUM SUBCATEGORY

(a) <u>Tantalum Alloy Leach</u> and <u>Rinse</u> BPT

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average
mg/kg (lb/mill)	ion lbs) of tant	alum powder produced
5, 5, 7, 7		
Antimony	661.800	295.200
*Copper	438.100	230.600
*Lead	96.850	46.120
*Nickel	442.800	292.900
Silver	94.550	39.200
*Zinc	336.700	140.700
*Tantalum	103.800	
*TSS	9,455.000	4,497.000
*pH Within the rang	ge of 7.5 to 10.	
(b) Capacitor Leach ar	nd Rinse BPT	
Pollutant or	Maximum for	Maximum for
Pollutant or	Maximum for any one day	
Pollutant or		Maximum for monthly average
Pollutant or pollutant property	any one day	monthly average
Pollutant or pollutant property	any one day	

*Copper	38.380	20.200
*Lead	8.484	4.040
*Nickel	38.780	25.650
Silver	8.282	3.434
*Zinc	29.490	12.320
*Tantalum	9.090	
*TSS	828.200	393.900
*pH Within the	e range of 7.5 to 10.0 at al	l times

*Regulated Pollutant

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TABLE IX-2 (Continued)

BPT MASS LIMITATIONS FOR THE SECONDARY TANTALUM SUBCATEGORY

(c) Tantalum Sludge Leach and Rinse BPT

Pollutant or	Maximum	for	Maximum	for
pollutant prope	rty any one	day	monthly	average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

Antimony	589.200	262.800
*Copper	390.100	205.300
*Lead	86.230	41.060
*Nickel	394.200	260.700
Silver	84.170	34.900
*Zinc	299.700	125.200
*Tantalum	92.390	— — —
*TSS	8,417.000	4,003.000
*pH Within the ran	nge of 7.5 to 10.0 at	all times

(d) Tantalum Powder Acid Wash and Rinse BPT

Pollutant or pollutant property		kimum for hthly average
mg/kg (lb/m	illion lbs) of tantalum	powder produced
Antimony *Copper *Lead *Nickel Silver *Zinc *Tantalum *TSS *pH Within the	1.005 0.665 0.147 0.672 0.144 0.511 0.158 14.350 range of 7.5 to 10.0 at	0.448 0.350 0.070 0.445 0.060 0.214 6.825 all times

*Regulated Pollutant

TABLE IX-2 (Continued)

BPT MASS LIMITATIONS FOR THE SECONDARY TANTALUM SUBCATEGORY

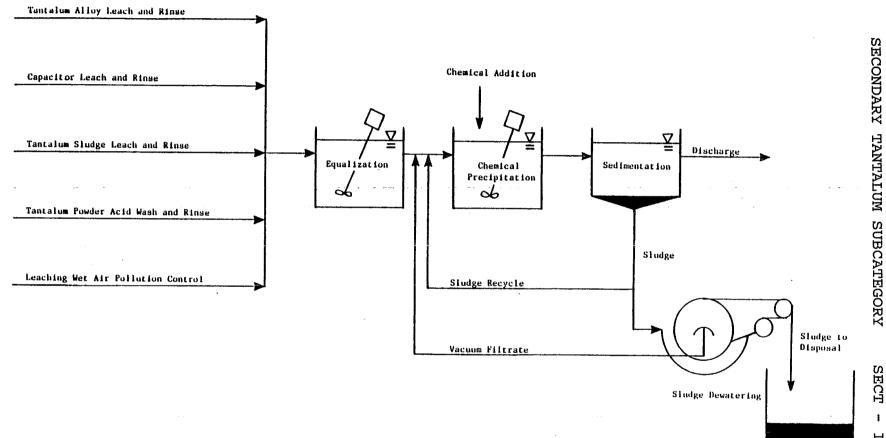
(e) Leaching Wet Air Pollution Control BPT

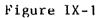
Pollutant		Maximum	for	Maximum	for
pollutant	property	any one	day	monthly	

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

Antimony *Copper *Lead *Nickel Silver *Zinc	14.010 9.272 2 050 9.370 2.001	6.246 4.880 0.976 6.198 0.830
*Tantalum *TSS *pH Within the	7.125 2.196 200.100 range of 7.5 to 10.0 at all	2.977 95.160 times

*Regulated Pollutant





BPT TREATMENT SCHEME FOR THE SECONDARY TANTALUM SUBCATEGORY

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SECTION X

BEST AVAILABLE TECHNOLOGY ECONOMICALLY ACHIEVABLE

These effluent limitations are based on the best control and treatment technology used by a specific point source within the industrial category or subcategory, or by another category from which it is transferable. Emphasis is placed on additional treatment techniques applied at the end of the treatment systems currently used, as well as reduction of the amount of water used and discharged, process control, and treatment technology optimization.

The factors considered in assessing best available technology economically achievable (BAT) include the age of equipment and facilities involved, the process used, process changes, nonwater quality environmental impacts (including energy requirements), and the costs of application of such technology BAT represents the best available technology economically achievable at plants of various ages, sizes, processes, or other characteristics. BAT may include feasible process changes or internal controls, even when not in common industry practice.

The required assessment of BAT considers costs, but does not require a balancing of costs against pollutant removals However, in assessing the proposed and promulgated BAT, the Agency has given substantial weight to the economic achievability of the technology.

TECHNICAL APPROACH TO BAT

The Agency reviewed a wide range of technology options and evaluated the available possibilities to ensure that the most effective and beneficial technologies were used as the basis of BAT. To accomplish this, the Agency elected to examine two technology options which could be applied to the secondary tantalum subcategory as alternatives for the basis of BAT effluent limitations.

For the development of BAT effluent limitations, mass loadings were calculated for each wastewater source or subdivision in the subcategory using the same technical approach as described in Section IX for BPT limitations development. The differences in the mass loadings for BPT and BAT are due to increased treatment.

POLLUTANT REMOVAL ESTIMATES

A complete description of the methodology used to calculate the estimated pollutant removal achieved by the application of the various treatment options is presented in Section X of Vol. I. In short, sampling data collected during the field sampling program were used to characterize the major wastewater streams considered for regulation. At each sampled facility, the sampling data were

SECONDARY TANTALUM SUBCATEGORY SECT - X

production normalized for each unit operation (i.e. mass of pollutant generated per mass of product manufactured). This value, referred to as the raw waste, was used to estimate the mass of toxic pollutants generated within the secondary tantalum subcategory. The pollutant removal estimates were calculated for each plant by first estimating the total mass of each pollutant in the untreated wastewater. This was calculated by first multiplying the raw waste values by the corresponding production value for that stream and then summing these values for each pollutant for every stream generated by the plant.

Next, the volume of wastewater discharged after the application of each treatment option was estimated for each operation at each plant by comparing the actual discharge to the regulatory flow. The smaller of the two values was selected and summed with the The mass of pollutant discharged was then other plant flows. estimated by multiplying the achievable concentration values attainable with the option (mq/1) by the estimated volume of process wastewater discharged by the subcategory. The mass of pollutant removed is the difference between the estimated mass of pollutant generated by each plant in the subcategory and the mass of pollutant discharged after application of the treatment option. The pollutant removal estimates for direct dischargers in the secondary tantalum subcategory are presented in Table X-1. (page 4618). These estimates are the same as those developed for the proposed regulation.

COMPLIANCE COSTS

In estimating subcategory-wide compliance costs, the first step was to develop a cost estimation model, relating the total costs with installation and operation associated of wastewater treatment technologies to plant process wastewater discharge. EPA applied the model to each plant. The plant's investment and operating costs are determined by what treatment it has in place and by its individual process wastewater discharge flow. As discussed above, this flow is either the actual or the BAT regulatory flow, whichever is lesser. The final step was to annualize the capital costs, and to sum the annualized capital costs, and the operating and maintenance costs for each plant, yielding the cost of compliance for the subcategory. The costs associated with the various options compliance are presented in Table X-2 (page 4619) for direct dischargers in the secondary tantalum subcategory. These costs were used in assessing economic achievability.

BAT OPTION SELECTION - PROPOSAL

EPA proposed BAT for the secondary tantalum subcategory based on Option C, chemical precipitation, sedimentation, and multimedia filtration technology.

The estimated capital cost of proposed BAT was \$13,474 and the annual cost was \$63,466 (1982 dollars). Implementation of the proposed BAT technology was estimated to remove 4.9 kilograms of

priority pollutants and 35.5 kilograms of suspended solids over the estimated BPT removal.

BAT OPTION SELECTION - PROMULGATION

EPA is promulgating BAT limitations for this subcategory based on Option C, which includes chemical precipitation, sedimentation, and multimedia filtration. The estimated capital cost of promulgated BAT is \$13,474 (1982 dollars) and the annual cost is \$63,466 (1982 dollars). The end-of-pipe treatment configuration for Option C is presented in Figure X-2 (page 4625).

EPA is promulgating BAT with multimedia filtration as part of the model technology because this technology results in additional removal of toxic metals. Filtration is also presently demonstrated at 25 plants throughout the nonferrous metals manufacturing category. Filtration adds reliability to the treatment system by making it less susceptible to operator error and to sudden changes in raw wastewater flow and concentrations.

Implementation of the control and treatment technologies of Option C would remove annually an estimated 26,273 kilograms of toxic metal pollutants and 20,115 kilograms of suspended solids, which is 4.9 kilograms of toxic metal pollutants and 35.5 kilograms of suspended solids over the estimated BPT removal.

WASTEWATER DISCHARGE RATES

A BAT discharge rate was calculated for each subdivision based upon the flows of the existing plants, as determined from analysis of the data collection portfolios. The discharge rate is used with the achievable treatment concentrations to determine BAT effluent limitations. Since the discharge rate may be different for each wastewater source, separate production normalized discharge rates for each of the five wastewater sources were determined and are summarized in Table X-3 (page 4620). The discharge rates are normalized on a production basis by relating the amount of wastewater generated to the mass of the intermediate product which is produced by the process associated with the waste stream in question. These production normalizing parameters, or PNPs, are also listed in Table X-3.

The BAT discharge rates reflect no flow reduction requirements as compared to the BPT option flows. In-process flow reduction beyond the BPT allowances is not achievable for any waste streams in this subcategory. As an example, the acid leach scrubber used at one of the secondary tantalum plants already operates at 92 percent recycle. Consequently, the BAT and BPT production normalized discharge flows are identical.

REGULATED POLLUTANT PARAMETERS

The raw wastewater concentrations from individual operations and the subcategory as a whole were examined to select certain pollutants and pollutant parameters for limitation. This examination and evaluation was presented in Section VI. The Agency, however, has chosen not to regulate all six priority pollutants selected in this analysis. The high cost associated with analysis for toxic metal pollutants has prompted EPA to develop an alternative method for regulating and monitoring metals priority pollutant discharges from the nonferrous manufacturing category. Rather than developing specific effluent mass limitations and standards for each of the toxic metals found in treatable concentrations in the raw wastewater from a given subcategory, the Agency is promulgating effluent mass limitations only for those pollutants generated in the greatest quantities as shown by the pollutant removal estimate analysis. The pollutants selected for specific limitation are listed below:

120. copper
122. lead
124. nickel
128. zinc
tantalum

By establishing limitations and standards for certain metal pollutants, dischargers will attain the same degree of control over toxic metal pollutants as they would have been required to achieve had all the toxic metal pollutants been directly limited.

This approach is technically justified since the achievable concentrations used for chemical precipitation and sedimentation technology are based on optimized treatment for concomitant multiple metals removal. Thus, even though metals have somewhat different theoretical solubilities, they will be removed at very precipitation and chemical same rate in a the nearly sedimentation treatment system operated for multiple metals removal. Filtration as part of the technology basis is likewise nonthis technology removes metals because justified preferentially.

The pollutants selected for specific limitation in the secondary tantalum subcategory are copper, lead, nickel, zinc, and tantalum. The following toxic metal pollutants are excluded from limitation on the basis that they are effectively controlled by the limitations developed for copper, lead, nickel, zinc, and tantalum:

114. antimony 126. silver

The priority metal pollutants copper, lead, nickel, and zinc, as well as the nonconventional metal pollutant tantalum, are specifically limited to ensure the control of the excluded priority metal pollutants These pollutants are indicators of the performance of the treatment technology.

EFFLUENT LIMITATIONS

The achievable concentrations, both one day maximum and monthly

SECONDARY TANTALUM SUBCATEGORY SECT - X

average values, are multiplied by the BAT normalized discharge flows summarized in Table X-3 (page 4620) to calculate the mass of pollutants allowed to be discharged per mass of product. The results of these calculations in milligrams of pollutant per kilogram of product represent the BAT effluent limitations and are presented in Table X-4 (page 4621) for each wastewater stream.

Table X-1

POLLUTANT REMOVAL ESTIMATES FOR DIRECT DISCHARGERS

Pollutant	RAW WASTE (kg/yr)	OPTION A DISCHARGE (kg/yr)	OPTION A Removed (kg/yr)	OPTION C DISCHARGE (kg/yr)	OPTION C REMOVED (kg/yr)
			3.0773	1.7765	8,7364
Antimony	10.5129	2.6462	7.8667	0.9541	0.6052
Arsenic	1.5593	1.4213	0.1380	0.1879	5.6579
Cadmium	5.8458	0.2985	5.5473	0.1879	8.8253
Chromium (total)	9.0866	0.3172	8.7694	1.4734	14523.2316
Copper	14524,7051	2.1924	14522.5126		0.1102
Cyanide (total)	0.2502	0.1664	0.0838	0.1400	4594.3368
Lead	- 4594.6398	0.4538	4594.1860	0.3030	0.0036
Mercury	0.0083	0.0048	0.0036	0.0048	
Nickel	4418.2463	2.7975	4415.4488	0.8358	4417.4105 22.8442
Selenium	23,1056	0.3782	22.7275	0.2614	0.1809
Silver	0.5509	0.3657	0.1853	0.3700	0.2633
Thallium	0.7842	0.5179	0.2663	0.5209	2690.7649
Zinc	2691.6323	1.2475	2690.3848	0.8674	2090.7049
Total Priority Pollutants	26280.9274	12.8072	26268.1202	7.9564	26272.9709
101200000				0.0750	0.1590
Ammonia	0.5249	0.3659	0.1590	0.3659	1491.5479
Cobalt	1491.6734	0.1885	1491.4849	0.1255	2.5610
Flouride	2.9343	0.3810	2.5534	0.3733	2.3010
Total Nonconv.	1495.1326	0.9354	1494.1972	0.8647	1494.2679
	20124.3335	45.3640	20078.9695	9.8329	20114.5006
TSS		93.7944	47.5923	93.7911	47.5955
Oil and Grease	141.3866	33+7344			
Total Conv.	20265.7201	139.1584	20126.5617	103.6240	20162.0961
Total Pollutants	48041.7801	152.9010	47888.8791	112.4452	47929.3350

SECONDARY TANTALUM SUBCATEGORY

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TABLE X-2

COST OF COMPLIANCE FOR THE SECONDARY TANTALUM SUBCATEGORY

Direct Dischargers

Option	Total Required Capital Cost (1982 dollars)	Total Annual Cost (1982 dollars)
A	6462	58854
С	13474	63466

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Table X-3

BAT WASTEWATER DISCHARGE RATES FOR THE SECONDARY TANTALUM SUBCATEGORY

	BAT Norr Discharg	ge Rate	Production Normalizing
Wastewater Stream	1/kkg	gal/ton	Parameter
Tantalum Alloy Leach and Rinse	230,600	55,261	Tantalum powder produced
Capacitor Leach and Rinse	20,200	4,841	Tantalum powder produced from leaching
Tantalum Sludge Leach and Rinse	205,300	49,198	Equivalent pure tantalum powder produced
Tantalum Powder Acid Wash and Rinse	350	84	Tantalum powder produced
Leaching Wet Air Pollution Control	4,880	1,169	Equivalent pure tantalum powder produced

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SECONDARY TANTALUM SUBCATEGORY SECT - X

TABLE X-4

BAT MASS LIMITATIONS FOR THE SECONDARY TANTALUM SUBCATEGORY

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(a) Tantalum Alloy Leach and Rinse BAT

Pollutant pollutant		Maximum for any one day	Maximum for monthly ave	rage	
mg/	′kg (lb∕milli	lon lbs) of tanta	lum powder p	roduced	
Antimony		445.100	198	3.300	
*Copper		295.200		0.700	
*Lead		64.570		9.980	
*Nickel	. ,	126.800		5.320	
Silver		66.870		7.670	
*Zinc		235.200		5.850	-
*Tantalum		103.800			
	tor Leach an	d Rinse BAT			
Pollutant	or	d <u>Rinse</u> BAT Maximum for any one day	Maximum for monthly aver	age	•
Pollutant pollutant	or property	Maximum for	monthly aver der produced 17 12 2 7 2	· · ·	hinç

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TABLE X-4 (Continued)

BAT MASS LIMITATIONS FOR THE SECONDARY TANTALUM SUBCATEGORY

(c) Tantalum Sludge Leach and Rinse BAT

Pollutant o	זר	Maximum	for	Maximum	for
pollutant p		any one		monthly	average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

Antimony *Copper *Lead *Nickel Silver *Zinc *Tantalum	396.200 262.800 57.480 112.900 59.540 209.400 92.390	176.600 125.200 26.690 75.960 24.640 86.230
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(d) Tantalum Powder Acid Wash and Rinse BAT

Pollutant or pollutant pro		laximum for ny one day	Maximum for monthly average
mg/kg	(lb/million	lbs) of tanta	lum powder produced
Antimony		0.676	0.301
*Copper		0.448	0.214
*Lead		0.098	0.046
*Nickel		0.193	0.130
Silver		0.102	0.042
		0.357	0.147
*Zinc *Tantalum		0.158	

*Regulated Pollutant

TABLE X-4 (Continued)

BAT MASS LIMITATIONS FOR THE SECONDARY TANTALUM SUBCATEGORY

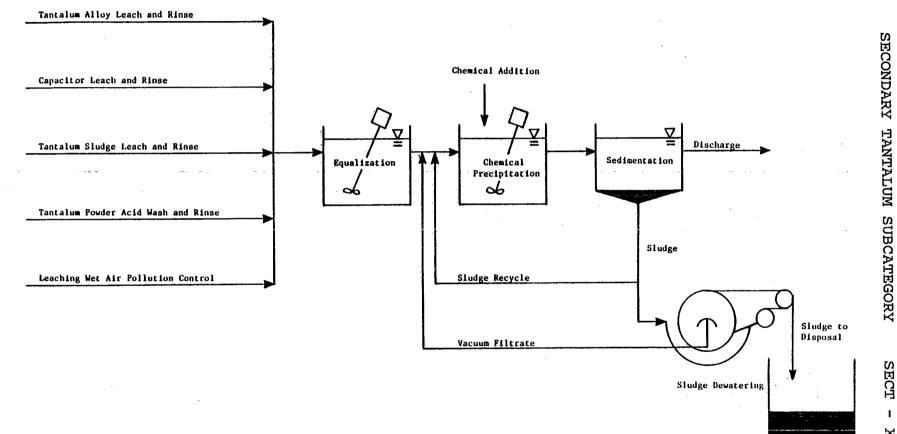
(e) Leaching Wet Air Pollution Control BAT

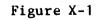
Pollutant or		Maxin	num	for	Maximum	for
pollutant pro	operty	any c	one	day	monthly	average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

Antimony	9.418	4.197
*Copper	6.246	2.977
*Lead	1.366	0.634
*Nickel	2.684	1.806
Silver	1.415	0.586
*Zinc	4.978	2.050
*Tantalum	2.196	

*Regulated Pollutant





BAT TREATMENT SCHEME FOR OPTION A

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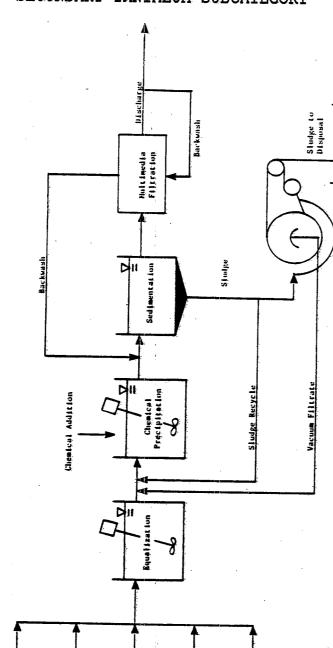


Figure X-2 BAT TREATMENT SCHEME FOR OFTION C

Shudge Dewatering

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Tantalum Powder Acid Wash and Rinse

Tantalym Sludge Leach and Rinse

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leaching Net Air Pollutiun Control

Capacitor Leach and Minue

Tantalue Alluy leach and Ainse

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SECONDARY TANTALUM SUBCATEGORY SECT - XI

SECTION XI

NEW SOURCE PERFORMANCE STANDARDS

The basis for new source performance standards (NSPS) is the best available demonstrated technology (BDT). New plants have the opportunity to design the best and most efficient production processes and wastewater treatment technologies without facing the added costs and restrictions encountered in retrofitting an existing plant. Therefore, EPA has considered the best demonstrated process changes, in-plant controls, and end-of-pipe treatment technologies which reduce pollution to the maximum extent feasible.

This section describes the technologies for treatment of wastewater from new sources and presents mass discharge standards for regulated pollutants for NSPS in the secondary tantalum subcategory, based on the selected treatment technology.

TECHNICAL APPROACH TO NSPS

New source performance standards are equivalent to the best available technology (BAT) selected for currently existing secondary tantalum plants. This result is a consequence of careful review by the Agency of a wide range of technical options for new source treatment systems which is discussed in Section IX of the General Development Document. Additionally, there was nothing found to indicate that the wastewater flows and characteristics of new plants would not be similar to those from existing plants, since the processes used by new sources are not expected to differ from those used at existing sources. Consequently, BAT production normalized discharge rates, which are based on the best existing practices of the subcategory, can also be applied to new sources. These rates are presented in Table XI-1 (page 4629).

Treatment technologies considered for the NSPS options are identical to the treatment technologies considered for the BAT options. These options are:

OPTION A

o Chemical precipitation and sedimentation

OPTION C

O Chemical precipitation and sedimentation
 Multimedia filtration

NSPS OPTION SELECTION - PROPOSAL

EPA proposed that the best available demonstrated technology for the secondary tantalum subcategory be equivalent to Option C (chemical precipitation, sedimentation, and multimedia filtration).

The wastewater flow rates for proposed NSPS are the same as the proposed BAT flow rates. Flow reduction measures for NSPS were not considered feasible because no new demonstrated technologies existed within the subcategory that improved on water use and Therefore, EPA concluded practices. flow discharge that reduction beyond the allowances proposed for BAT was unachievable, and NSPS flow rates should be equal to those for BAT.

NSPS OPTION SELECTION - PROMULGATION

EPA is promulgating NSPS for the secondary tantalum subcategory based on Option C (chemical precipitation, sedimentation, and This technology is demonstrated by 25 multimedia filtration). plants in the nonferrous metals manufacturing category. The wastewater flow rates for promulgated NSPS are the same as the promulgated BAT flow rates. Flow reduction measures for NSPS are not feasible, because dry scrubbing is not demonstrated for controlling emissions from acid leaching operations. The nature these emissions (acid fumes, hot particulate of matter) technically precludes the use of dry scrubbers. Therefore, EPA is including an allowance from this source at NSPS equivalent to that promulgated for BAT. EPA also does not believe that new plants could achieve any additional flow reduction beyond the 92 percent scrubber effluent recycle presently practiced in the industry.

REGULATED POLLUTANT PARAMETERS

The Agency has no reason to believe that the pollutants that will be found in treatable concentrations in processes within new sources will be any different than with existing sources. Accordingly, pollutants and pollutant parameters selected for limitation under NSPS, in accordance with the rationale of Sections VI and X, are identical to those selected for BAT. The conventional pollutant parameters TSS and pH are also selected for limitation.

NEW SOURCE PERFORMANCE STANDARDS

The NSPS discharge flows for each wastewater source are the same as the discharge rates for BAT and are shown in Table XI-1. The mass of pollutant allowed to be discharged per mass of product is calculated multiplying the appropriate by achievable (mg/l) by the production normalized wastewater concentration discharge flows (l/kkg). The results of these calculations are the production-based new source performance standards. These standards are presented in Table XI-2.

Table XI-1

NSPS WASTEWATER DISCHARGE RATES FOR THE SECONDARY TANTALUM SUBCATEGORY

Wastewater Stream	NSPS Normalized Discharge Rate		Production Normalizing
wastewater Stream	1/kkg	gal/ton	Parameter
Tantalum Alloy Leach and Rinse	230,600	55,261	Tantalum powder produced
Capacitor Leach and Rinse	20,200	4,841	Tantalum powder produced from leaching
Tantalum Sludge Leach and Rinse	205,300	49,198	Equivalent pure tantalum powder produced
Tantalum Powder Acid Wash and Rinse	350	84	Tantalum powder produced
Leaching Wet Air Pollution Control	4,880	1,169	Equivalent pure tantalum powder produced

SECONDARY TANTALUM SUBCATEGORY SECT - XI

TABLE XI-2

NSPS FOR THE SECONDARY TANTALUM SUBCATEGORY

(a) Tantalum Alloy Leach and Rinse NSPS

Pollutant	or	Maximum	for	Maximum	for
pollutant		any one	day	monthly	average

mg/kg	(lb/million 1	lbs) of tantalur	n powder produced
Antimony		445.100	198.300
*Copper		295.200	140.700
*Lead		64.570	29.980
*Nickel		126.800	85.320
Silver		66.870	27.670
*Zinc		235.200	96.850
*Tantalum		103.800	
*TSS		3,459.000	2,767.000
*pH Within		E 7.5 to 10.0 at	t all times

(b) Capacitor Leach and Rinse NSPS

Pollutant	or	Maximum	for	Maximum	for
pollutant		any one	day	monthly	average

mg/kg (lb/million lbs) of tantalum powder produced from leaching

Antimony	38.990	17.370 12.320
*Copper	25.860 5.656	2.626
*Lead *Nickel	11.110	7.474
Silver	5.858	2.424
*Zinc	20.600	8.484
*Tantalum	9.090	
*TSS	303.000	242.400
*pH Within the ran	ge of 7.5 to 10.0 at	all times

*Regulated Pollutant

TABLE XI-2 (Continued)

NSPS FOR THE SECONDARY TANTALUM SUBCATEGORY

(c) Tantalum Sludge Leach and Rinse NSPS

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Pollutant or	Maximum for	Maximum for	
pollutant property	any one day	monthly average	<i>,</i> .

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

Antim	ony	396.200	176.600
*Coppe	r	262.800	125.200
	*Lead 57.480		26.690
*Nickel		112.900	75.960
Silve	r	59.540	24.640
	*Zinc 209.400		86.230
	*Tantalum 92.390		
*TSS		. 3,080.000	2,464.000
*pH	Within	the range of 7.5 to 10.0 at all	times

(d) Tantalum Powder Acid Wash and Rinse NSPS

Pollutant		Maximum	for	Maximum	for
pollutant	property	any one	day	monthly	average

mg/kg (lb/million lbs) of tantalum powder produced Antimony 0.676 0.301 *Copper 0.448 0.214 *Lead 0.098 0.046 *Nickel 0.193 0.130 Silver 0.102 0.042 *Zinc 0.357 0.147 *Tantalum 0.158 _ _ _ *TSS 5.250 4.200 Within the range of 7.5 to 10.0 at all times *pH

*Regulated Pollutant

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SECONDARY TANTALUM SUBCATEGORY SECT - XI

Table XI-2 (Continued)

NSPS FOR THE SECONDARY TANTALUM SUBCATEGORY

(e) Leaching Wet Air Pollution Control NSPS

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Pollutant or	Maximum for	Maximum for	
pollutant property	any one day	monthly average	
mg/kg (lb/million lbs)	of equivalent	pure tantalum powder	produced
Antimony	9.418	4.197	
*Copper	6.246	2.977	
*Lead	1.366	0.634	
*Nickel	2.684	1.806	
Silver	1.415	0.586	
*Zinc	4.978	2.050	
*Tantalum	2.196		
*TSS	73.200	58.560	
*pH Within the range	of 7.5 to 10.0	D at all times	

*Regulated Pollutant

SECONDARY TANTALUM SUBCATEGORY SECT - XII

SECTION XII

PRETREATMENT STANDARDS

This section describes the control and treatment technologies for pretreatment of process wastewaters from new sources in the secondary tantalum subcategory. PSES are designed to prevent the discharge of pollutants which pass through, interfere with, or are otherwise incompatible with the operation of publicly owned treatment works (POTW). The Clean Water Act requires pretreatment for pollutants, such as toxic metals, that limit POTW sludge management alternatives. New indirect discharge facilities, like new direct discharge facilities, have the opportunity to incorporate the best available demonstrated technologies, including process changes, in-plant controls, and end-of-pipe treatment technologies, and to use plant site selection to ensure adequate treatment system installation. Pretreatment standards are to be technology based, analogous to the best available technology for removal of toxic pollutants. Pretreatment standards for regulated pollutants are presented based on the selected control and treatment technology.

Pretreatment standards for existing sources (PSES) will not be promulgated for the secondary tantalum subcategory because there are no existing indirect dischargers in this subcategory. However, pretreatment standards for new sources (PSNS) will be promulgated.

TECHNICAL APPROACH TO PRETREATMENT

Before proposing and promulgating pretreatment standards, the Agency examines whether the pollutants discharged by the industry pass through the POTW or interfere with the POTW operation or its sludge disposal practices. In determining chosen whether pollutants pass through a well-operated POTW achieving secondary treatment, the Agency compares the percentage of a pollutant removed by POTW with the percentage removed by direct dischargers applying the best available technology economically achievable. A pollutant is deemed to pass through the POTW when the average percentage removed nationwide by well-operated POTW meeting secondary treatment requirements, is less than the percentage removed by direct dischargers complying with BAT effluent standards guidelines for that pollutant.

This definition of pass through satisfies the two competing objectives set by Congress that standards for indirect dischargers be equivalent to standards for direct dischargers while at the same time the treatment capability and performance of the POTW be recognized and taken into account in regulating the discharge of pollutants from indirect dischargers. The Agency compares percentage removal rather than the mass or concentration of pollutants discharged because the latter would not take into account the mass of pollutants discharged to the POTW from nonindustrial sources or the dilution of the pollutants in the POTW effluent to lower concentrations due to the addition of large amounts of non-industrial wastewater.

PRETREATMENT STANDARDS FOR NEW SOURCES

Options for pretreatment of wastewaters from new sources are based on increasing the effectiveness of end-of-pipe treatment technologies. All in-plant changes and applicable end-of-pipe treatment processes have been discussed previously in Sections X and XI. The options for PSNS, therefore, are the same as the BAT options discussed in Section X.

A description of each option is presented in Section X. Treatment technologies considered for the PSNS options are:

OPTION A

o Chemical precipitation and sedimentation

OPTION C

- o Chemical precipitation and sedimentation
- o Multimedia filtration

PSNS OPTION SELECTION - PROPOSAL

EPA proposed PSNS for the secondary tantalum subcategory based on Option C, chemical precipitation, sedimentation, and multimedia filtration. The wastewater discharge rates proposed for PSNS are equivalent to the proposed BAT discharge rates. No flow reduction measures for PSNS were considered feasible beyond the rates proposed for BAT.

PSNS OPTION SELECTION - PROMULGATION

EPA has selected Option C (chemical precipitation, sedimentation, and multimedia filtration) as the regulatory approach for pretreatment standards for new sources on the basis that it achieves effective removal of toxic pollutants and is demonstrated by 25 plants throughout the nonferrous metals manufacturing category.

The wastewater discharge rates for PSNS are identical to the promulgated BAT discharge rates for each waste stream. The PSNS discharge rates are shown in Table XII-1 (page 4636). No additional flow reduction measures for PSNS are feasible. EPA does not believe that new plants could achieve flow reduction beyond the 92 percent scrubber effluent recycle presently practiced in the industry.

REGULATED POLLUTANT PARAMETERS

Pollutants selected for limitation, in accordance with the rationale of Sections VI and X, are identical to those selected

for limitation for BAT. It is necessary to promulgate PSNS to prevent the pass-through of copper, lead, nickel, and zinc, which are the limited pollutants. These toxic pollutants are removed by a well-operated POTW achieving secondary treatment at an average of 48 percent while BAT level technology removes approximately 99 percent.

PRETREATMENT STANDARDS

Pretreatment standards are based on the pollutant concentrations achievable from the selected treatment technology, (Option C), and the discharge rates determined in Section X for BAT. A mass of pollutant per mass of product (mg/kg) allocation is given for each subdivision within the subcategory. This pollutant allocation is based on the product of the treatable concentration from the promulgated treatment (mg/l) and the production normalized wastewater discharge rate (l/kkg). The achievable treatment concentrations for BAT are identical to those for PSNS. PSNS are presented in Table XII-2 (page 4637).

Table XII-1

PSNS WASTEWATER DISCHARGE RATES FOR THE SECONDARY TANTALUM SUBCATEGORY

	PSNS Normalized Discharge Rate		Production Normalizing		
Wastewater Stream	1/kkg	gal/ton	Parameter		
Tantalum Alloy Leach and Rinse	230,600	55,261	Tantalum powder produced		
Capacitor Leach and Rinse	_20_,200	4,841	Tantalum powder produced from leaching		
Tantalum Sludge Leach and Rinse	205,300	49,198	Equivalent pure tantalum powder produced		
Tantalum Powder Acid Wash and Rinse	350	84	Tantalum powder produced		
Leaching Wet Air Pollution Control	4,880	1,169	Equivalent pure tantalum powder produced		

SECONDARY TANTALUM SUBCATEGORY SECT - XII

TABLE XII-2

PSNS FOR THE SECONDARY TANTALUM SUBCATEGORY

(a) Tantalum Alloy Leach and Rinse PSNS

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average
· · · · · · · · · · · · · · · · · · ·		
mg/kg (1b/mil.	lion 1bs) of tant	alum powder produced
Antimony	445.100	198.300
*Copper	295.200	140.700
*Lead	64.570	29.980
*Nickel	126.800	85.320
Silver	66.870	27.670
*Zinc	235.200	96.850
*Tantalum	103.800	
Pollutant or	and <u>Rinse</u> PSNS Maximum for	Maximum for
Pollutant or		Maximum for monthly average
(b) <u>Capacitor Leach</u> Pollutant or pollutant property mg/kg (lb/million lbs	Maximum for any one day	monthly average
Pollutant or pollutant property mg/kg (lb/million lbs	Maximum for any one day s) of tantalum po	monthly average wder produced from lea
Pollutant or pollutant property mg/kg (lb/million lbs Antimony	Maximum for any one day s) of tantalum po 38.990	monthly average wder produced from lea 17.370
Pollutant or pollutant property mg/kg (lb/million lbs Antimony *Copper	Maximum for any one day s) of tantalum po 38.990 25.860	monthly average wder produced from lea 17.370 12.320
Pollutant or pollutant property mg/kg (lb/million lbs Antimony *Copper *Lead	Maximum for any one day s) of tantalum po 38.990 25.860 5.656	monthly average wder produced from lea 17.370 12.320 2.626
Pollutant or pollutant property mg/kg (lb/million lbs Antimony *Copper *Lead *Nickel	Maximum for any one day s) of tantalum po 38.990 25.860 5.656 11.110	monthly average wder produced from lea 17.370 12.320 2.626 7.474
Pollutant or pollutant property mg/kg (lb/million lbs Antimony *Copper *Lead *Nickel Silver	Maximum for any one day s) of tantalum po 38.990 25.860 5.656 11.110 5.858	monthly average wder produced from lea 17.370 12.320 2.626 7.474 2.424
Pollutant or pollutant property mg/kg (lb/million lbs Antimony *Copper *Lead *Nickel	Maximum for any one day s) of tantalum po 38.990 25.860 5.656 11.110	monthly average wder produced from lea 17.370 12.320 2.626 7.474

*Regulated Pollutant

TABLE XII-2 (Continued)

PSNS FOR THE SECONDARY TANTALUM SUBCATEGORY

(c) Tantalum Sludge Leach and Rinse PSNS

Pollutant o	or	Maximum	for	Maximum	for
pollutant p		any one	day	monthly	average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

(d) Tantalum Powder Acid Wash and Rinse PSNS

-		
Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average

mg/kg (lb/million lbs) of tantalum powder produced 0.301 0.676 Antimony 0.214 0.448 *Copper 0.046 0.098 *Lead 0.130 0.193 *Nickel 0.042 0.102 Silver 0.357 0.147 *Zinc 0.158 -----*Tantalum

*Regulated Pollutant

SECONDARY TANTALUM SUBCATEGORY SECT - XII

TABLE XII-2 (Continued)

PSNS FOR THE SECONDARY TANTALUM SUBCATEGORY

(e) Leaching Wet Air Pollution Control PSNS

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average

mg/kg (lb/million lbs) of equivalent pure tantalum powder produced

Antimony	9.418	4.197
*Copper	6.246	2.977
*Lead	1.366	0.634
*Nickel	2.684	1.806
Silver	1.415	0.586
*Zinc	4.978	2.050
*Tantalum	2.196	

*Regulated Pollutant

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SECONDARY TANTALUM SUBCATEGORY SECT - XIII

SECTION XIII

BEST CONVENTIONAL POLLUTANT CONTROL TECHNOLOGY

EPA is not promulgating best conventional pollutant control technology (BCT) for the secondary tantalum subcategory at this time.

SECONDARY TANTALUM SUBCATEGORY SECT - XIII

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NONFERROUS METALS MANUFACTURING POINT SOURCE CATEGORY

DEVELOPMENT DOCUMENT SUPPLEMENT

for the

Secondary Uranium Subcategory

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May 1989

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SECTION I

SUMMARY

This document provides the technical basis for promulgating effluent limitations based on best practicable technology (BPT) and best available technology (BAT) for existing direct dischargers, standards of performance for new source direct dischargers (NSPS), and pretreatment standards for new indirect dischargers (PSNS).

The secondary uranium subcategory consists of three plants. Of the three plants, two discharge directly to surface waters, and one achieves zero discharge of process wastewater.

EPA first studied the secondary uranium subcategory to determine differences in raw materials, final whether products, manufacturing processes, equipment, age and size of plants, or water usage required the development of separate effluent limitations and standards for different segments of the This involved a detailed analysis of wastewater subcategory. discharge and treated effluent characteristics, including the sources and volume of water used, the processes used, the sources of pollutants and wastewaters in the plant, and the constituents of wastewaters including priority pollutants. As a result, nine subdivisions or building blocks have been identified for this subcategory that warrant separate effluent limitations. These include:

- (a) Refinery sump filtrate,
- (b) Slag leach reslurry,
- (c) Digestion wet air pollution control,
- (d) Solvent extraction raffinate filtrate,
- (e) Evaporation and denitration wet air pollution control,
- (f) Hydrofluorination water scrubber,
- (g) Hydrofluorination alkaline scrubber,
- (h) Magnesium reduction and casting floor wash water, and
- (i) Laundry wastewater.

EPA also identified several distinct control and treatment technologies (both in-plant and end-of-pipe) applicable to the secondary uranium subcategory. The Agency analyzed both historical and newly generated data on the performance of these technologies, including their nonwater quality environmental impacts and air quality, solid waste generation, and energy requirements. EPA also studied various flow reduction techniques reported in the data collection portfolios (dcp) and plant visits.

Engineering costs were prepared for each of the control and treatment options considered for the subcategory. These costs were then used by the Agency to estimate the impact of implementing the various options on the subcategory. For each

SECT - I

control and treatment option that the Agency found to be most effective and technically feasible in controlling the discharge of pollutants, we estimated the number of potential closures, number of employees affected, and impact on price. These results are reported in a separate document entitled "The Economic Impact Analysis of Effluent Limitations and Standards for the Nonferrous Metals Manufacturing Industry."

After examining the various treatment technologies, 'the Agency has identified BPT to represent the average of the best existing technology. Metals removal based on chemical precipitation and sedimentation technology is the basis for the BPT limitations. To meet the BPT effluent limitations based on this technology, the secondary uranium subcategory is expected to incur an estimated capital cost of \$54,800 and an annual cost of \$90,400.

For BAT, filtration is added as an effluent polishing step to the BPT end-of-pipe treatment scheme. To meet the BAT effluent limitations based on this technology, the secondary uranium subcategory is estimated to incur a capital cost of \$88,000 and an annual cost of \$106,700.

NSPS is equivalent to BAT. In selecting NSPS, EPA recognizes that new plants have the opportunity to implement the best and most efficient manufacturing processes and treatment technology. As such, the technology basis of BAT has been determined as the best demonstrated technology.

PSES is not being promulgated for this subcategory because there are no existing indirect dischargers in the secondary uranium subcategory. For PSNS, the Agency selected pretreatment and endof-pipe treatment techniques equivalent to BAT.

The best conventional technology (BCT) replaces BAT for the control of conventional pollutants. BCT is not being promulgated at this time because the methodology for BCT has not yet been finalized.

The mass limitations and standards for BPT, BAT, NSPS, and PSNS are presented in Section II.

SECTION II

CONCLUSIONS

EPA has divided the secondary uranium subcategory into nine subdivisions for the purpose of effluent limitations and standards. These subdivisions are:

- (a) Refinery sump filtrate,
- (b) Slag leach reslurry,
- (c) Digestion wet air pollution control,
- (d) Solvent extraction raffinate filtrate,
- (e) Evaporation and denitration wet air pollution control,
- (f) Hydrofluorination water scrubber,
- (g) Hydrofluorination alkaline scrubber,
- (h) Magnesium reduction and casting floor wash water, and
- (i) Laundry wastewater.

BPT is promulgated based on the performance achievable by the application of chemical precipitation and sedimentation technology. The following BPT effluent limitations are promulgated:

(a) <u>Refinery Sump Filtrate</u> BPT

Pollutan	tant or	Maximum for	Maximum for
	Property	Any One Day	Monthly Average
mg/kg	(lb/million	lbs) of uranium	processed in the refinery
Chromium		32.270	13.200
Copper		139.300	73.340
Nickel		140.800	93.140
Fluoride		2,567.000	1,459.000
TSS		3,007.000	1,430.000
pH		ne range of 7.5	to 10.0 at all times

b) Slag Leach Reslurry BPT

	tant or	Maximum for	Maximum for
	t Property	Any One Day	MonthlY Average
mg/kg	(lb/million	lbs) of uranium	processed in the refinery
Chromium		2.009	0.822
Copper		8.675	4.566
Nickel		8.767	5.799
Fluoride		1?9.800	90.860
TSS		187.200	89.0?0
pH		the range of 7.5	5 to 10.0 at all times

(c) Digestion Wet Air Pollution Control BPT

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million	lbs) of uranium	processed in the refinery
Chromium (total)	0.000	0.000
Copper	0.000	0.000
Nickel	0.000	0.000
Fluoride	0.000	0.000
TSS	0.000	0.000
pH Within t	the range of 7.5	to 10.0 at all times

(d) Solvent Extraction Raffinate Filtrate BPT

	ant or	Maximum for	Maximum for
	Property	Any One Day	Monthly Average
mg/kg	(lb/million	lbs) of uranium	processed in the refinery
Chromium		2.802	1.146
Copper		12.100	6.369
Nickel		12.230	8.089
Fluoride		222.900	126.700
TSS		261.100	124.200
pH		ne range of 7.5	to 10.0 at all times

(e) Evaporation and Denitration Wet Air Pollution Control BPT

Pollutant or Pollutant Property		Maximum for Maximum for Any One Day Monthly Average		
mg	/kg (lb/mil	lion lbs) of ura	nium trioxide produced	
Chromium Copper Nickel Fluoride TSS pH		0.000 0.000 0.000 0.000 0.000 e range of 7.5 to	0.000 0.000 0.000 0.000 0.000 0.000 0 10.0 at all times	

(f) Hydrofluorination	<u>Water</u> Scrubber	<u>c</u> BPT
Pollutant or	Maximum for	Maximum for
Pollutant Property		
mg/kg (lb/million	lbs) of uranium	n tetrafluoride produced
Chromium (total)	0.000	0.000
Copper	0.000	0.000
Nickel	0.000	0.000
Fluoride	0.000	0.000
TSS	0.000	0.000
pH Within the	range of 7.5 to	0 10.0 at all times
(g) <u>Hydrofluorination</u>	Alkaline Scrub	ober BPT
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million	lbs) of uranium	a tetrafluoride produced
Chromium (total)	0.009	0.004
Copper	0.038	0.020
Nickel	0.038	0.025
Fluoride	0.700	0.398
TSS	0.820	0.390
pH Within the	range of 7.5 to	0 10.0 at all times
(h) <u>Magnesium</u> Reducti	on and Casting	<u>Floor Wash</u> BPT
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million lbs)	of uranium pro	oduced by magnesium reduction
Chromium (total)	0.013	0.005
Copper	0.057	0.030
Nickel	0.058	0.038
Fluoride	1.054	0.599
TSS	1.234	0.587
pH Within the	range of 7.5 t	o 10.0 at all times
· · · · · · · · · · · · · · · · · · ·	· · · · · · · · · · · · · · · · · · ·	· · · · · · · · · · · · · · · · · · ·

(i) Laundry Wastewater BPT

Pollutant or	Maximum for	Maximum for	· · · ·
Pollutant Property	Any One Day	Monthly Average	
mg/kg (lb/million lbs)	of uranium produ	iced by magnesium	reduction
Chromium (total)	0.084	0.035	÷
Copper	0.365	0.192	
Nickel	0.369	0.244	
Fluoride	6.720	3.821	
TSS	7.872	3.744	
pH Within the P	cange of 7.5 to 1	L0.0 at all times	

BAT is promulgated based on the performance achievable by the application of chemical precipitation, sedimentation, and multimedia filtration technology. The following BAT effluent limitations are promulgated:

(a) Refinery Sump Filtrate BAT

Pollutant		Maximu Any On		Maxim Monthly			9
mg/kg	(lb/million	lbs) of	uranium	processed	in	the	refinery
Chromium Copper Nickel Fluoride	(total)	27.1 93.8 40.3 2,567.0	80 40	11.0 44. 27.1 1,459.0	740 140		

(b) Slag Leach Reslurry BAT

Pollutant or		Maximum for		Maximum for		
Pollutant	Property	Any	One Day	Monthly	Averag	e
mg/kg	(lb/million	lbs) o	of uranium	processed	in the	refinery
Chromium Copper Nickel Fluoride	(total)	5 2	.689 .844 .511 .800	2.	685 785 689 860	

Digestion Wet Air Pollution Control (C) BAT Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average mg/kg (lb/million lbs) of uranium processed in the refinery 0.000 Chromium (total) 0.000 Copper 0.000 0.000 Nickel 0.000 0.000 Fluoride 0.000 0 000 (đ) Solvent Extraction Raffinate Filtrate BAT

Pollutant or Maximum for Maximum for Monthly Average Pollutant Property Any One Day mg/kg (lb/million lbs) of uranium processed in the refinery Chromium (total) 2.357 0.955 Copper 8.152 3.885 Nickel 3.503 2.357 Fluoride 222.900 126.700

(e) Evaporation and Denitration Wet Air Pollution Control BAT

Pollutant or Pollutant Property	Maximum for Any One Day	Maximum for Monthly Average		
mg/kg (lb/mill	ion lbs) of urar	nium trioxide produce	ed	
Chromium (total) Copper	0.000	0.000 0.000		
Nickel Fluoride	0.000	0.000		

(f) Hydrofluorination Water Scrubber BAT

		· · ·		
Pollutant or			Maximum for	
Pollutant Property		e Day	Monthly Average	
b/million	lbs) of	uranium	tetrafluoride	produced
tal)	0.00	00	0.000	
·	0.00)0	0.000	
	0.00	00	0.000	
	0.00	00	0.000	
	operty b/million	operty Any One b/million lbs) of tal) 0.00 0.00 0.00	operty Any One Day b/million lbs) of uranium	operty Any One Day Monthly Avera b/million lbs) of uranium tetrafluoride tal) 0.000 0.000 0.000 0.000 0.000 0.000

(g) Hydrofluorination Alkaline Scrubber BAT

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million	lbs) of uraniu	m tetrafluoride produce
Chromium (total)	0.007	0.003
Copper	0.026	0.012
Nickel	0.011	0.007
Fluoride	0.700	0.398
(h) <u>Magnesium</u> <u>Reducti</u>	on and Casting	Floor Wash BAT
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million lbs) of uranium pro	oduced by magnesium red
Chromium (total)	0.011	0.005
Copper	0.039	0.018
Nickel	0.017	0.011
Fluoride	1.054	0.599
(i) Laundry Wastewat	er BAT	
Pollutant or	Maximum for	Maximum for
Pollutant Property		
mg/kg (lb/million lbs) of uranium pro	oduced by magnesium red
Chromium (total)	0.036	0.014
Copper	0.123	0.059
Nickel	0.053	0.036
Fluoride	3.360	1.910

NSPS are promulgated based on the performance achievable by the application of chemical precipitation, sedimentation, and multimedia filtration technology. The following effluent standards are promulgated for new sources:

(a) <u>Refinery Sump</u> Filtrate NSPS

	ant or	Maximum for	Maximum for
	Property	Any One Day	Monthly Average
mg/kg	(lb/million	lbs) of uranium	processed in the refinery
Chromium		27.140	11.000
Copper		93.880	44.740
Nickel		40.340	27.140
Fluoride		2,567.000	1,459.000
TSS		1,100.000	880.100
pH		the range of 7.5	to 10.0 at all times

(b) Slag Leach Reslurry NSPS

	tant or	Maximum for	Maximum for
	Property	Any One Day	Monthly Average
mg/kg	(lb/million	lbs) of uranium	processed in the refinery
Chromium	· · ·	1.689	0.685
Copper		5.844	2.785
Nickel		2.511	1.689
Fluoride		159.800	90.860
TSS		68.490	54.790
pH		e range of 7.5 t	o 10.0 at all times

(c) <u>Digestion</u> <u>Wet</u> <u>Air</u> <u>Pollution</u> <u>Control</u> NSPS

Pollutant or Pollutant Property		Maximum for Any One Day	Maximum for Monthly Average		
mg/kg	(lb/million	lbs) of uranium	processed in the refinery		
Chromium Copper Nickel Fluoride TSS pH		0.000 0.000 0.000 0.000 0.000 e range of 7.5 to	0.000 0.000 0.000 0.000 0.000 0.000 0 10.0 at all times		

(d) Solvent Extraction Raffinate Filtrate NSPS Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average mg/kg (lb/million lbs) of uranium processed in the refinery Chromium (total) 2.357 0.955 8.152 3.885 Copper Nickel 3.503 2.357 Fluoride 222.900 126.700 TSS 95.540 76.430 Within the range of 7.5 to 10 at all times Hα Evaporation and Denitration Wet Air Pollution Control (e) NSPS Pollutant or Maximum for Maximum for Pollutant Property Any One Day Monthly Average mg/kg (lb/million lbs) of uranium trioxide produced Chromium (total) 0.000 0.000 0.000 0.000 Copper 0.000 0.000 Nickel Fluoride 0.000 0.000 TSS 0.000 0.000 Within the range of 7.5 to 10.0 at all times рH f) Hydrofluorination Water Scrubber NSPS

Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million	lbs) of uranium	tetrafluoride produced
Chromium (total)	0.000	0.000
Copper	0.000	0.000
Nickel	0.000	0.000
Fluoride	0.000	0.000
TSS	0.000	0.000
pH Within the	range of 7.5 to 1	10.0 at all times

(g) Hydrofluorination	Alkaline Scru	bber NSPS	
Pollutant or Pollutant Property			
mg/kg (lb/million	lbs) of uraniu	m tetrafluoride pro	duced
-	0.007 0.026 0.011 0.700 0.300 ange of 7.5 to	0.003 0.012 0.007 0.398 0.240 10.0 at all times	
(h) <u>Magnesium</u> <u>Reducti</u>	on and Casting	Floor Wash NSPS	,
Pollutant or Pollutant Property			<u> </u>
mg/kg (lb/million lbs)	of uranium pr	oduced by magnesium	reduction
Chromium (total) Copper Nickel Fluoride TSS pH Within the r	0.011 0.039 0.017 1.054 0.452 ange of 7.5 to	0.005 0.018 0.011 0.599 0.361 10.0 at all times	· · ·
(i) Laundry Wastewate	r NSPS	, i i i i i i i i i i i i i i i i i i i	
Pollutant or Pollutant Property	Any One Day		
mg/kg (lb/million lbs)	of uranium pr	oduced by magnesium	reduction
Chromium (total) Copper Nickel Fluoride TSS pH Within the r	0.036 0.123 0.053 3.360 1.440 ange of 7.5 to	0.014 0.059 0.036 1.910 1.152 10.0 at all times	
PSES is not being pro	mulgated for t	nis subcategory at 1	this time

PSES is not being promulgated for this subcategory at this time because there are no existing indirect dischargers in the secondary uranium subcategory.

PSNS are promulgated based on the performance achievable by the application of chemical precipitation, sedimentation, and multimedia filtration technology. The following pretreatment standards are promulgated for new sources:

(a) <u>Refinery</u> <u>Sump</u> <u>Filtrate</u> PSNS

1			
Maximum for	Maximu	m for	
Any One Day	Monthly	Average	
bs) of uranium	processed	in the ref	iner
27.140	11.0	00	
ry PSNS			
Maximum for	Maximu	m for	
Any One Day	Monthly	Average	
		in the ref	
.DS) OF UTANIUM	processed	in the rer	iner
1.689	0.6	85	
5.844	2.7		
2.511	1.6	89	
2.511 159.800 Pollution Cont	90.8	60	·
159.800	90.8	60 m for	x
159.800 Pollution Cont Maximum for	90.8 rol PSNS Maximu Monthly	60 m for Average	iner
159.800 Pollution Cont Maximum for Any One Day bs) of uranium	90.8 rol PSNS Maximu Monthly processed	60 m for Average in the ref	iner
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.000	90.8 rol PSNS Maximu Monthly processed 0.0	60 m for Average in the ref 00	iner
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.000 0.000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0	60 m for Average in the ref 00 00	iner
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.000	90.8 rol PSNS Maximu Monthly processed 0.0	60 m for Average in the ref 00 00 00	iner
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.000 0.000 0.000 0.000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0 0.0 0.0 0.0	60 m for Average in the ref 00 00 00 00 S	iner
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.	60 m for Average in the ref 00 00 00 00 00 S m for	iner
159.800 <u>Pollution Cont</u> <u>Maximum for</u> Any One Day bs) of uranium 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0 0.0 0.0 0.0	60 m for Average in the ref 00 00 00 00 00 S m for	iner
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.	60 m for Average in the ref 00 00 00 00 S m for Average	
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.0000 0.00000 0.00000 0.00000 0.00000 0.00000 0.00000 0.00000 0.00000 0.0000000 0.00000000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.	60 m for Average in the ref 00 00 00 00 S m for Average in the ref	
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.0000 0.00000 0.00000 0.00000 0.00000 0.00000 0.000000 0.00000000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.	60 m for Average in the ref 00 00 00 00 S m for Average in the ref 55	
159.800 Pollution Cont Maximum for Any One Day bs) of uranium 0.0000 0.00000 0.00000 0.00000 0.00000 0.00000 0.000000 0.00000000	90.8 rol PSNS Maximu Monthly processed 0.0 0.0 0.0 0.0 trate PSN Maximu Monthly processed 0.9	60 m for Average in the ref 00 00 00 00 S m for Average in the ref 55 85	
	bs) of uranium 27.140 93.880 40.340 2,567.000 Ty PSNS Maximum for Any One Day bs) of uranium 1.689	Any One DayMonthlybs) of uranium processed27.14093.88044.740.34027.12,567.0001,459.0ryPSNSMaximum for Any One DayMonthlybs) of uranium processed1.6890.6	Any One DayMonthly Averagebs) of uranium processed in the ref27.14093.88044.74040.34027.1402,567.0001,459.000ryPSNSMaximum for Any One DayMonthly Averagebs) of uranium processed in the ref1.6890.685

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(e) Evaporation and	Denitration Wet	Air Pollution Control PSNS
Pollutant or	Maximum for	Maximum for
Pollutant Property		
mg/kg (lb/mil	lion lbs) of ura	nium trioxide produced
Chromium (total)	0.000	0.000
Copper	0.000	0.000
Nickel	0 000	0 000
Fluoride	0.000	0.000
(f) <u>Hydrofluorinatio</u>	on <u>Water</u> Scrubbe	r PSNS
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million	n lbs) of uraniu	m tetrafluoride produced
Chromium (total)	0.000	0.000
Copper	0.000	0.000
Nickel	0.000	0.000
Fluoride	0.000	0.000
(g) <u>Hydrofluorinatic</u> Pollutant or	on Alkaline Scru Maximum for	bber PSNS Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million	n lbs) of uraniu	m tetrafluoride produced
Chromium (total)	0.007	0.003
Copper	0.026	0.018
Nickel	0.011	0.007
Fluoride	0.700	0.398
(h) <u>Magnesium</u> <u>Reduct</u>	tion and Casting	Floor Wash PSNS
Pollutant or	Maximum for	Maximum for
Pollutant Property	Any One Day	Monthly Average
mg/kg (lb/million lbs	s) of uranium pro	oduced by magnesium reduction
Chromium (total)	0.011	0.005
Copper	0.039	0.018
Nickel	0.017	0.011
Fluoride	1.054	0.599
Fluoride	1.054	0.599

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(i) Laundry Wastewater PSNS

Pollutant or Pollutant Property	Maximum for Any One Day		
mg/kg (lb/million lbs)	of uranium	produced by magnesium redu	ction
Chromium (total) Copper Nickel Fluoride	0.036 0.123 0.053 0.360	0.014 0.059 0.036 1.910	

EPA is not promulgating BCT at this time for the secondary uranium subcategory.

SECTION III

SUBCATEGORY PROFILE

This section of the secondary uranium supplement describes the raw materials and processes used in producing secondary uranium and presents a profile of the secondary uranium plants identified in this study.

Secondary uranium is processed domestically as two general types of materials; slightly enriched with approximately 0.95 percent U_{235} , and depleted uranium with approximately 0.20 percent U_{235} . Natural grade uranium contains approximately 0.70 percent U_{235} . The slightly enriched material is processed at one facility operated by the U.S. Department of Energy. This material is used to fabricate fuel cores for "production reactors" which are used to produce plutonium.

The major use of depleted uranium is in ordinance applications. The source of depleted uranium is depleted uranium hexafluoride, UF_6 resulting from enrichment of natural uranium for nuclear applications. The high density and pyrophoricity of uranium metal reduced from depleted UF_6 make it ideal for use in armor penetrating ammunition. Other uses of secondary uranium are containers for spent nuclear reactor residues, radiation shielding applications, ballast and counterweights on aircraft control surfaces, and research.

DESCRIPTION OF SECONDARY URANIUM PRODUCTION

The production of secondary uranium can be divided into two distinct stages. The first stage is production of uranium tetrafluoride, UF4, from secondary materials, and the second stage is magnesium reduction of uranium tetrafluoride to pure uranium metal. All the plants in this subcategory perform the second stage process, but only one plant produces uranium tetrafluoride from secondary materials. The secondary uranium production processes are shown schematically in Figures III-1 and III-2 (pages 4674 and 4675), and are described in the following paragraphs.

RAW MATERIALS

The raw material necessary for the production of uranium by the magnesium reduction process is uranium tetrafluoride, UF₄. This material is generally obtained from enrichment plants which produce uranium for nuclear energy applications. The enrichment process involves separation of enriched UF6 from depleted UF₆. Much of the depleted uranium hexafluoride is converted to UF4 which is subsequently used as a raw material in the magnesium reduction process. Uranium tetrafluoride is also produced from uranium-bearing scrap. One of the plants in this subcategory uses uranium scrap (mainly off-spec product or machining scrap),

residues, and magnesium reduction slag as raw materials in addition to using uranium tetrafluoride. The following discussions describe the production of uranium from secondary sources and the production of uranium metal from uranium tetrafluoride in more detail.

URANIUM TETRAFLUORIDE PRODUCTION

One plant in the secondary uranium subcategory has the capacity ure uranium tetrafluoride from scrap This plant uses the manufactured UF4 to manufacture uranium materials. in its magnesium reduction operation as a supplement to UF_A obtained from other sources. This process is primarily a uranium recovery operation, as the raw materials are scrap from machining operations, and slag generated by magnesium reduction. The magnesium fluoride slag is recycled to the recovery process whenever its residual uranium content is economically recoverable.

The first step in the recovery process is acid leaching the raw materials to dissolve uranium. Nitric acid is used in all digestion, leaching, and dissolving operations. The resultant uranyl nitrate solution is filtered and undissolved solids are discarded.

Solvent extraction follows the dissolution operation. In the solvent extraction process, uranyl nitrate is extracted into a solvent phase from the impure solution with an organic solvent such as tributyl phosphate in kerosene. The solvent extraction raffinate is discharged to treatment.

Following the solvent extraction operation the uranyl nitrate is stripped from the organic phase with deionized water. The aqueous uranyl nitrate solution undergoes evaporation to produce a dry uranyl nitrate product which is calcined causing the nitrate to burn off as gaseous nitrogen oxides. The resulting product is yellow uranium trioxide, UO₃.

The final stages of uranium tetrafluoride production involve two operations; hydrogen reduction and hydrofluorination. Uranium trioxide is reduced by hydrogen to produce uranium dioxide, UO₂. Hydrogen for this process is obtained by cracking ammonia. Then, uranium dioxide is contacted with vaporized hydrofluoric acid at elevated temperatures. The resulting product is uranium tetrafluoride, UF₄, which is feed material for the magnesium reduction operation.

The potential waste streams associated with the production of uranium tetrafluoride are generated in the preliminary acid leaching steps and the solvent extraction and purification operations. Wet air pollution controls are also used in this process to scrub gases from the acid leaching, evaporation and denitration, and hydrofluorination operations.

MAGNESIUM REDUCTION PROCESS

The magnesium reduction process is widely used to produce uranium metal from uranium tetrafluoride. Uranium tetrafluoride is mixed with magnesium and reduced to uranium metal in a thermite-type bomb reduction vessel. The reduction reaction requires about three minutes and reaches a temperature around 1,900°C. After the magnesium fluoride slag and uranium metal are allowed to cool, the uranium metal is mechanically separated from the slag. No process water is associated with this process, therefore no waste streams are generated.

PROCESS WASTEWATER SOURCES

Although a variety of processes are involved in secondary uranium production, the process wastewater sources can be subdivided as follows:

- (a) Refinery sump filtrate,
- (b) Slag leach reslurry,
- (c) Digestion wet air pollution control,
- (d) Solvent extraction raffinate filtrate,
- (e) Evaporation and denitration wet air pollution control,
- (f) Hydrofluorination water scrubber,
- (g) Hydrofluorination alkaline scrubber,
- (h) Magnesium reduction and casting floor wash, and
- (i) Laundry wastewater.

OTHER WASTEWATER SOURCES

Other wastewaters may be associated with the secondary uranium subcategory. These wastewater streams include stormwater runoff, and maintenance and cleanup water. These waste streams are not considered as a part of this rulemak-ing. EPA believes that the flows and pollutant loadings associated with these waste streams are insignificant relative to the waste streams selected and are best handled by the appropriate permit authority on a case-bycase basis under authority of Section 402 of the Clean Water Act.

AGE, PRODUCTION, AND PROCESS PROFILE

Figure III-3 (page 4676) shows the location of the three secondary uranium plants operating in the United States. All three plants are on the eastern part of the country. Table III-1 (page 4671) shows the relative ages of the three plants. This shows that two plants were built in the early years of the uranium industry, while the third plant was built in the early It was probably built in anticipation of the growth of 1)70's. the uranium industry due to commercial uses of uranium, primarily in power generation. Table III-2 (page 4672) gives the yearly production ranges for the three plants in this subcategory.

Table III-3 (page 4673) provides a summary of the number of plants generating wastewater for the waste streams associated with various processes and the number of plants with the process.

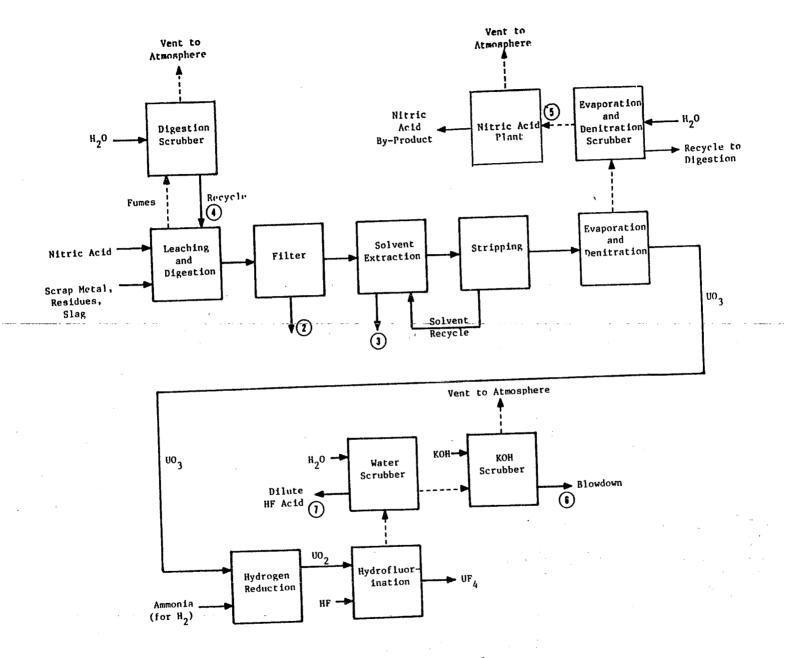
Table III-1

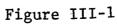
INITIAL OPERATING YEAR (RANGE) SUMMARY OF PLANTS IN THE SECONDARY URANIUM SUBCATEGORY BY DISCHARGE TYPE

Initial Operating Year (Range) Summary of Plants (Plant Age in Years)						
Type of Plant <u>Discharge</u>	1983- 1974 (0-10)	1973- 1969 <u>(11-15)</u>	1968- 1959 <u>(16-25)</u>	1958- 1954 <u>(26-30)</u>	1953- 1949 <u>(31-35)</u>	<u>Total</u>
Direct	0	1	0	0	1	2
Indirect		Q	0			. 0
Zero	<u>0</u>	<u>0</u>	<u>0</u>	<u>1</u>	<u>0</u>	<u>1</u>
TOTAL	0	. 1	. 0	1	1	3

SECT - III

SECONDARY URANIUM SUBCATEGORY





URANIUM TETRAFLUORIDE PRODUCTION PROCESS IN THE SECONDARY URANIUM SUBCATEGORY SECONDARY URANIUM SUBCATEGORY SECT

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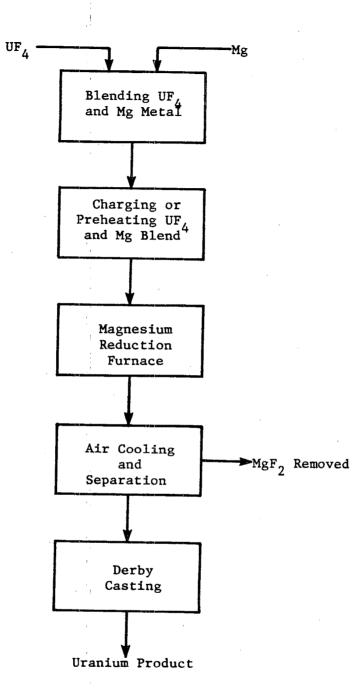
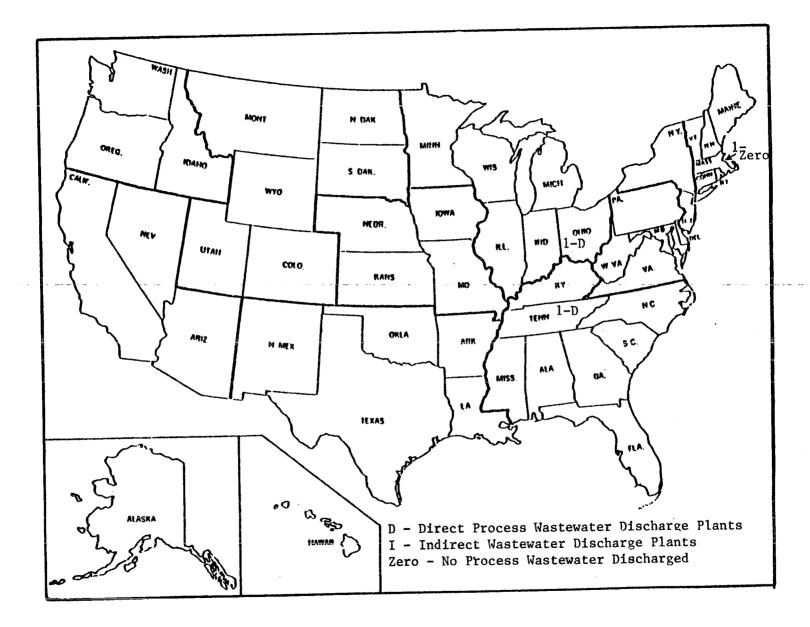


Figure III-2

MAGNESIUM REDUCTION PROCESS IN THE SECONDARY URANIUM SUBCATEGORY



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Figure III-3

GEOGRAPHIC LOCATIONS OF THE SECONDARY URANIUM SUBCATEGORY PLANTS

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SECTION IV

SUBCATEGORIZATION

This section summarizes the factors considered during the designation of the related subdivisions of the secondary uranium subcategory. Production normalizing parameters for each subdivision are also discussed.

FACTORS CONSIDERED IN SUBDIVIDING THE SECONDARY URANIUM SUBCATEGORY

The factors listed previously under general subcategorization were each evaluated when considering subdivision of the secondary uranium subcategory. In the discussion that follows, the factors will be described as they pertain to this particular subcategory.

The rationale for considering further subdivision of the secondary uranium subcategory is based primarily on differences in the production processes and raw materials used. Within this subcategory a number of different operations are performed which may or may not have a water use or discharge, and which may require the establishment of separate effluent limitations. While secondary uranium is still considered a single subcategory, a more thorough examination of the production processes has illustrated the need for limitations and standards based on specific flow allowances for the following subdivisions:

- 1. Refinery sump filtrate,
- 2. Slag leach reslurry,
- 3. Digestion wet air pollution control,
- 4. Solvent extraction raffinate filtrate,
- 5. Evaporation and denitration wet air pollution control,
- 6. Hydrofluorination water scrubber,
- 7. Hydrofluorination alkaline scrubber,
- 8. Magnesium reduction and casting floor wash, and
- 9. Laundry wastewater.

These subdivisions follow directly from differences within the process of refining scrap, residues, and slag to produce uranium tetrafluoride for use in magnesium reduction to uranium metal.

Leaching of the raw materials gives rise to the first, second, and fourth subdivisions. A major source of wastewater is the filtrate that is generated by leaching uranium from the raw materials. When slag is processed, the residual solids are discharged as a slurry which may be a significant source of pollutants. Wastewater from scrubbers which are used to control acid fumes in the leaching operation is also a source of pollutants.

Solvent extraction is used in the refining process to purify a uranium intermediate product. Solvent extraction results in a raffinate waste stream that contains significant quantities of pollutants.

Subdivisions five through seven arise from wet air pollution controls which control emissions from the process used to refine scrap, residues, and slag to a usable product. Evaporation, denitration, and hydrofluorination are all operations that necessitate air pollution control systems. In some cases, water use is recycled to the process rather than discharged. The potential sources of wastewater and associated pollutants require that each subdivision be examined and handled on an individual basis. Subdivisions eight and nine result from floor washing in the magnesium reduction and casting area and laundering of plant personnel clothing.

OTHER FACTORS

The other factors considered in this evaluation either support the establishment of the seven subdivisions or were shown to be inappropriate bases for subdivision. Air pollution control methods, treatment costs, and total energy requirements are functions of the selected subcategorization factors -- metal product, raw materials, and production processes. Therefore, they are not independent factors and do not affect the subcategorization which has been applied. As discussed in Section IV of the General Development Document, certain other factors, such as plant age, plant size, and the number of employees, were also evaluated and determined to be inappropriate for use as bases for subdivision of nonferrous metals plants.

PRODUCTION NORMALIZING PARAMETERS

As discussed previously, the effluent limitations and standards developed in this document establish mass limitations on the discharge of specific pollutant parameters. To allow these regulations to be applied to plants with various production capacities, the mass of pollutant discharged must be related to a unit of production. This factor is known as the production normalizing parameter (PNP).

In general, for each production process which has a wastewater associated with it, the actual mass of uranium intermediate product produced will be used as the PNP. Thus, the PNPs for the nine subdivisions are as follows:

Subdivision

PNP

1.	Refinery sump filtrate	kkg of uranium processed in the refinery	
2.	Slag leach reslurry	kkg of uranium processed in the refinery	
3.	Digestion wet air pollution control	kkg of uranium processed in the refinery	

- 4. Solvent extraction raffinate filtrate
- 5. Evaporation and denitration wet air pollution control
- 6. Hydrofluorination water scrubber
- 7. Hydrofluorination alkaline scrubber
- 8. Magnesium reduction and casting floor wash

kkg of uranium processed in the refinery

kkg of uranium trioxide produced

kkg of uranium tetrafluoride produced

kkg of uranium tetrafluoride produced

kkg of uranium produced by magnesium reduction

9. Laundry wastewater

kkg of uranium produced by magnesium reduction

The last two subdivisions were added after proposal to account for additional waste streams documented by the plants in their comments. For proposal, the Agency had insufficient data to quantify the flow associated with these operations. Data obtained during post proposal sampling visits have enabled the Agency to promulgate discharge allowances for these building blocks.

Based on comments from the industry received between proposal and promulgation, the Agency revised the production normalizing parameters for the first four subdivisions. The PNPs were changed from the mass of uranium trioxide produced to the mass of uranium processed within the subdivision operation. This enables plants to calculate their regulatory discharge allowances when they perform operations on a batch or campaign basis.

Several subdivision names have also been altered since proposal. This was done in response to industry comments requesting that subdivision names be modified to reflect actual practice within the industry.

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SECTION V

WATER USE AND WASTEWATER CHARACTERISTICS

This section describes the characteristics of the wastewaters associated with the secondary uranium subcategory. Water use and discharge rates are explained and then summarized. Data used to characterize the wastewaters are presented, and the specific source, water use and discharge flows, and wastewater characteristics for each separate wastewater source are discussed.

Section V of the General Development Document contains a detailed description of the data sources and methods of analysis used to characterize wastewater from the nonferrous metals manufacturing category. To summarize this information briefly, two principal data sources were used: data collection portfolios (dcp) and field sampling results. Data collection portfolios contain information regarding wastewater flows and production levels.

In order to quantify the pollutant discharge from secondary uranium plants, a field sampling program was conducted after proposal. A complete list of the pollutants considered and a summary of the techniques used in sampling and laboratory analyses are included in Section V of Vol. I. Wastewater samples were analyzed for 124 of the 126 priority pollutants and other pollutants deemed appropriate. Because the analytical standard for TCDD was judged to be too hazardous to be made generally available, samples were never analyzed for this pollutant. Samples were also never analyzed for asbestos. There is no reason to expect that TCDD or asbestos would be present in nonferrous metals manufacturing wastewater. Two plants were selected for sampling in the subcategory. In general, the samples were analyzed for three classes of pollutants: toxic pollutants, toxic metal pollutants, and organic criteria pollutants (which includes both conventional and nonconventional pollutants).

Additional wastewater flow and production data were received through industry comments and sampling activities between proposal and promulgation. This aided EPA in promulgating discharge allowances for magnesium reduction and casting floor wash and laundry wastewater which had not previously been proposed. It also aided EPA in revising the production normalized flows for several subdivisions.

As described in Section IV of this supplement, the secondary uranium subcategory has been split into nine subdivisions or wastewater sources, so that the proposed regulation contains mass discharge limitations and standards for nine unit processes discharging process wastewater. Differences in the wastewater characteristics associated with these subdivisions are to be expected. For this reason, wastewater streams corresponding to each subdivision are addressed separately in the discussions that follow. These wastewater sources are:

- 1. Refinery sump filtrate,
- 2. Slag leach reslurry,
- 3. Digestion wet air pollution control,
- 4. Solvent extraction raffinate filtrate,
- 5. Evaporation and denitration wet air pollution control,
- 6. Hydrofluorination water scrubber,
- 7. Hydrofluorination alkaline scrubber,
- 8. Magnesium reduction and casting floor wash, and
- 9. Laundry wastewater.

WASTEWATER FLOW RATES

Data supplied by dcp responses were evaluated, and two flow-toproduction ratios, water use and wastewater discharge flow, were calculated for each stream. The two ratios are differentiated by the flow value used in calculation. Water use is defined as the volume of water or other fluid required for a given process per mass of uranium product and is therefore based on the sum of recycle and make-up flows to a given process. Wastewater flow discharged after pretreatment or recycle (if these are present) is used in calculating the production normalized flow -- the volume of wastewater discharged from a given process to further treatment, disposal, or discharge per mass of uranium produced. Differences between the water use and wastewater flows associated with a given stream result from recycle, evaporation, and The production values used in carry-over on the product. calculation correspond to the production normalizing parameter, assigned to each stream, as outlined in Section IV. As an PNP, example, refinery sump filtrate wastewater flow is related to the mass of uranium processed in the refinery. As such, the discharge rate liters of refinery sump filtrate per metric ton of uranium processed in the refinery (gallons of refinery sump filtrate per ton of uranium processed in the refinery).

The production normalized discharge flows were compiled and statistically analyzed by stream type. These production normalized water use and discharge flows are presented by subdivision in Tables V-1 through V-9 at the end of this section. Where appropriate, an attempt was made to identify factors that could account for variations in water use and discharge rates. These variations are discussed later in this section by subdivision. A similar analysis of factors affecting the wastewater flows is presented in Sections X, XI, and XII where representative BAT, NSPS, and pretreatment flows are selected for use in calculating the effluent limitations.

The water use and discharge rates shown do not include nonprocess wastewater, such as rainfall runoff and noncontact cooling water.

WASTEWATER CHARACTERISTICS DATA

Data used to characterize the various wastewaters associated with secondary uranium production come from two sources -- data

collection portfolios and analytical data from sampling.

DATA COLLECTION PORTFOLIOS

In the data collection portfolios, the secondary uranium plants were asked to specify the presence or absence of toxic pollutants in their wastewater. The one plant responding to this questionnaire did not report the presence of any toxic organic pollutants. The responses for the toxic metals and cyanide are summarized below:

		Believed Present
		(Based on Raw Materials and
Pollutant	Known Present	Process Chemicals Used)
	······································	
Antimony	0	0
Arsenic	0	Q
Beryllium	0	0
Cadmium	0	0
Chromium	1	· 0
Copper	1	0
Cyanide	0	0
Lead	0	0
Mercury	0	0
Nickel	1	0
Selenium	0	0
Silver	0	0
Thallium	0	0
Zinc	0	0

FIELD SAMPLING DATA

In order to quantify the concentrations of pollutants in wastewater from secondary uranium plants, wastewater samples were collected at two of the plants belonging to this subcategory. Diagrams indicating the sampling sites, waste streams and production processes are shown in Figures V-1 and V-2 (pages 4726 and 4727).

Tables V-10 through V-16 summarize the data for 124 priority pollutants as well as other pollutants that were considered appropriate to this subcategory.

Table V-10 (page 4691) presents the data for refinery sump Tables V-11 and V-12 (pages 4694 and 4702) summarize filtrate. the data for solvent extraction raffinate after lime addition and sedimentation (V-11), and after additional treatment consisting of pH adjustment and filtration (V-12). Tables V-13 (page 4706), V-14 (page 4709), and V-15 (page 4719) show the data for hydrofluorination alkaline scrubber wastewater, reduction and casting floor wash, and laundry wastewater, respectively. Finally, Table V-16 (page 4723) presents data for treated wastewater from reduction and casting as well as machining Note that the stream numbers listed in the tables operations. correspond to those given in the plant sampling site diagrams,

Figures V-1 and V-2 (pages 4727 and 4727). Where no data are listed for a specific day of sampling, the wastewater samples for the stream were not collected.

The data tables include some samples measured at concentrations considered not quantifiable. Metal values reported as less than a certain value were considered not quantifiable.

The detection limits shown on the data tables for metals and conventional and nonconventional pollutants are not the same in all cases as the published detection limits for these pollutants by the same analytical methods. The detection limits used were reported with the analytical data and hence are the appropriate as a result of a number of laboratory-specific, equipmentspecific, and daily operator-specific factors. These factors can include day-to-day differences in machine calibration, variation in stock solutions, and variation in operators.

The statistical analysis of data includes some samples measured concentrations considered not quantifiable. For at data considered as detected but below quantifiable concentrations, а value of zero is used for averaging. Nonconventional and conventional pollutant data reported with a "less than" sign are considered as detected, but not further quantifiable. A value of zero is used for averaging. Metal values reported as less than a certain value were considered as below quantification, and consequently were assigned a value of zero in the calculation of the average.

WASTEWATER CHARACTERISTICS AND FLOWS BY SUBDIVISION

Since secondary uranium production involves nine principal sources of wastewater and each has potentially different characteristics and flows, the wastewater characteristics and discharge rates corresponding to each subdivision will be described separately. A brief description of why the associated production processes generate a wastewater will also be discussed.

REFINERY SUMP FILTRATE

The source of this waste stream is in the refinery digestion operation. Here the uranium scrap, residues, and compounds are acid leached, dissolving the uranium into solution. The primary sources of wastewater in the digestion and dissolving operations are pump leakage, pump seal water, spills, and hosedown water. The latter is required for health and safety reasons. The production normalized water use and discharge rates for refinery sump filtrate are given in Table V-1 (page 4688) in liters per metric ton of uranium processed in the refinery.

The sampling data for refinery sump filtrate are presented in Table V-10 (page 4691). The data show that this wastewater is characterized by treatable concentrations of chromium, magnesium, and suspended solids.

SLAG LEACH RESLURRY

This waste stream originates in the refinery digestion operation. Magnesium fluoride slag containing residual levels of uranium is acid leached to recover the uranium values. After leaching, the undissolved solids are filtered and discharged to treatment as a slurry. The production normalized water use and discharge rates for slag leach reslurry are given in Table V-2 (page 4688) in liters per metric ton of uranium processed in the refinery.

Although no sampling data are available for this wastewater, it is assumed to be similar to the refinery sump filtrate with treatable concentrations of magnesium and suspended solids.

DIGESTION WET AIR POLLUTION CONTROL

The acid leach operation, at the start of the uranium scrap, residue, and slag refining process, includes a water scrubbing system to control the discharge of acidic fumes and particulate matter. The scrubber liquor is recycled within the scrubber system, but a blowdown stream prevents build-up of acid and particulates. The blowdown stream is reused in the acid digestion and dissolution operation. Table V-4 (page 4689) shows the production normalized water use and discharge rates for digestion wet air pollution control.

Because the scrubber liquor is entirely recycled and reused, no discharge of wastewater results from the use of digestion wet air pollution control.

SOLVENT EXTRACTION RAFFINATE FILTRATE

Solvent extraction follows the acid leaching operation and is used to purify the uranium compound in solution. An organic solvent, tributyl phosphate in a kerosene carrier, selectively extracts the uranium compound from an acid solution. The solvent extraction raffinate filtrate is discharged to treatment. Table V-3 (page 4688) presents the production normalized water use and discharge rates for the solvent extraction raffinate filtrate in liters per metric ton of uranium processed in solvent extraction.

Although the Agency was not able to obtain samples of this wastewater prior to treatment, the data in Table V-11 (page 4694) for solvent extraction raffinate after lime addition and sedimentation show that this wastewater contains concentrations of several priority metals far in excess of their treatable concentrations. These metals include antimony, chromium, copper, lead, selenium, and zinc. Treatable concentrations of magnesium, uranium, and suspended solids are also present.

EVAPORATION AND DENITRATICS WET AIR POLLUTION CONTROL

A water scrubber is used to control vapors and fumes from the evaporation and denitration operations. Evaporation is used to

concentrate the uranium solution (uranyl nitrate) after it has been stripped from the organic phase into an aqueous phase. After evaporation, the concentrated intermediate uranium product is calcined to drive off the nitrate bound to the uranium and to produce dry uranium trioxide. The nitrates in the air react to form nitric acid, and the scrubber is used to control these acid fumes. Table V-5 (page 4689) shows the production normalized water use and discharge rates for the evaporation and denitration fume scrubber.

Because the scrubber liquor has a high acid content, it is recycled for use in the digestion operation. There it is used to dilute fresh acid used for leaching and dissolution. Since the scrubber liquor is entirely reused, no discharge of wastewater is practiced in the evaporation and denitration operations.

HYDROFLUORINATION WATER SCRUBBER

The hydrofluorination unit produces uranium tetrafluoride by contacting uranium dioxide with vaporized hydrofluoric acid at elevated temperatures. The off-gases from this operation contain significant quantities of unreacted hydrofluoric acid. The scrubber on this unit scrubs the acid fumes from the operation by absorbing the hydrofluoric acid into water. Scrubbed gases are vented to the alkaline scrubber. Table V-7 (page 4690) shows the production normalized water use and discharge rates in liters per metric ton of uranium tetrafluoride produced.

Since the hydrofluorination scrubber cleans what is predominantly vaporized unreacted hydrofluoric acid, the scrubber liquor concentrates this acid as it is recycled through the system. When the desired concentration of hydrofluoric acid is attained, the liquor is drawn off and sold for industrial use. For this reason, no discharge of wastewater occurs from the hydrofluorination water scrubber.

HYDROFLUORINATION ALKALINE SCRUBBER

This scrubber handles vent gases from the hydrofluorination water scrubber. These gases originated in the hydrofluorination operation where uranium dioxide is converted to uranium tetrafluoride. Hydrofluoric acid fumes that were not absorbed by the water scrubber are cleaned and neutralized by the KOH scrubber prior to venting the exhaust gases to the atmosphere. Scrubber blowdown is discharged to treatment. Production normalized water use and discharge rates are presented in Table V-6 (page 4689) in liters per metric ton of uranium tetrafluoride produced.

The sampling data for hydrofluorination alkaline scrubber wastewater are presented in Table V-13 (page 4706). These data show that this wastewater is characterized by an alkaline pH and treatable concentrations of arsenic, copper, and nickel.

MAGNESIUM REDUCTION AND CASTING FLOOR WASH WATER

Water is used to wash floors and equipment in the magnesium reduction and casting area. This water is eventually discharged as a wastewater stream. Table V-8 (page 4690) presents the production normalized water use and discharge rates for magnesium reduction and casting floor wash water in liters per metric ton of uranium produced by magnesium reduction.

The analytical data for this waste stream are presented in Table V-14 (page 4709). The data show that this wastewater is characterized by treatable concentrations of copper, lead, zinc, magnesium, uranium, and suspended solids.

LAUNDRY WASTEWATER

Water is used to wash the clothing of plant personnel working in process areas. This practice helps to minimize the amount of uranium which leaves the plant site on workers and their clothes. This water is eventually discharged as a wastewater stream. Water use and discharge rates for laundry wastewater are presented in Table V-9 (page 4790) in liters per metric ton of uranium produced by magnesium reduction.

The analytical data for this waste stream are presented in Table V-15 (page 4719).

Table V-l

WATER USE AND DISCHARGE RATES FOR REFINERY SUMP FILTRATE

(1/kkg of uranium processed in the refinery)

Plant Code	Percent Recycle	Production Normalized Water <u>Use</u>	Production Normalized Discharge <u>Flow</u>
1175	0	73,340	73,340

TABLE V-2

WATER USE AND DISCHARGE RATES FOR SLAG LEACH RESLURRY

(1/kkg of uranium processed in the refinery)

Plant Code	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge <u>Flow</u>
1175	0	4,566	4,566
	· .		

TABLE V-3

WATER USE AND DISCHARGE RATES FOR DIGESTION WET AIR POLLUTION CONTROL

(l/kkg of uranium processed in the refinery) Production Production Normalized Percent Normalized Discharge <u>Plant Code Recycle Water Use Flow</u> 1175 100 NR 0

TABLE V-4

WATER USE AND DISCHARGE RATES FOR SOLVENT EXTRACTION RAFFINATE FILTRATE

(1/kkg of uranium processed in the refinery)

Plant Code	Percent Recycle	Production Normalized Water <u>Use</u>	Production Normalized Discharge <u>Flow</u>
1175	0	6,369	6,369
	÷		

TABLE V-5

WATER USE AND DISCHARGE RATES FOR EVAPORATION AND DENITRATION WET AIR POLLUTION CONTROL

(1/kkg of uranium trioxide produced)

<u>Plant</u> Code	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge <u>Flow</u>
1175	100	NR	0
、	r		
		·	

TABLE V-6

WATER USE AND DISCHARGE RATES FOR HYDROFLUORINATION WATER SCRUBBER

(1/kkg of uranium tetrafluoride produced)

	<u>Plant</u> Code	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge <u>Flow</u>
,	1175	100	NR	0
			•	

TABLE V-7

WATER USE AND DISCHARGE RATES FOR HYDROFLUORINATION ALKALINE SCRUBBER

(1/kkg of uranium tetrafluoride produced)

<u>Plant</u> Code	Percent Recycle	Production Normalized Water <u>Use</u>	Production Normalized Discharge <u>Flow</u>
1175	NR	NR	20

TABLE V-8

WATER USE AND DISCHARGE RATES FOR MAGNESIUM REDUCTION AND CASTING FLOOR WASH WATER

(1/kkg of uranium produced by magnesium reduction)

<u>Plant</u> <u>Code</u>	Percent Recycle	Production Normalized Water Use	Production Normalized Discharge <u>Flow</u>
1175	0	331	331
1066	0	30.1	30.1

TABLE V-9

WATER USE AND DISCHARGE RATES FOR LAUNDRY WASTEWATER

(1/kkg of uranium produced by magnesium reduction)

<u>Plant</u> Code	Percent Recycle	Production Normalized Water <u>Use</u>	Production Normalized Discharge <u>Flow</u>
1175	NR	NR	NR
1066	0	192	192

Table V-10

REFINERY SUMP FILTRATE SAMPLING DATA

•	Pollutant	Stream Code	Sample Type*	<u>Conc</u>	<u>centration</u> Day 1	ns (mg/1) Day 2	Day 3	
<u>Toxic Po</u>	<u>llutants</u>					<u></u>	V	ノコン
114. an	timony	124	1	<0.02	0.08	0.11	0.033	//////////////////////////////////////
115. ar	senic	124	- 1	<0.002	<0.002	<0.002	Z0 000	
117. be	ryllium	124	1	<0.001	<0.001	<0.001	0.001 AN (0.002 GRAN]
118. ca	dmium	124	1	<0.002	<0.002	<0.002	<0.002 ⊠	
119. ch	romium (total)	124	1	<0.005	0:034	0.067	0.072 tu	j
120. co	pper	124	1	<0.001	0.027	0.028	0.072 BCATEGORY 0.027 TEGORY	2
121. су	anide (total)	124	1	<0.01	0.18	0.01	0.01 g	
122. le	ad	124	1	<0.002	<0.002	<0.002	<0.002	i
123. me:	rcury	124	1	0.0003	0.0003	0.0002	0.0002 m	2 1
124. ni	ckel	124	1	<0.003	<0.003	<0.003	<0.003 H	
125. se	lenium	124	1, .	<0.08	<0.08	0.10	<0.08 <) I
126. si	lver	124	1	<0.002	<0.002	<0.002	<0.002	
127. the	allium	124	1	<0.09	<0.09	<0.09	<0.09	
128. zim	nc	124	1	<0.003	0.006	0.017	0.019	

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REFINERY SUMP FILTRATE SAMPLING DATA

	Stream	Sample	Concentrations (mg/1)
Pollutant	Code	Type*	Source Day 1 Day 2 Day 3
Nonconventional Pollutants			
Acidity	124	1	114 23 <10 <10
Alkalinity	124	1	17 18 35 19
Ammonia Nitrogen	124	1	0.03 0.51 1.0 0.97
Chloride	124	1	13 22 34 28
Fluoride	124	1	0.3 7.7 6.2 2.3
Iron	124	1	0.37 0.079 0.044 0.12
Magnesium	124	1	9.6 560 1,700 1,700
Manganese	124	1	0.002 0.002 0.007 0.006
Nitrate	124	1	0.11 1,600 4,300 4,900
Sulfate	124	1	74 78 78 <11
Total Dissolved Solids (TDS)	124	1	160 2,900 13,000 12,000
Total Organic Carbon (TOC)	124	. 1	<1 6 12 23
Total Solids (TS)	124	1	200 2,900 15,000 12,000
Uranium	124	. 1	0.013 0.026 0.036 0.028

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SECONDARY URANIUM SUBCATEGORY SECT -

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REFINERY SUMP FILTRATE SAMPLING DATA

Pollutant	Stream Code	Sample <u>Type</u> *	<u>Cor</u> Source	Day 1	ons (mg/1) Day 2	Day 3
Conventional Pollutants						
Oil and Grease	124	1	14	290	140	41
Total Suspended Solids (TSS)	124	1	<1	16	35	45
pH	124	. 1	9.3	8.4	9.7	9.6

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 One-time grab
 Manual composite during intermittent process operation
 8-hour manual composite
 8-hour automatic composite
 - 24-hour manual composite
 - 24-hour automatic composite *Sample Type Code:

Table V-11

SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

<u>Pollutant</u>	Stream Code	Sample Type*	Conc Source	entration Day 1	s (mg/l) Day 2	Day 3	SEC
<u>Toxic Pollutants</u>							SECONDARY
1. acenaphthene	135	1	ND	ND			ARY
2. acrolein	135	1	ND	ND			UR
3. acrylonitrile	135	1	ND	ND			URANIUM
4. benzene	135	1	ND	ND			
5. benzidine	135	1	ND	ND			UBC
6. carbon tetrachloride	135	1	ND	ND			SUBCATEGORY
 chlorobenzene 	135	1	ND	ND			30RY
 8. 1,2,4-trichlorobenzene 	135	1	ND	ND			
 9. hexachlorobenzene 	135	1	ND	ND			SECT
	135	1	ND	ND			CH H
	135	1	ND	ND	. .		4
11. 1,1,1-trichloroethane		•	ND	ND			
12. hexachloroethane	135	1		•			
13. 1,1-dichloroethane	135	1	ND	ND			•
14. 1,1,2-trichloroethane	135	1	ND	ND			

SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

	<u>Pollutant</u>	Stream <u>Code</u>	Sample Type*	<u>Conc</u> Source	entration Day 1	<u>s (mg/1)</u> Day 2	Day 3	70
Toxic Po	ollutants (Cont.)							SECONDARY
15. 1,	1,2,2-tetrachloroethane	135	1	ND	ND			NDA:
16. ch	loroethane	135	1	ND	ND			
17. bi	s(chloromethyl)ether	135	1	ND	ND			URANIUM
18. bi	s(2-chloroethyl)ether	135	1	ND	ND			ΤUM
19. 2-	chloroethyl vinyl ether	135	1	ND	ND			SUE
20. 2-	chloronaphthalene	135	1	ND	ND			SUBCATEGORY
21. 2,	4,6-trichlorophenol	135	1	ND	ND		,	EGOI
22. p-	chloro-m-cresol	135	1	ND	ND			ХХ
23. ch	loroform	135	1	ND	ND			ល
24. 2-	chlorophenol	135	1	ND	ND			SECT
25. 1,	2-dichlorobenzene	135) 1	ND	ND -		. ж . К	- √
26. 1,	3-dichlorobenzene	135	1	ND	ND			
27. 1,	4-dichlorobenzene	135	⁻ 1	ND	ND			
28. 3,	3'-dichlorobenzidine	135	1	ND	ND			

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SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

	<u>Pollutant</u>	Stream Code	Sample Type*	<u>Conc</u> Source	entrations Day 1	<u>mg/1)</u> Day 2	Day 3	70
<u>Toxic</u>	Pollutants (Cont.)							SECONDARY
29.	1,1-dichloroethylene	135	1	ND	ND			NDAF
30.	1,2- <u>trans</u> -dichloroethylene	135	1	ND	ND			
- 31.	2,4-dichlorophenol	- 135 -	· · · 1 . · · · · · ·	ND	. ND		a se ta anna a se	URANIUM
32.	1,2-dichloropropane	135	1	ND	ND			
33.	1,3-dichloropropene	135	1	ND	ND			SUBCATEGORY
34.	2,4-dimethylphenol	135	1	ND	ND			CATE
35.	2,4-dinitrotoluene	135	1	ND	ND			GOR
36.	2,6-dinitrotoluene	135	1	ND	ND			К
37.	1,2-diphenylhydrazine	135	1	ND	ND			SE
38.	ethylbenzene	135	1	ND	ND			SECT
39.	fluoranthene	135	1	ND	ND	r		۱ ۲
40.	4-chlorophenyl phenyl ether	135	1	ND	ND			
41.	4-bromophenyl phenyl ether	135	1	ND	ND	,		
42.	bis(2-chloroisopropyl)ether	135	. 1	ND	ND			

SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

×	Pollutant	Stream <u>Code</u>	Sample Type*	<u>Conc</u> Source	entration Day 1	<u>ns (mg/1)</u> Day 2	Day 3	-
Toxic	Pollutants (Cont.)							SEC
43.	bis(2-chloroethoxy)methane	135	1	ND	ND			SECONDARY
44.	methylene chloride	135	1	ND	ND			
45.	methyl chloride (chloromethane)	135	1	ND	ND			URAN
46.	methyl bromide (bromomethane)	135	1	ND	ND			URANIUM
47.	bromoform (tribromomethane)	135	1	ND	ND			
48.	dichlorobromomethane	135	1	ND	ND			SUBCATEGORY
49.	trichlorofluoromethane	135	1	ND	ND			EGO
50.	dichlorodifluoromethane	135	1	ND	ND			RY
51.	chlorodibromomethane	135	1	ND	ND .			ß
52,	hexachlorobutadiene	135	1	ND	ND			SECT
53,	hexachlorocyclopentadiene	135	1.	ND	ND	-		- V
54.	isophorone	135	1	ND	ND			
55.	naphthalene	135	1	ND	ND			
56.	nitrobenzene	135	1	ND	ND			

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SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

	Pollutant	Stream Code	Sample Type*	<u>Conc</u> Source	entrations Day 1	(mg/1) Day 2	Day 3
Toxic	Pollutants (Cont.)						
57.	2-nitrophenol	135	1	ND	ND		
58.	4-nitrophenol	135	1	ND	ND		
5.9.	2,4-dinitrophenol	135	1	ND	ND	• ··· .	
60.	4,6-dinitro-o-cresol	135	1	ND	ND		
61.	N-nitrosodimethylamine	135	1	ND	ND		
62.	N-nitrosodiphenylamine	135	1	ND	ND		
63.	N-nitrosodi-n-propylamine	135	1	ND	ND		
64.	pentachlorophenol	135	1	ND	ND		
65.	phenol	135	1	ND	ND		
66.	bis(2-ethylhexyl)phthalate	135	1	ND	ND		
67.	butyl benzyl phthalate	135	1	ND	ND		*
68.	di-n-butyl phthalate	135	- 1 -	ND	ND .		· ·
69.	di-n-octyl phthalate	135	1	ND	ND		
70.	diethyl phthalate	135	1	ND	ND		

SECONDARY URANIUM SUBCATEGORY

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SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

	Pollutant	Stream <u>Code</u>	Sample <u>Type*</u>	<u>Conc</u> Source	entration Day 1	ns (mg/1) Day 2	Day 3	70
Toxic	Pollutants (Cont.)				• •		-	SECONDARY
71.	dimethyl phthalate	135	1	ND	ND			NDAJ
72.	benzo(a)anthracene	135	1	ND	ND			
73.	benzo(a)pyrene	135	1	ND	ND			URANIUM
74.	benzo(b)fluoranthene	135	1	ND	ND	•. * •		IUM
75.	benzo(k)fluoranthene	135	1	ND	ND			SUB
76.	chrysene	135	1	ND	ND			SUBCATEGORY
77.	acenaphthylene	135	1	ND	ND			EGOF
78.	anthracene	135	1	ND	ND			Я
79.	benzo(ghi)perylene	135	1	ND	ND			ß
80.	fluorene	135	1	ND	ND			SECT
81.	phenanthrene	135	1	ND	ND			י ל
82.	dibenzo(a,h)anthracene	135	1	ND	ND			
83.	indeno(1,2,3-c,d)pyrene	135	1	ND	ND	•		
84.	pyrene	135	2 1 1 1 2 2	ND	ND			

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SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

	Pollutant	Stream Code	Sample Type*	Con Source	centrations Day 1	(mg/1) Day 2	Day 3	SE
<u>Toxic</u>	Pollutants (Cont.)							SECONDARY
85.	tetrachloroethylene	135	1	ND	ND			DARY
86.	toluene	135	1	ND	. ND			
87.	trichloroethylene	135	· · · · · · · · · · · · · · · · · · ·	ND	ND		• . • • • • • • • •	URANIUM
88.	vinyl chloride (chloroethylene)	135	1	ND	ND			
114.	antimony	135	1	<0.02	10			UBC
115.	arsenic	135	1	<0.002	<0.02			SUBCATEGORY
117.	beryllium	135	1	<0.001	0.17			JORY
118.	cadmium	135	_1	<0.002	1.2			
119.	chromium (total)	135	1	<0.005	25			SECT
120.	copper	135	1	<0.001	4,160			Hi.
122.	lead	135	1	<0.002	14		-	4
123.	mercury	135	···· 1	0.0003	0.011			
124.	nickel	135	· 1	<0.002	49	, ·		
125.	selenium	135	1	<0.08	13			

SOLVENT EXTRACTION RAFFINATE AFTER LIME ADDITION AND SEDIMENTATION SAMPLING DATA

Pollutant	Stream Code	Sample Type*	<u>Concentrations (mg/1)</u> Source Day 1 Day 2 Day 3	
Toxic Pollutants (Cont.)				SECONDARY
126. silver	135	1	<0.002 1.7	IDAR
127. thallium	135	1	<0.09 3	•
128. zinc	135	1	<0.003 42	URANIUM
Nonconventional Pollutants				MD
Iron	135	1	0.37 920	SUBCATEGORY
Magnesium	135	1	9.6 18,000	ATE
Manganese	135	1	0.002 55	GOR
Uranium	135	1	0.013 20.0	Ň
Conventional Pollutant				SECT
рН	135	1	9.3 8.3	CH H

- *Sample Type Code: 1 One-time grab 2 Manual composite during intermittent process operation 3 8-hour manual composite 4 8-hour automatic composite 5 24-hour manual composite 6 24-hour automatic composite

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Table V-12

SOLVENT EXTRACTION RAFFINATE FILTRATE SAMPLING DATA

Pollutant	Stream Code	Sample Type*	Conc Source	entration Day 1	<u>s (mg/1)</u> Day 2	Day 3	ß
<u>Toxic Pollutants</u>							ECON
114. antimony	123	1	<0.02	0.02	1.5 1.5	2.3	SECONDARY
115. arsenic	123	1	<0.002	<0.002	<0.002 <0.002	0.008	URANIUM
117. beryllium	123	1	<0.001	<0.001	<0.001 <0.001	<0.001	
118. cadmium	123	1	<0.002	0.020	0.008 0.010	<0.002	SUBCATEGORY
119. chromium (total)	123	1	<0.005	<0.005	0.083 0.087	0.086	GORY
120. copper	123	1	<0.001	0.18	0.096 0.10	1.8	SECT
121. cyanide (total)	123	1	<0.01	<0.01	<0.01 <0.01	<0.01	I
122, lead	123	1	<0.002	0.058	2.2 2.3	2.3	V
123. mercury	123	. 1	0.0003	0.0009	0.0003	0.0003	
124. nickel	123	• 1	<0.003	0.009	0.026 0.027	<0.003	

SOLVENT EXTRACTION RAFFINATE FILTRATE SAMPLING DATA

Pollutant	Stream Code	Sample Type*	<u>Cor</u> Source	<u>centratic</u> Day 1	ons (mg/1) Day 2) Day 3	-
Toxic Pollutants (Continued)				. .			SEC
125. selenium	123	1	<0.08	<0.08	0.36	<0.08	SECONDARY URANIUM
126. silver	123	. 1	<0.002	<0.002	0.069	0.008	Y URA
127. thallium	123	1	<0.09	<0.09	0.09	<0.09	
128. zinc	123	1	<0.003	0.011	0.017 0.017	0.091	SUBCATEGORY
Nonconventional Pollutants							TEGO
Acidity	123	1	114	340	<10 <10	<10	DRY
Alkalinity	123	1	17	<1	770 7 770	,800	SECT
Ammonium Nitrogen	123	1 1	0.03	0.08	0.46	0.19	י ע
Chloride	123	1	13	14	320 300	210	
Fluoride	123	1	0.3	1.2	7.8 8.1	31	

SOLVENT EXTRACTION RAFFINATE FILTRATE SAMPLING DATA

D 11	Stream	Sample		centration		Dour 3	
<u>Pollutant</u>	Code	<u>Type*</u>	Source	Day 1	Day 2	Day 3	
Nonconventional Pollutants (Continued)						SECO
Iron	123	1	0.37	0.19	0.12 0.11	0.057	SECONDARY
Magnesium	123	1	9.6	12	1.9	0.38	
Manganese	123	1	0.002	<0.001	0.008	<0.001	URANIUM ;
Nitrate	123	1	0.11		000 8 300	,100	SUBCATEGORY
Sulfate	123	1	74	310	240 230	4	regory
Total Dissolved Solids (TDS)	123	1	160	800 110, 110,		,900	·
Total Organic Carbon (TOC)	123	1.	<1	1	23 21	6	SECT -
Total Solids (TS)	123	1	200	940 120 120		,400	- V
Uranium	123	1	0.013	0.014	0.017 0.018	0.020	

4704

SOLVENT EXTRACTION RAFFINATE FILTRATE SAMPLING DATA

D 11	Stream	Sample	Concentrations (mg/1)			
Pollutant	Code	Type*	Source	Day 1	Day 2	Day 3
Conventional Pollutants		-				
Oil and Grease	123	1	14	9	10 2	3
Total Suspended Solids (TSS)	123	1	<1	<1 1	870 ,100	45 5
рН	123	1	9.3	9.4	11.5 11.5	13.1 g

*Sample Type Code:

- 1 One-time grab
 2 Manual composite during intermittent process operation
 3 8-hour manual composite
 4 8-hour automatic composite
 5 24-hour manual composite
 6 24-hour automatic composite

SECONDARY URANIUM SUBCATEGORY SECT ı.

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Table V-13

HYDROFLUORINATION ALKALINE (KOH) SCRUBBER RAW WASTEWATER SAMPLING DATA

	Pollutant	Stream Code	Sample Type*	Conc Source	entration Day 1	s (mg/1) Day 2	Day 3
<u>Toxic</u>	Pollutants						CONDA
114.	antimony	121	1	<0.02	0.35	0.39	Uns.
115.	arsenic	121	1	<0.002	12	14	
117.	beryllium	121		<0.001	0.003	0.003	
118.	cadmium	121	1	<0.002	0.073	0.055	
119.	chromium (total)	121	1	<0.005	0.063	0.075	
120.	copper	121	1	<0.001	2.0	3.6	.)
121.	cyanide (total)	121	1	<0.01	<0.01	0.01	
122.	lead	121	1	<0.002	0.011	0.003	F
123.	mercury	121	1	0.0003	0.0014	0.0020	C I
124.	nickel	121	1	<0.003	3.6	1.9	(
125.	selenium	121	1	<0.08	0.46	0.54	
126.	silver	121	.1 .	<0.002	0.10	0.12	
127.	thallium	121	1	<0.09	0.14	0.18	
128.	zinc	121	1	<0.003	0.31	0.30	

SECONDARY URANIUM SUBCATEGORY SECT -

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HYDROFLUORINATION ALKALINE (KOH) SCRUBBER RAW WASTEWATER SAMPLING DATA

<u>Pollutant</u>	Stream Code	Sample Type*	<u>Concentrations (mg/1)</u> Source <u>Day 1 Day 2 Day 3</u> 10
Nonconventional Pollutants			114
Acidity	121	1	114 NDA _F
Alkalinity	121	1	17
Ammonia Nitrogen	121	1.	0.03 3.8 ANIM 13 21 18
Chloride	121	1	
Fluoride	121	1	0.3 19,00032,000SUBCATE0.370.120.0799.60.650.78
Iron	121	1	0.37 0.12 0.079
Magnesium	121	1	9.6 0.65 0.78 ^G
Manganese	121	1	0.002 0.066 0.054
Nitrate	121	1	0.11 0.03 0.05 m
Sulfate	121	1.	0.11 0.03 0.05 они на
Total Dissolved Solids (TDS)	121	1	160 <
Total Organic Carbon (TOC)	121	1	<1 4,000
Total Solids (TS)	121	1 · · · ·	200
Uranium	121	1	0.013 0.015 0.014

4707

HYDROFLUORINATION ALKALINE (KOH) SCRUBBER RAW WASTEWATER SAMPLING DATA

	Stream	Sample	С	oncentrat	ions $(mg/1)$	
Pollutant	Code	<u>Type*</u>	Sourc	<u>e Day</u>	1 <u>Day 2</u>	Day 3 M
Conventional Pollutants	• P					OND
Oil and Grease	121	1	14	7,300	6,600	IDARY
Total Suspended Solids (TSS)	121	1	<1		·	
рН	121	1	9.3	14.8	15.3	UT NF

*Sample Type Code:	 1 - One-time grab 2 - Manual composite during intermittent process operation 3 - 8-hour manual composite 4 - 8-hour automatic composite 5 - 24-hour manual composite 6 - 24-hour automatic composite

Table V-14

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

	<u>Pollutant</u>	Stream Code	Sample Type*	<u>Conce</u>	<u>ntrations (mg/l)</u> <u>Day 1 Day 2</u>	Day 3	SI
Toxic	Pollutants			1. 		÷	SECONDARY
1.	acenaphthene	405	1	ND		ND	DARS
2.	acrolein	405	1	ND	• •	ND	
3.	acrylonitrile	405	. 1	ND	:	ND	URANIUM
4.	benzene	405	1	ND		ND	
5.	benzidine	405	1	ND		ND	SUBC
6.	carbon tetrachloride	405	1	ND		ND	ATE
7.	chlorobenzene	405	1	ND		ND	SUBCATEGORY
8.	1,2,4-trichlorobenzene	405	1	ND		ND	
9.	hexachlorobenzene	405	1	ND		ND	SECT
10.	1,2-dichloroethane	405	1	ND		ND	H
11.	1,1,1-trichloroethane	405	1	ND		ND	4
12.	hexachloroethane	405	1	ND		ND	
13.	1,1-dichloroethane	405	i	ND		ND	
14.	1,1,2-trichloroethane	405	1	ND		ND	

4709

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

	<u>Pollutant</u>	Stream <u>Code</u>	Sample Type*	Conce Source	entrations Day 1	(mg/1) Day 2	Day 3	SI
<u>Toxic</u>	Pollutants (Cont.)							ECON
15.	1,1,2,2-tetrachloroethane	405	1	ND		v	ND	SECONDARY
16.	chloroethane	405	1	ND			ND	
17.	bis(chloromethyl)ether	405	1	ND			ND	URANIUM
18.	bis(2-chloroethyl)ether	405	1	ND		· • · · · · · · · · · · · · · · · · · ·	ND	
19.	2-chloroethyl vinyl ether	405	1	ND			ND	SUBO
20.	2-chloronaphthalene	405	1	ND			ND	SUBCATEGORY
21.	2,4,6-trichlorophenol	405	1	ND			ND	GOR
22.	p-chloro-m-cresol	405	1	ND			ND	κ _ι
23.	chloroform	405	1	ND			ND	SECT
24.	2-chlorophenol	405	1	ND			ND	CT T
25.	1,2-dichlorobenzene	405	1	ND			ND	- <
26.	1,3-dichlorobenzene	405	1	ND			ND	
27.	1,4-dichlorobenzene	405	1	ND			ND	
28.	3,3'-dichlorobenzidine	405	1	ND			ND	

4710

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

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	Pollutant	Stream Code	Sample Type*	<u>Conc</u> Source	entrations (mg, Day 1 Day		Day :	<u>a</u>
<u>Toxi</u>	<u>c Pollutants</u> (Cont.)						<u></u>	SECO
29.	1,1-dichloroethylene	405	1	ND	·		ND	SECONDARY
30.	1,2- <u>trans</u> -dichloroethylene	405	1	ND			ND	
31.	2,4-dichlorophenol	405	. 1	ND		-	ND	JRAN
32.	1,2-dichloropropane	405	1	ND			ND	URANIUM
33.	1,3-dichloropropene	405	1	ND			ND	
34.	2,4-dimethylphenol	405	1	ND			ND	SUBCATEGORY
35.	2,4-dinitrotoluene	405	1	ND		•	ND	EGOI
36.	2,6-dinitrotoluene	405	1	ND			ND	RΥ
37.	1,2-diphenylhydrazine	405	1	ND			ND	S
38.	ethylbenzene	405	. 1	ND			ND	SECT
39.	fluoranthene	405	1	ND			ND	ו ל
40.	4-chlorophenyl phenyl ether	405	1	ND	-i		ND	
41.	4-bromophenyl phenyl ether	405	1	ND			ND	
42.	bis(2-chloroisopropyl)ether	405	1	ND			ND	

4711

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

		Stream	Sample	the second s	entrations		Day 3	
	Pollutant	Code	Type*	Source	Day 1	<u>Day 2</u>	Day J	SEC
<u>Toxic</u>	Pollutants (Cont.)							SECONDARY
43.	bis(2-chloroethoxy)methane	405	1	ND			ND	ARY
44.	methylene chloride	405	1	ND	,	A	ND	UR
45.	methyl chloride (chloromethane)	405	· 1· ····	ND and		·· ·· · · · ·	ND	URANIUM
46.	methyl bromide (bromomethane)	405	1	ND			ND	
	bromoform (tribromomethane)	405	1	ND			ND	SUBC
47.	dichlorobromomethane	405	1	ND			ND	SUBCATEGORY
48.		405	1	ND			ND	GOR
49.	trichlorofluoromethane	405	1	ND			ND	К
50.	dichlorodifluoromethane	·	I				ND	Ŋ
51.	chlorodibromomethane	405	1	ND				SECT
52.	hexachlorobutadiene	405	1	ND -			ND	i I
53.	hexachlorocyclopentadiene	405	1	ND			ND	4
54.	isophorone	405	.1	- ND -			ND	
	naphthalene	405	· · · 1	ND			ND	
55. 56.	naphthalene	405	1	ND			ND	

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REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

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Pollutant	Stream	Sample	Conc	entration	s (mg/1)		
TOTTALAIL	Code	Type*	Source	Day 1	Day 2	Day 3	_ v
Toxic Pollutants (Cont.)							
57. 2-nitrophenol	405	1	ND			ND	DECONDART
58. 4-nitrophenol	405	1	ND			ND	
59. 2,4-dinitrophenol	405	1	ND		•	ND	UND T NIGYO
60. 4,6-dinitro-o-cresol	405	1	ND			ND	LON
61. N-nitrosodimethylamine	405	1	ND			ND	SADORITEOOK X
62. N-nitrosodiphenylamine	405	1	ND			ND	C M T T
63. N-nitrosodi-n-propylamine	405	1	ND			ND	GON
64. pentachlorophenol	405	1	ND			ND	÷.
65. phenol	405	1	ND		۰.	ND	Ũ
66. bis(2-ethylhexyl)phthalate	405	1	ND			ND	DECT.
67. butyl benzyl phthalate	405	1	ND			ND	ו <
68. di-n-butyl phthalate	405	1	ND			ND	
69. di-n-octyl phthalate	405	1	ND		,	ND	
70. diethyl phthalate	405	1 1	ND			ND	
			e				

4713

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REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

Pollutant	Stream Code	Sample Type*	Conc Source	entration: Day 1	s (mg/1) Day 2	Day 3	SEC
Toxic Pollutants (Cont.)							OND
71. dimethyl phthalate	405	1	ND			ND	SECONDARY
72. benzo(a)anthracene	405	1	ND			ND	URI
73. benzo(a)pyrene	405		ND .			ND	URANIUM
74. benzo(b)fluoranthene	405	1	ND			ND	
75. benzo(k)fluoranthene	405	1	ND			ND	SUBCATEGORY
76. chrysene	405	1	ND			ND	ATEO
77. acenaphthylene	405	1	ND			ND	JORY
78. anthracene	405	1	ND			ND	••
79. benzo(ghi)perylene	405	1	ND			ND	SECT
80. fluorene	405	1	ND			ND	H
	405	1	ND			ND	<
81. phenanthrene	405	1	ND			ND	
82. dibenzo(a,h)anthracene		4				ND	
83. indeno(1,2,3-c,d)pyrene	405	1 · ·	ND				
84. pyrene	405	1	ND			ND	

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

	Pollutant	Stream Code	Sample Type*	<u>Con</u> Source	<u>centratio</u> Day 1	ns (mg/1) Day 2	Day 3	
Toxic	Pollutants (Cont.)			· .				SEC
85.	tetrachloroethylene	405	1	ND			ND	SECONDARY
86.	toluene	405	1	ND			ND	ARY
87.	trichloroethylene	405	1	ND			ND	URA
88.	vinyl chloride (chloroethylene)	405	1	ND			ND	URANIUM
114.	antimony	126 405	1 1	<0.02 <0.0006	0.55	0.54	<0.0006	
115.	arsenic	126 405	1	<0.002 <0.001	<0.002	<0.002	0.0028	SUBCATEGORY
117.	beryllium	126 405	1 1	<0.001 0.012	<0.01	<0.01	0.051	DRY
118.	cadmium	126 405	1	<0.002 <0.003	<0.02	<0.02	0.064	SECT
119.	chromium (total)	126 405	1 1	<0.005 0.061	0.065	<0.05	0.6	- V
120.	copper	126 405	1	<0.001 0.088	0.092	0.084	2.3	
121.	cyanide (total)	126 405	1 12	<0.01 <0.01	<0.01	≪0.01	<0.01	

4715 .

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

Pollutant	Stream Code	Sample Type*	<u>Conc</u> Source	entration Day 1	<u>s (mg/1)</u> Day 2	Day 3	ß
· ·		and all all all all all all all all all al					EC
<u>Toxic Pollutants</u> (Cont.)							ONI
122. lead	126 405	1 1	<0.002 0.036	0.089	0.071	4.1	SECONDARY
123. mercury	126 405	1 1	0.0003 <0.005	0.0002	<0.0002	<0.0005	URANIUM
124. nickel	126 405	1 1	<0.003 0.055	<0.03	<0.03	0.5	
125. selenium	126 405	1 1	<0.08 <0.001	0.87	0.88	0.0033	SUBCATEGORY
126. silver	126 405	1 1	<0.002 <0.0005	<0.02	<0.02	0.0008	JORY
127. thallium	126 405	1 1	<0.09 <0.001	<0.9	<0.9	<0.001	SECT
128. zinc	126 405	1 1	<0.003 0.101	0.32	0.38	4.0	Ч - V
Nonconventional Pollutants		•	•	•			·
Acidity	126 405	1	114 <10.0	23	110	<10	• ·
Alkalinity	126 405	1	17 33.0	100	79	618	

4716

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

Pollutant	Stream <u>Code</u>	Sample Type*	Co Source	ncentratio Day 1	ns (mg/ Day	1) 2 Day 3	- 70
Nonconventional Pollutants (Cont.)			X.) SECO
Ammonia Nitrogen	126 405	1 1	0.03	084	1.2	2.1	SECONDARY
Chloride	126 405	1	13 36.0	21	22	74	URANIUM
Fluoride	126 405	1 1	0.3 0.41	18	13	1.8	
Iron	126 405	1 1	0.37 0.16	5.4	3.9	48.0	SUBCATEGORY
Magnesium	126 405	.1	9.6 8.0	26	20	1,499.0	GORY
Manganese	126 405	1 1	0.002 0.058	0.047	0.037	2.3	SECT
Nitrate	126 405	1	0.11 <0.09	0.36	0.31	75	.1
Sulfate	126 405	1 1	74 2.8	160	740	8.8	V
Total Dissolved Solids (TDS)	126 405	1 1	160 300	310 13	,000	680	

REDUCTION AND CASTING FLOOR WASH RAW WASTEWATER SAMPLING DATA

Pollutant	Stream <u>Code</u>	Sample Type*	Con Source	<u>centrati</u> Day 1	ons (mg/ Day		<u>3</u> N
Toxic Pollutants (Cont.)							CONI
Total Organic Carbon (TOC)	126 405	1 1	<1 <10.0	740	1,900	2 ·	SECONDARY
Total Solids (TS)	126 405	1 1	200 330.0	340 1	5,000	-3;000-	3, MIUM
Uranium	126 405	1 1	0.013 0.089	0.054	0.05	9 79	
Conventional Pollutants							CAT
Oil and Grease	126 405	1 1	14 <1.0	380	440	25	SUBCATEGORY
Total Suspended Solids (TSS)	126 405	1 1	<1 <1.0	9	35	1,600	Ø
pH	126 405	1 1	9.3 6	7.4	7.6	9	SECT -

*Sample Type Code:

- 1 One-time grab
- 2 Manual composite during intermittent process operation
 3 8-hour manual composite
 4 8-hour automatic composite
 5 24-hour manual composite
 6 24-hour automatic composite

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Table V-15

URANIUM LAUNDRY/LAB WASTE RAW WASTEWATER SAMPLING DATA

	1 A.							
	Stream		<u> </u>					
Pollutant	<u>_Code</u>	<u>Type#</u>	Source	Day 1	Day 2	Day 3		
Toxic Pollutants		- ,			r			
114. antimony	908	1	<0.0006	<0.0006				
115. arsenic	908	1	<0.001	0.028				
117. beryllium	908	1 .	0.012	0.015		:		
418. cadmium	908	1	<0.03	<0.03				
119. chromium (total)	908	1	0.061	<0.03				
120. copper	908	1	0.088	0.25				
121. cyanide (total)	908	1	<0.01	<0.1				
122. lead	908	1	0.036	0.042				
123. mercury	908	· 1	<0.005	<0.005				
124. nickel	908	1	0.055	<0.03				
125. selenium	908	1	<0.001	<0.001	. •			
126. silver	908	1	<0.0005	0.0048	·			
127. thallium	908	- 1	<0.001	<0.001				
128. zinc	908	1	0.101	0.7		-		
		· .						

URANIUM LAUNDRY/LAB WASTE RAW WASTEWATER SAMPLING DATA

	Stream	Sample	Concentrations (mg/1)			
Pollutant	<u>Code</u>	Type#	Source	Day 1	Day 2	Day 3
Nonconventional Pollutants						
Acidity	908	1	<10.0	<10 ¹		,
Alkalinity	908	1	33.0	59		
Aluminum	908	1	0.131	0.9		
Ammonia Nitrogen	908	1	0.07	2.3		
Barium	908	1	0.2	0.2		
Boron	908	1	<0.2	0.3		:
Calcium	908	1	0.045	17.0		
Chemical Oxygen Demand (COD)	908	1	<50.0	<50		
Chloride	908	1	36.0	210		
Cobalt	908	1	0.044	0.25		
Fluoride	908	1 1	0.41	0.79		
Iron	908	1	0.16	0.16		
Magnesium	908	1	8.0	5.3		
Manganese	908	1	0.058	0.2		
Molybdenum	908	1	<0.03	<0.03		

SECONDARY URANIUM SUBCATEGORY SECT -

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URANIUM LAUNDRY/LAB WASTE RAW WASTEWATER SAMPLING DATA

		Sample	C	an the game in the descent of		
Pollutant	<u>Code</u>	<u>Type#</u>	Source	<u>Day 1</u>	Day 2	Day 3
Nonconventional Pollutants (Con	t.)					
Nitrate	908	1	<0.09	<0.09		
Phosphorus	908	1	0.5	12		-
Sodium	908	1	74.0	133.0		
Sulfate	908	1	2.8	14.		
Tin	908	1	<0.25	<0.2		
Titanium	908	1	<0.2	<0.2		
Total Dissolved Solids (TDS)	908	1	300.0	590		
Total Organic Carbon (TOC)	908	1	<10.0	46		
Total Solids (TS)	908	1	330.0	630	u ,	,
Uranium	908	1	0.89	0.51		
Vanadium	908	1	<0.03	<0.03		
Yttrium	908		<0.1	7.3		,
				ų		

SECONDARY URANIUM SUBCATEGORY SECT - V

URANIUM LAUNDRY/LAB WASTE RAW WASTEWATER SAMPLING DATA

	Stream	Sample	Concentrations (nCi/L)				
<u>Pollutant</u>	<u>Code</u>	Type#	Source	Day 1	Day 2	<u>Day 3</u>	
Nonconventional Pollutants (Co	nt.)	v					
Gross Alpha	908	1	0.014	13.7			
Gross Beta	908	1 .	<0.013	18.5			
Radium-226	908	1	<0.0008	3.6	- araa maa aanaa waa -	••••	
		•	Co	ncentration	s (mg/l)	·····	
Conventional Pollutants							
Oil and Grease	908	1	<1.0	42			
Total Suspended Solids (TSS)	908	. 1	<1.0	11			
pH (standard units)	908	1	б	6			

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*Sample Type Code: 1 - One-time grab

SECONDARY URANIUM SUBCATEGORY SECT

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Table V-16

PLANT 6 FILTRATE RAW WASTEWATER SAMPLING DATA

Pollutant		Stream	Sample	Con			
		Code	Type*	Source	Day 1	<u>ns (mg/l)</u> <u>Day 2</u>	Day 3
<u>Toxic</u>	e Pollutants	·					
114.	antimony	130	1	<0.02	0.14	0.083	
115.	arsenic	130	1	<0.002	<0.002	<0.002	
117.	beryllium	130	1	<0.001	<0.001	<0.001	
118.	cadmium	130	1	<0.002	0.017	<0.002	
119.	chromium (total)	130	1	<0.005	0.080	0.016	
120.	copper	130	1	<0.001	1.1	0.94	
121.	cyanide (total)	130	1	<0.01	0.30	0.54	
122.	lead	130	1	<0.002	0.21	0.005	
123.	mercury	130	1	0.0003	<0.0002	0.0004	
124.	nickel	130	1	<0.003	0.70	0.092	-
125.	selenium	130	1	<0.08	0.18	0.12	
126.	silver	130	1	<0.002	0.057	0.017	
127.	thallium	130	1	<0.09	<0.09	<0.09	
128.	zinc	130	1	<0.003	0.15	0.037	

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SECONDARY URANIUM SUBCATEGORY SECT - V

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PLANT 6 FILTRATE RAW WASTEWATER SAMPLING DATA

Pollutant	Stream <u>Code</u>	Sample Type*	Con Source	<u>Day 1</u>	ns (mg/1) Day 2	Day 3
Nonconventional Pollutants						
Acidity	130	1	114	20	<10	
Alkalinity	130	1	17	100	310	
Ammonia Nitrogen	130		0.03	. 1.7		
Chloride	130	1	13	380	130	
Fluoride	130	1	0.3	19	8.6	
Iron	130	1	0.37	4.3	0.31	
Magnesium	130	1	9.6	11	0.59	
Manganese	130	1	0.002	0.086	0.002	
Nitrate	130	1	0.11 2	0,000 11	,000	
Sulfate	130	1	74	63	91	
Total Dissolved Solids (TDS)	130	1	160 3	1,000 14	+,000	
Total Organic Carbon (TOC)	130	••• 1 ••	<1	73	66	
Total Solids (TS)	130	· · · 1	200 3	1,000 14	4,000	
Uranium	130	1	0.013	0.063	0.037	

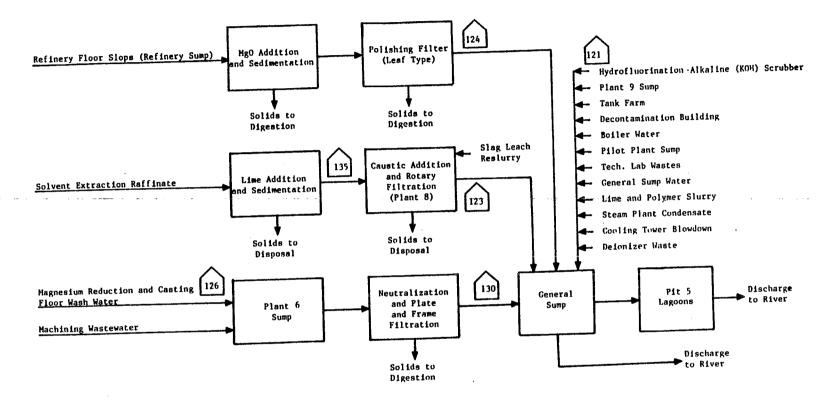
SECONDARY URANIUM SUBCATEGORY SECT - V

PLANT 6 FILTRATE RAW WASTEWATER SAMPLING DATA

Pollutant	Stream	Sample		ncentrati		
<u>Conventional Pollutants</u>	_Code	<u>Type*</u>	Source	Day 1	Day 2	Day 3
Oil and Grease	130	1	14	52	31	
Total Suspended Solids (TSS)	130	- 1	<1	170	100	
рН	130	1	9. 3	8.2	11.8	

*Sample Type Code:

- 1 One-time grab
- 2 Manual composite during intermittent process operation
 3 8-hour manual composite
 4 8-hour automatic composite
 5 24-hour manual composite
 6 24-hour automatic composite



NOTE: Flows indicated are in gallons per day

Figure V-1

SAMPLING LOCATIONS AT SECONDARY URANIUM PLANT A

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SECONDARY URANIUM SUBCATEGORY

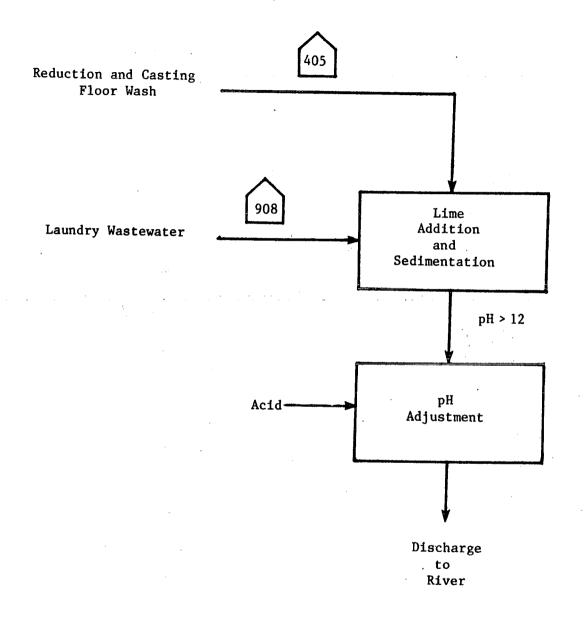


Figure V-2

SAMPLING LOCATIONS AT SECONDARY URANIUM PLANT B

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SECTION VI

SELECTION OF POLLUTANT PARAMETERS

This section examines the chemical analysis data presented in Section V and discusses the selection or exclusion of pollutants for potential limitation. The discussion that follows presents briefly discusses the selection of conventional and and pollutants for effluent limitations. nonconventional Also described is the analysis that was performed to select or exclude pollutants for further consideration toxic priority for limitations and standards. Pollutants will be considered for limitation if they are present in concentrations treatable by the technologies considered in this analysis. The treatable concentrations used for the priority metals were the long-term achievable by performance values chemical precipitation, sedimentation, and filtration. The treatable concentrations used for the priority organics were the long-term performance values achievable by carbon adsorption.

CONVENTIONAL AND NONCONVENTIONAL POLLUTANT PARAMETERS SELECTED

As part of this study, The Agency examined samples from two plants in this subcategory for toxic, conventional and nonconventional pollutants. The conventional and nonconventional pollutants or pollutant parameters selected for limitation in the secondary uranium subcategory are:

fluoride total suspended solids (TSS) pH

Fluoride was found in the two samples of hydrofluorination alkaline scrubber wastewater at concentrations of 19,000 mg/l and 32,000 mg/l, significantly higher than the 14.5 mg/l achievable with identified treatment technology. For this reason, fluoride is selected for limitation in this subcategory.

concentrations ranging from less than 1 to 1,600 mg/l TSS were observed in the raw waste samples analyzed for this study. Most these concentrations are well above the 2.6 mg/l treatable of concentration. Most of the specific methods used to remove priority metals do so by converting these metals to precipitates, and these metal-containing precipitates should not be discharged. Meeting a limitation on total suspended solids helps ensure that removal of these precipitated toxic metals has been effective. For these reasons, total suspended solids are selected for limitation in this subcategory.

The pH values observed during this study ranged from 6.0 to 15.3. These values suggest that the pH of secondary uranium wastewaters may be outside the 7.5 to 10.0 range considered desirable for discharge to receiving waters. Many deleterious effects are caused by extreme pH values or rapid changes in pH. Also, effective removal of toxic metals by precipitation requires careful control of pH. Since pH control within the desirable limits is readily attainable by available treatment, pH is selected for limitation in this subcategory.

Ammonia and uranium have been considered for regulation in this subcategory. Neither pollutant has been selected for regulation at promulgation. Data collected after proposal indicated that ammonia is no longer used in the secondary uranium processing. Therefore, EPA is not promulgating limitations for ammonia. Effluent limitations for uranium have not been established in this subcategory. Pursuant to the Supreme Court's ruling in Train v. Colorado Public Interest Research Group, 426 U.S. 1 (1976). EPA lacks the authority under the Clean Water Act to regulate this pollutant since it is a "source" material as defined by the Nuclear Regulatory Commission under the Atomic Energy Act.

TOXIC PRIORITY POLLUTANTS

The frequency of occurrence of the toxic pollutants in the raw wastewater samples taken is presented in Table VI-1 (page 4373). Table VI-1 is based on the raw wastewater data from streams 124, 135, 121, 126, 405, and 908 (see Section V). These data provide the basis for the categorization of specific pollutants, as discussed below. Treatment plant samples were not considered in the frequency count.

TOXIC POLLUTANTS NEVER DETECTED

The toxic pollutants listed in Table VI-2 (page 4736) were not detected in any raw wastewater samples from this subcategory. Therefore, they are not selected for consideration in establishing limitations:

TOXIC POLLUTANTS PRESENT BELOW CONCENTRATIONS ACHIEVABLE BY TREATMENT

The toxic pollutants listed below are not selected for consideration in establishing limitations because they were not found in any raw wastewater samples from this subcategory above concentrations considered achievable by existing or available treatment technologies.

117. beryllium
121. cyanide
123. mercury

TOXIC POLLUTANTS DETECTED IN A SMALL NUMBER OF SOURCES

The following pollutants were not selected for limitation on the basis that they were detected in the effluent from only a small number of sources within the subcategory, and are uniquely related to only these sources:

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66. bis(2-ethylhexyl) phthalate 127. thallium

Although these pollutants were not selected for limitation in establishing national guidelines, it may be appropriate on a case-by-case basis for the local permitting authority to specify effluent limitations.

Bis(2-ethylhexyl) phthalate was detected in one of two samples of secondary uranium raw wastewater analyzed. The observed concentration is 0.989 mg/l. This compound is a plasticizer commonly used in field sampling equipment and is not used or formed as a by-product in this subcategory. For this reason, bis(2-ethyl-hexyl) phthalate is not selected for further consideration for limitation.

Thallium was detected above its treatable concentration of 0.34 mg/l in only one out of 10 samples analyzed. The observed treatable concentration is 3 mg/l. The Agency has no reason to believe that treatable concentrations of thallium should be present in secondary uranium wastewaters and does not believe that the one observed treatable value is representative of the subcategory. For these reasons, thallium is not selected for further consideration for limitation.

PRIORITY POLLUTANTS SELECTED FOR FURTHER CONSIDERATION IN ESTABLISHING LIMITATIONS AND STANDARDS

The toxic pollutants listed below are selected for further consideration in establishing limitations and standards for this subcategory. The toxic pollutants selected for further consideration for limitation are each discussed following the list.

114. antimony
115. arsenic
118. cadmium
119. chromium
120. copper
122. lead
124. nickel
125. selenium
126. silver
128. zinc

Antimony was detected above its treatability concentration of 0.47 mg/l in three out of 10 samples analyzed. Antimony is selected for further consideration for limitation.

Arsenic was detected above its treatability concentration of 0.34 mg/l in two out of 10 samples. These samples showed 12.0 and 14.0 mg/l arsenic in the raw wastewater. Therefore, arsenic is selected for further consideration for limitation.

Cadmium was detected above its treatable concentration of 0.049

mg/l in four samples containing 1.2, 0.73, 0.055, and 0.064 mg/l. Cadmium is selected for further consideration for limitation.

Chromium was detected above its treatability concentration of 0.07 mg/l in three out of 10 samples analyzed. The analytical data showed 0.072, 0.075, and 25 mg/l chromium in the raw wastewater. Chromium is selected for further consideration for limitation.

Copper was detected above its treatability concentration of 0.39 mg/l in four out of 10 samples analyzed. The observed treatable concentrations ranged from 0.084 to 4,160 mg/l. Copper is selected for further consideration for limitation.

Lead was detected above its treatable concentration of 0.08 mg/l in three out of 10 samples analyzed. The samples indicated 0.089, 4.1, and 14 mg/l of lead in the raw wastewater. Lead is selected for further consideration for limitation.

Nickel was detected above its treatability concentration of 0.22 mg/l in four out of 10 samples analyzed. Nickel is selected for further consideration for limitation.

Selenium was detected above its treatability concentration of 0.20 mg/l in five out of 10 raw wastewater samples analyzed. Therefore, selenium is selected for further consideration for limitation.

Silver was detected above its treatability concentration of 0.07 mg/l in three out of 10 samples analyzed. Silver is selected for further consideration for limitation.

Zinc was detected above its treatability concentration of 0.23 mg/l in seven out of 10 samples analyzed. Zinc is selected for further consideration for limitation.

Table VI-1

FREQUENCY OF OCCURRENCE OF PRIORITY POLLUTANTS SECONDARY URANIUM SUBCATEGORY RAW WASTEWATER

	Pollutant	Analytical Quantification Concentration (mg/l)(a)	Treatable Concentra- tion (mg/l)(b)	Number of Streams Analyzed	Number of Samples Analyzed	ND	Detected Below Quantification Concentration	Detected Below Treat- able Concen- tration	Detected Above Treat- able Concen- tration
1.	acenaphthene	0.010	0.01	2	2	2		-	
2.	acrolein	0.010	0.01	2	2 `	2 2			
3.	acrylonitrile	0.010	0.01	2	2	ž			i
4.	benzene	0.010	0.01	<u>.</u> 2	2	2			
5.	benzidine	0.010	0.01	2	2	2			
6.	carbon tetrachloride	0.010	0.01	2	$\overline{2}$	$\overline{2}$			
7.	chlorobenzene	0.010	0.01	2	2	2			
8.	1,2,4-trichlorobenzene	0.010	0.01	2	2	$\overline{2}$			
. 9.	hexachlorobenzene	0.010	0.01	2	ž	2		1. 1. Mar. 1. M	
10.	1,2-dichlorobenzene	0.010	0.01	2	2	2			
11.	1,1,1-trichloroethane	0.010	0.01	2	$\overline{2}$	2			
12.	hexachloroethane	0.010	0.01	ž	$\overline{2}$	$\overline{2}$			i
13.	1,1-dichloroethane	0.010	0.01	2	2	$\overline{2}$			
14.	1,1,2-trichloroethane	0.010	0.01	2	2		•		
15.	1,1,2,2-tetrachloroethane	0.010	0.01	2	2	2 2			
16.	chloroethane	0.010	0.01	2	2	2			
17.	bis(chloromethyl) ether	0.010	0.01	2	2	2			
18.	bis(2-chloroethyl) ether	0.010	0.01	2	2	2 2			
19.	2-chloroethyl vinyl ether	0.010	0.01	2	2	2			
20.	2-chloronaphthalene	0.010	0.01	2	2				i
21.	2,4,6-trichlorophenol	0.010	0.01	2	2	2 2			
22.	parachlorometa cresol	0.010	0.01	2	2	2			
23.	chloroform	0.010	0.01	2	2	2			
24.	2-chlorophenol	0.010	0.01	2	2	2			
25.	1,2-dichlorobenzene	0.010	0.01	2	2	2 2			
26.	1,3-dichlorobenzene	0.010	0.01	2	2	2			·
27.	1,4-dichlorobenzene	0.010	0.01	2	2	2			
28.	3,3'-dichlorobenzidine	0.010	0.01	2	2	2			
29.	1,1-dichloroethylene	0.010	0.01	2 2	2	2			
30.	1,2- <u>trans</u> -dichloroethylene	0.010	0.01		2	2			1
31.	2,4-dichlorophenol	0.010	0.01	2	2	2			
32.	1,2-dichloropropane	0.010	0.01	2	2	2			
33.	1,3-dichloropropylene	0.010	0.01	2	2	2			
34.	2,4-dimethylphenol	0.010	0.01	2	2	2			
35.	2,4-dinitrotoluene	0.010	0.01	2	2	2			
36.	2,6-dinitrotoluene	0.010	0.01	2	2			-	
37.	1,2-diphenylhydrazine	0.010	0.01	2	2	2 2			
38.	ethylbenzene	0.010	0.01	2	2	2			
39.	fluoranthene	0.010	0.01	2	2	2			

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FREQUENCY OF OCCURRENCE OF PRIORITY POLLUTANTS SECONDARY URANIUM SUBCATEGORY RAW WASTEWATER

	Pollutant	Analytical Quantification Concentration (mg/l)(a)	Treatable Concentra- tion (mg/1)(b)	Number of Streams Analyzed	Number of Samples <u>Analyzed</u>	ND	Detected Below Quantification Concentration	Detected Below Treat- able Concen- tration	Detected Above Treat- able Concen-
40.	4-chlorophenyl phenyl ether	0.010	0.01	2	2	2			
41.	4-bromophenyl phenyl ether	0.010	0.01	2	2	2			
42.	bis(2-chloroisopropyl) ether	0.010	0.01	2	2	2			
43.	bis(2-chloroethoxy) methane	0.010	0.01	2	2	2			
44.	methylene chloride	0.010	0.01	2	2	2			
45.	methyl chloride	0.010	0.01	2	2	2			
46.	methyl bromide	0.010	0.01	2	2	2			
47.	bromoform	0.010	0.01	2	2	. 2			
48.	dichlorobromomethane	. 0.010	0.01	2	2	2			
49.	trichlorofluoromethane	0.010	0.01	2	2	2			
50.	dichlorodifluoromethane	0.010	0.01	2	2	2			
51.	chlorodibromomethane	0.010	0.01	2	2	2			
52.	hexachlorobutadiene	0.010	0.01	2	2	2 2			
53.	hexachlorocyclopentadiene	0.010	0.01	2	2	2			
54.	isophorone	0.010	0.01	2	2	2			
55.	naphthalene	0.010	0.01	2	2 2	2 2			•
56.	nitrobenzene	0.010	0.01	2	2	2			
57.	2-nitrophenol	0.010	0.01	2	2	2			
58.	4-nitrophenol	0.010	0.01	2	2 2	2 2			
59.	2,4-dinitrophenol	0.010	0.01	2	2	2			
60.	4,6-dinitro-o-cresol	0.010	0.01	2	2	2			
61.	N-nitrosodimethylamine	0.010	0.01	2	22	2			
62.	N-nitrosodiphenylamine	0.010	0.01	2	2	2			
63.	N-nitroso-n-propylamine	0.010	0.01	2	2	2			
64.	pentachlorophenol	0.010	0.01	2	2	2			
65.	phenol	0.010	0.01	2	2	2			
66.	bis(2-ethylhexyl) phthalate	0.010	0.01	2	2	1			
67.	butyl benzyl phthalate	0.010	0.01	2	2	2			
68.	di-n-butyl phthalate	0.010	0.01	2	2	2			
69.	di-n-octyl phthalate	0.010	0.01	2	2	2			
70.	diethyl phthalate	0.010	0.01	2	2	2			
71.	dimethyl phthalate	0.010	0.01	2	2	2			
72.	benzo(a)anthracene	0.010	0.01	2	2	2			
73.	benzo(a)pyrene	0.010	0.01	2	2	2			
74.	3,4-benzofluoranthene	0.010	0.01	2	2	2			
75.	benzo(k)fluoranthene	0.010	0.01	2	2	2			
76.	chrysene	0.010	0.01	2	2	2			
77.	acenaphthylene	0.010	0.01	2	2	2			
78.	anthracene	0.010	0.01	2	2	2			

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SECONDARY URANIUM SUBCATEGORY

FREQUENCY OF OCCURRENCE OF PRIORITY POLLUTANTS SECONDARY URANIUM SUBCATEGORY RAW WASTEWATER

	Pollutant	Analytical Quantification Concentration (mg/1)(a)	Treatable Concentra- tion (mg/1)(b)	Number of Streams Analyzed	Number of Samples Analyzed	<u>ND</u>	Detected Below Quantification Concentration	Detected Below Treat- able Concen- tration	Detected Above Treat- able Concen- tration
79.	benzo(ghi)perylene	0.010	0.01	2	2	2			
80.	fluorene	0.010	0.01	$\overline{2}$	$\overline{2}$	2			
81.	phenanthrene	0.010	0.01	$\tilde{2}$	2	2			
82.	dibenzo(a, h)anthracene	0.010	0.01	2	$\overline{2}$	2			
83.	indeno(1,2,3-c,d)pyrene	0.010	0.01	2	$\overline{2}$	2			
84.	pyrene	0.010	0.01	$\overline{2}$	2	2	· · · · · · · · ·		
85.	tetrachloroethylene	0.010	0.01	2	2	2			
86.	toluene	0.010	0.01	2	2	2			
87.	trichloroethylene	0.010	0.01	2	2	$\overline{2}$			
88.	vinyl chloride	0.010	0.01	2	$\overline{2}$	$\overline{2}$			
114.	antimony	0.100	0.47	5	10	-	4	3	3
115.	arsenic	0.010	0.34	5	10		i	ĭ	2
117.	beryllium	0.010	0.20	5	10		7		4
118.	cadmium	0.002	0.049	5	10		6	3	4
119.	chromium	0.005	0.07	5	10		2	5	3
120.	copper	0.009	0.39	5	10		-	6	4
121.	cyanide	0.02	0.047	4	9		8	ĩ	7
122.	lead	0.020	0.08	5	10		5	2	3
123.	mercury	0.0001	0.036	5	10		3	7	5
124.	nickel	0.005	0.22	5	10		6	•	4
125.	selenium	0.01	0.20	5	10		4	1	Š
126.	silver	0.02	0.07	5.	10		7		3
127.	thallium	0.100	0.34	5	10		7	2	1
128.	zine	0.050	0.23	5	10		3	-	7
129.	2,3,7,8-tetrachlorodibenzo- p-dioxin (TCDD)	Not Analyzed					-		

(a) Analytical quantification concentration was reported with the data (see Section V).

(b) Treatable concentrations are based on performance of chemical precipitation, sedimentation, and filtration.

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- VI

TABLE VI-2

TOXIC POLLUTANTS NEVER DETECTED

1.	acenaphthene
2.	acrolein
3.	acrylonitrile
4.	benzene
5.	benzidine
б.	carbon tetrachloride (tetrachloromethane)
7.	chlorobenzene
8.	1,2,4-trichlorobenzene
9.	hexachlorobenzene
10.	1,2-dichloroethane
11.	1,1,1-trichloroethane
12.	hexachloroethane
13.	1,1-dichloroethane
14.	1,1,2-trichloroethane
15.	1,1,2,2-tetrachloroethane
16.	chloroethane
17.	bis (chloromethyl) ether (deleted)
18.	bis (2-chloroethyl) ether
19.	2-chloroethyl vinyl ether (mixed)
20.	2-chloronaphthalene
21.	2,4,6-trichlorophenol
22.	parachlorometa cresol
23.	chloroform (trichloromethane)
2?.	2-chlorophenol
25.	1,2-dichlorobenzene
26.	1,3-dichlorobenzene
27.	1,4-dichlorobenzene
28.	3,3'-dichlorobenzidine
29.	1,1-dichloroethylene
30.	1,2-trans-dichloroethylene
31.	2,4-dichlorophenol
32.	1,2-dichloropropane
33.	1,2-dichloropropylene (1,3-dichloropropene)
34.	2,4-dimethylphenol
35.	2,4-dinitrotoluene
36.	2,6-dinitrotoluene
37.	1,2-diphenylhydrazine
38.	ethylbenzene
39.	fluoranthene
40.	4-chlorophenyl phenyl ether
41.	4-bromophenyl phenyl ether
42.	bis(2-chloroisopropyl) ether
43.	bis(2-choroethoxy) methane
44.	methylene chloride (dichloromethane)
45.	methyl chloride (chloromethane)
46.	methyl bromide (bromomethane)
	bromoform (tribromomethane)
47.	
48.	dichlorobromomethane
49.	trichlorofluoromethane (deleted)

TABLE VI-2 (Continued)

TOXIC POLLUTANTS NEVER DETECTED

50. dichlorodifluoromethane (deleted) 51. chlorodibromomethane hexachlorobutadiene 52. 53. hexachlorocyclopentadiene 54. isophorone 55. naphthalene 56. nitrobenzene 57. 2-nitrophenol 58. 4-nitrophenol 59. 2,4-dinitrophenol 60. 4,6-dinitro-o-cresol 61. N-nitrosodimethylamine 62. N-nitrosodiphenylamine 63. N-nitrosodi-n-propylamine 64. pentachlorophenol 65. phenol 67. butyl benzyl phthalate 68. di-n-butyl phthalate 69. di-n-octyl phthalate 70. diethyl phthalate 71. dimethyl phthalate 72. benzo (a)anthracene (1,2-benzanthracene) 73. benzo (a)pyrene (3,4-benzopyrene) 74. 3,4-benzofluoranthene 75. benzo(k)fluoranthane (11,12-benzofluoranthene) 76. chrysene 77. acenaphthylene 78. anthracene 79. benzo(ghi)perylene (1,11-benzoperylene) 80. fluorene 81. phenanthrene 82. dibenzo (a,h)anthracene (1,2,5,6-dibenzanthracene)
83. indeno (1,2,3-cd)pyrene (w,e,-o-phenylenepyrene) 84. pyrene 85. tetrachloroethylene 86. toluene 87. trichloroethylene 88. vinyl chloride (chloroethylene) 89. aldrin* 90. dieldrin* 91. chlordane (technical mixture and metabolites)* 92. 4,4'-DDT* 93. 4,4'-DDE(p,p'DDX)*4,4'-DDD(p,p'TDE)* 94. 95. a-endosulfan.Alpha* 96. b-endosulfan-8eta* 97. endosulfan sulfate* 98. endrin* 99. endrin aldehyde*

TABLE VI-2 (Continued)

TOXIC POLLUTANTS NEVER DETECTED

100. heptachlor* 101. heptachlor epoxide* 102. a-Alpha-BHC* 103. b-Beta-BHC* 104. r-Gamma-BHC(lindane)* 105. g-Delta-BHC* 106. PCB-1242 (Arochlor 1242)* 107. PCB-1254 (Arochlor 1254)* 108. PCB-1221 (Arochlor 1221)* 109. PCB-1232 (Arochlor 1232)* 110. PCB-1248 (Arochlor 1248)* 111. PCB-1260 (Arochlor 1260)* 112. PCB-1016 (Arochlor 1016)* 113. toxaphene* 116. asbestos (Fibrous) 129. 2,3,7,8-tetra chlorodibenzo-p-dioxin (TCDD)

*We did not analyze for these pollutants in samples of raw wastewater from this subcategory. These pollutants are not believed to be present based on the Agency's best engineering judgment which includes consideration of raw materials and process operations.

SECONDARY URANIUM SUBCATEGORY SECT - VII

SECTION VII

CONTROL AND TREATMENT TECHNOLOGIES

The preceding sections of this supplement discussed the sources, flows, and characteristics of the wastewaters from the secondary uranium plants. This section summarizes the description of these wastewaters and indicates the treatment technologies which are currently practiced in the secondary uranium subcategory for each wastewater stream. Also, this section presents the control and treatment technology options which were examined by the Agency for possible application to the secondary uranium subcategory.

CURRENT CONTROL AND TREATMENT PRACTICES

This section presents a summary of the control and treatment technologies that are currently being applied to each of the sources generating wastewater in this subcategory. As discussed Section V, wastewater associated with the secondary uranium in subcategory is characterized by the presence of the toxic metal pollutants and suspended solids. This analysis is supported by the raw (untreated) wastewater data, presented in Section V, from It is expected that these pollutants are uranium ore mill. present in each of the waste streams at concentrations above treatability, and these waste streams are commonly combined for Construction of one wastewater treatment system for treatment. combined treatment allows plants to take advantage of economic scale and in some instances to combine streams of different The direct alkalinity to reduce treatment chemical requirements. discharging plant in this subcategory currently has a combined wastewater treatment system including chemical precipitation and The options selected for consideration for BPT, sedimentation. BAT, NSPS, and pretreatment will be summarized toward the end of this section.

REFINERY SUMP FILTRATE

Refinery sump filtrate wastewater has its source in the digestion operation. Components of this wastewater include pump leakage, pump seal water, spills, and washdown water. All these flows are collected in a sump to which chemicals are added to precipitate uranium. After filtration with a leaf filter, the filtrate is discharged to a general sump for treatment consisting of neutralization, flocculation and sedimentation, and discharge to a surface water.

SLAG LEACH RESLURRY

In addition to solid uranium scrap and uranium residues, magnesium fluoride slag from the magnesium reduction operation is used as a raw material for uranium recovery. The recovery process involves acid leaching the slag to dissolve uranium. Separation of the uranium-containing acid and the leached slag is done by filtration, after which the slag solids are reslurried with water. The slurry is treated by neutralization and rotary precoat filtration. The filtrate is discharged to a surface water.

DIGESTION WET AIR POLLUTION CONTROL

The acid leaching operation includes a water scrubber for control of acid fumes generated from leaching. The system recirculates water to absorb particulates and acid gases, and a blowdown stream prevents build-up of acid and particulates. The blowdown stream is reused in the acid digestion and dissolution operation. Since the scrubber liquor is entirely recycled and reused, no discharge to the treatment system occurs.

SOLVENT EXTRACTION RAFFINATE FILTRATE

Purification of the uranium compound that results from acid leaching is done by solvent extraction. An organic solvent is used to selectively extract the uranium compound from the acid solution. Impurities from acid leaching are left in the acid solution. This solvent extraction raffinate filtrate is discharged to combined treatment consisting of neutralization and sedimentation, followed by discharge to a surface water.

EVAPORATION AND DENITRATION WET AIR POLLUTION CONTROL

After purification by solvent extraction, the uranyl nitrate solution is concentrated by evaporation. The calcination step which follows converts the uranium compound to uranium trioxide. Calcination off-gases contain much nitric acid. Since the scrubber liquor absorbs the nitric acid, the liquor is not discharged as a wastewater but used to dilute fresh acid in the digestion operation. Therefore, no wastewater is discharged from the evaporation and denitration operations.

HYDROFLUORINATION WATER SCRUBBER

Hydrofluorination, as described above, involves contacting uranium dioxide with vaporized hydrofluoric acid at an elevated temperature. Unreacted hydrofluoric acid fumes are passed through a water scrubber which absorbs much of the hydrofluoric acid. Vent gases pass to the second scrubber noted above. Since the scrubber liquor over the hydrofluorination unit absorbs acid, the liquor is circulated until a specified concentration of hydrofluoric acid is attained. Then the solution is drawn off and sold for industrial use. Therefore, the hydrofluorination water scrubber discharges no wastewater to treatment.

HYDROFLUORINATION ALKALINE SCRUBBER

Hydrofluorination involves contacting uranium dioxide with hydrofluoric acid to produce uranium tetrafluoride. The offgases from this operation, after passing through a water scrubber, are scrubbed by a circulating KOH solution which neutralizes and scrubs the acidic fumes. The scrubber liquor is completely recycled until scrubber efficiency diminishes; then the liquor is batch discharged to combined treatment. Treatment consists of neutralization and sedimentation, followed by direct discharge to a surface water.

MAGNESIUM REDUCTION AND CASTING FLOOR WASH WATER

Water is used to wash floors and equipment in the magnesium reduction and casting area. One plant uses a floor washing machine and a second plant washes floors manually. The plant which uses a floor washing machine discharges the wash water to a batch chemical precipitation and sedimentation treatment system prior to discharge to surface waters. At the other plant reporting this stream, the reduction and casting wash water is combined with machining wastewater and treated using chemical precipitation and rotary precoat filtration prior to discharge to surface waters.

LAUNDRY WASTEWATER

Two plants reported the use of water to wash the clothing of plant personnel working in production areas. One facility treats the resulting wastewater in a batch chemical precipitation and sedimentation treatment system prior to discharge to surface waters. The other facility reporting this practice, discharges laundry wastewater to surface waters after treatment consisting of chemical precipitation and sedimentation.

CONTROL AND TREATMENT OPTIONS

The Agency examined two control and treatment technology options that are applicable to the secondary uranium subcategory. The options selected for evaluation represent applicable end-of-pipe treatment technologies.

Examination of the waste streams in this subcategory shows that no further in-process flow reduction is achievable. Recycle of laundry wastewater has been considered in Section X, BAT. On the VI), options including activated carbon adsorption were not considered.

OPTION A

Option A for the secondary uranium subcategory requires control and treatment technologies to reduce the discharge of pollutant mass.

The Option A treatment scheme consists of chemical precipitation and sedimentation technology. Specifically, lime or some other alkaline compound is used to precipitate dissolved metals as metal hydroxides. The metal hydroxides and suspended solids settle out and the sludge is collected. Vacuum filtration is used to dewater sludge. OPTION C

Option C for the secondary uranium subcategory consists of all control and treatment requirements of Option A (chemical precipitation and sedimentation) plus multimedia filtration technology added at the end of the Option A treatment scheme. Multimedia filtration is used to remove suspended solids, including precipitates of metals, beyond the concentration attainable by gravity sedimentation. The filter suggested is of the gravity, mixed-media type, although other forms of filters, such as rapid sand filters or pressure filters would perform satisfactorily. The addition of filters also provides consistent removal during periods of time in which there are rapid increases in flows or loadings of pollutants to the treatment system.

SECTION VIII

COSTS, ENERGY, AND NONWATER QUALITY ASPECTS

This section presents a summary of compliance costs for the secondary uranium subcategory and a description of the treatment options and subcategory-specific assumptions used to develop these estimates. Together with the estimated pollutant removal performance presented in Section X of this supplement, these cost estimates provide a basis for evaluating each regulatory option. These cost estimates are also used in determining the probable economic impact of regulation on the subcategory at different pollutant discharge levels. In addition, this section addresses nonwater quality environmental impacts of wastewater treatment and control alternatives, including air pollution, solid wastes, and energy requirements, which are specific to the secondary uranium subcategory.

TREATMENT OPTIONS FOR EXISTING SOURCES

As discussed in Section VII, two treatment options have been developed and considered in promulgating limitations and standards for the secondary uranium subcategory. These options are summarized below and schematically presented in Figures X-1 and X-2 (pages 4777 and 4778).

OPTION A

The Option A treatment scheme consists of chemical precipitation and sedimentation technology.

OPTION C

Option C for the secondary uranium subcategory consists of all control and treatment requirements of Option A (chemical precipitation and sedimentation) plus multimedia filtration technology added at the end of the Option A treatment scheme.

COST METHODOLOGY

A detailed discussion of the methodology used to develop the compliance costs is presented in Section VIII of Vol. I. Plantby-plant compliance costs for the nonferrous metals manufacturing category have been revised as necessary following proposal. These revisions calculate incremental costs, above treatment already in place, necessary to comply with the promulgated effluent limitations and standards and are presented in the administrative record supporting this regulation. A comparison of the costs developed for proposal and the revised costs for the final regulation are presented in Table VIII-1 (page 4746) for the direct discharger in this subcategory.

Each of the general assumptions used to develop compliance costs

is presented in Section VIII of the General Development Document. Subcategory-specific assumptions were used in developing compliance costs for the secondary uranium subcategory are listed below.

(1) Costs for purchasing a floor washing machine were included in the compliance cost estimates for plant 1175 because the Agency believes that this piece of equipment is necessary for this facility to achieve the flow allowance for magnesium reduction and casting floor wash water.

NONWATER QUALITY ASPECTS

A general discussion of the nonwater quality aspects of the control and treatment options considered for the nonferrous metals category is contained in Section VIII of the General Development Document. Nonwater quality impacts specific to the secondary uranium subcategory, including energy requirements, solid waste and air pollution are discussed below.

ENERGY REQUIREMENTS

The methodology used for determining the energy requirements for the various options is discussed in Section VIII of the General Energy requirements for Option A are Development Document. estimated at 52,000 kwh/yr, and for Option C the estimated requirement is 62,000 kwh/yr. Option C energy requirements increased over those for Option A because filtration is being added as an end-of-pipe treatment technology. Since recycle and scrubber liquor is already practiced in this energy requirement savings resulting from flow reuse of subcategory, energy requirement reduction measures are not reflected in this analysis. Both Option A and Option C energy requirements represent less than 1 percent of the energy usage in the secondary uranium industry. is therefore concluded that the energy requirements of the It treatment options considered will have no significant impact on total plant energy consumption.

SOLID WASTE

Sludge generated in the secondary uranium subcategory is due to the precipitation of metals as hydroxides and carbonates using lime. Sludges associated with the secondary uranium subcategory will necessarily contain quantities of toxic metal pollutants.

Wastes generated by secondary metal industries can be regulated as hazardous. However, the Agency examined the solid wastes that would be generated at secondary nonferrous metals manufacturing plants by the suggested treatment technologies and believes they not hazardous wastes under the Agency's regulations are implementing Section 3001 of the Resource Conservation and Recovery Act. None of the secondary uranium subcategory wastes are listed specifically as hazardous, nor are they likely to exhibit a characteristic of hazardous waste. This judgment is

made based on the recommended technology of lime precipitation and filtration. By the addition of a small excess of lime during treatment, similar sludges, specifically toxic metal bearing sludges, generated by other industries such as the iron and steel industry passed the Extraction Procedure (EP) toxicity test. See 40 CFR §261.24. Thus, the Agency believes that the wastewater sludges will similarly not be EP toxic if the recommended technology is applied.

Although it is the Agency's view that solid wastes generated as a result of these guidelines are not expected to be hazardous, generators of these wastes must test the waste to determine if the wastes meet any of the characteristics of hazardous waste (see 40 CFR 262.11).

If these wastes should be identified or are listed as hazardous, they will come within the scope of RCRA's "cradle to grave" hazardous waste management program, requiring regulation from, the point of generation to point of final disposition. EPA's generator standards would require generators to hazardous nonferrous metals manufacturing wastes to meet containerization, labeling, recordkeeping, and reporting requirements; if plants dispose of hazardous wastes off-site, they would have to prepare a manifest which would track the movement of the wastes from the generator's premises to a permitted off-site treatment, storage, or disposal facility. See 40 CFR 262.20 45 FR 33142 (May 19, 1980), as amended at 45 FR 86973 (December 31, 1980). The transporter regulations require transporters of hazardous wastes to comply with the manifest system to assure that the wastes are delivered to a permitted facility. See 40 CFR 263.20 45 FR 33151 (May 19, 1980), as amended at 45 FR 86973 (December 31, 1980). Finally, RCRA regulations establish standards for hazardous waste treatment, storage, and disposal facilities allowed to receive such wastes. See 40 CFR Part 464 46 FR 2802 (January 12, 1981), 47 FR 32274 (July 26, 1982).

Even if these wastes are not identified as hazardous, they still must be disposed of in compliance with the Subtitle D open dumping standards, implementing 4004 of RCRA. See 44 FR 53438 (September 13, 1979). It is estimated that the secondary uranium subcategory will generate 285 metric tons of sludge per year when implementing the BPT treatment technology. The Agency has calculated as part of the costs for wastewater treatment the cost of hauling and disposing of these wastes.

AIR POLLUTION

There is no reason to believe that any substantial air pollution problems will result from implementation of chemical precipitation, sedimentation, and multimedia filtration. The model technologies transfer pollutants to solid waste and are not likely to transfer pollutants to air.

TABLE VIII-1

COST OF COMPLIANCE FOR THE SECONDARY URANIUM SUBCATEGORY DIRECT DISCHARGERS

(March 1982 Dollars)

Option	Proposal <u>Capital</u> Cost	Cost <u>Annual</u> Cost	Promulgati <u>Capital</u> Cost	ion Cost <u>Annual</u> Cost
А	28,600	73,600	54,800	90,400
С	54,300	86,500	88,000	106,700

SECTION IX

BEST PRACTICABLE CONTROL TECHNOLOGY CURRENTLY AVAILABLE

This section defines the effluent characteristics attainable through the application of best practicable control technology currently available (BPT). BPT reflects the existing performance by plants of various sizes, ages, and manufacturing processes within the secondary uranium subcategory, as well as the established performance of the model BPT systems. Particular consideration is given to the treatment already in place at plants within the data base.

The factors considered in identifying BPT include the total cost of applying the technology in relation to the effluent reduction benefits from such application, the age of equipment and facilities involved, the manufacturing processes used, nonwater quality environmental impacts (including energy requirements), and other factors the Administrator considers appropriate. In general, the BPT level represents the average of the existing performances of plants of various ages, sizes, processes, or common characteristics. Where existing performance other is uniformly inadequate, BPT may be transferred from a different subcategory or category. Limitations based on transfer technology are supported by a rationale concluding that of the technology is, indeed, transferable, and a reasonable prediction that it will be capable of achieving the prescribed effluent limits BPT focuses on end-of-pipe treatment rather than process changes or internal controls, except where such practices are common industry practice.

TECHNICAL APPROACH TO BPT

The Agency studied the nonferrous metals category to identify the processes used, the wastewaters generated, and the treatment processes installed. Information was collected from industry using data collection portfolios, and specific plants were sampled and the wastewaters analyzed. In making technical assessments of data, reviewing manufacturing processes, and assessing wastewater treatment technology options, both indirect and direct dischargers have been considered as a single group. An examination of plants and processes did not indicate any process differences based on the type of discharge, whether it be direct or indirect.

As explained in Section IV, the secondary uranium subcategory has been subdivided into seven potential wastewater sources. Since the water use, discharge rates, and pollutant characteristics of each of these wastewaters is potentially unique, effluent limitations will be developed for each of the seven subdivisions.

For each of the subdivisions, a specific approach was followed

The first the development of BPT mass limitations. for requirement to calculate these limitations is to account for production and flow variability from plant to plant. Therefore, unit of production or production normalizing parameter (PNP) a determined for each wastewater stream which could then be was related to the flow from the process to determine a production normalized flow. Selection of the PNP for each process element is discussed in Section IV. Each plant within the subcategory was then analyzed to determine which subdivisions were present, the specific flow rates generated for each subdivision, and the specific production normalized flows for each subdivision. This analysis is discussed in detail in Section V. Nonprocess wastewaters such as rainfall runoff and noncontact cooling water are not considered in the analysis.

Production normalized flows for each subdivision were then analyzed to determine the flow to be used as part of the basis for BPT mass limitations. The selected flow (sometimes referred to as the BPT regulatory flow or BPT discharge rate) reflects the water use controls which are common practices within the category. The BPT regulatory flow is based on the average of all applicable data. Plants with normalized flows above the average may have to implement some method of flow reduction to achieve the BPT limitations.

The second requirement to calculate mass limitations is the set of concentrations that are achievable by application of the BPT level of treatment technology. Section VII discusses the various control and treatment technologies which are currently in place for each wastewater source. In most cases, the current control and treatment technologies consist of chemical precipitation and sedimentation (lime and settle technology) and a combination of reuse and recycle to reduce flow.

Using these regulatory flows and the achievable concentrations, the next step is to calculate mass loadings for each wastewater source or subdivision. This calculation was made on a stream-bystream basis, primarily because plants in this subcategory may perform one or more of the operations in various combinations. The mass loadings (milligrams of pollutant per kilogram of production - mg/kg) were calculated based on the BPT regulatory flow (l/kkg) and the concentration achievable by the BPT level of treatment technology (mg/l) for each pollutant parameter to be limited under BPT. These mass loadings are published in the Federal Register and in 40 CFR Part 421 as the effluent limitations.

The mass loadings which are allowed under BPT for each plant will be the sum of the individual mass loadings for the various wastewater sources which are found at particular plants. Accordingly, all the wastewater generated within a plant may be combined for treatment in a single or common treatment system, but the effluent limitations for these combined wastewaters are based on the various wastewater sources which actually contribute to the combined flow. This method accounts for the variety of

combinations of wastewater sources and production processes which may be found at secondary uranium plants.

The Agency usually establishes wastewater limitations in terms of mass rather than concentration. This approach prevents the use of dilution as a treatment method (except for controlling pH). The production normalized wastewater flow (l/kkg) is a link between the production operations and the effluent limitations. The pollutant discharge attributable to each operation can be calculated from the normalized flow and effluent concentration achievable by the treatment technology and summed to derive an appropriate limitation for each plant.

INDUSTRY COST AND POLLUTANT REMOVAL ESTIMATES

In balancing costs in relation to pollutant removal estimates, EPA considers the volume and nature of existing discharges, the volume and nature of discharges expected after application of BPT, the general environmental effects of the pollutants, and the cost and economic impacts of the required pollution control level. The Act does not require or permit consideration of water quality problems attributable to particular point sources or industries, or water quality improvements in particular water quality bodies. Accordingly, water quality considerations were not the basis for selecting the proposed or promulgated BPT.

The methodology for calculating pollutant removal estimates and plant compliance costs is discussed in Section X. The pollutant removal estimates have been revised since proposal. Table X-1 (page 4769) shows the pollutant removal estimates for each treatment option for direct dischargers. Compliance costs for direct dischargers are presented in Table X-2 (page 4770).

BPT OPTION SELECTION - PROPOSAL

EPA proposed BPT requirements for the secondary uranium subcategory. The technology basis for the BPT limitations is lime precipitation and sedimentation technology to remove metals and solids from combined wastewaters and to control pH. BPT also includes ammonia steam stripping. These technologies are already in-place at the one discharger in the subcategory. The pollutants specifically proposed for regulation at BPT are chromium, copper, nickel, ammonia, fluoride, uranium, TSS, and pH.

Implementation of the proposed BPT limitations will remove annually an estimated 1,280 kg of toxic metals, 12,000 kg of ammonia and 1,763 kg of TSS. While the one discharging plant has the equipment in-place to comply with BPT, we do not believe that the plant is currently achieving the proposed BPT limitations. We project capital and annual costs of \$28,600 and \$37,644 (1982 dollars) respectively for modifications to technology presented in-place at the discharging facility to achieve proposed BPT regulations.

BPT OPTION SELECTION - PROMULGATION

The promulgated technology basis for the BPT limitations is Option A. chemical precipitation and sedimentation technology to remove metals and solids from combined wastewaters and to control pH. The promulgated technology basis for BPT limitations differs from the proposed technology basis. These technologies are demonstrated and economically achievable since they are already in place at several discharging plants throughout the nonferrous metals manufacturing category.

Implementation of the promulgated BPT limitations will remove annually an estimated 100 kg of toxic metals, 4,283 kg of nonconventional pollutants, and 651 kg of TSS. While the one discharging plant has most of the equipment in-place to comply plant --The Agency project. (1982 dollars) with BPT, EPA does not believe that the plant is currently achieving the promulgated BPT limitations. capital and annual costs of \$54,800 and \$90,400 (1982 respectively for modifications to technology presently in-place discharging facility to achieve promulgated the BPT at regulations. The end-of-pipe treatment configuration for Option A is presented in Figure IX-1 (page 4761).

WASTEWATER DISCHARGE RATES

A BPT discharge rate is calculated for each subdivision based on the average of the flows of the existing plants, as determined from analysis of data collection portfolios. The discharge rate is used with the achievable treatment concentrations to determine BPT effluent limitations. Since the discharge rate may be different for each wastewater source, separate production normalized discharge rates for each of the seven wastewater sources are discussed below and summarized in Table IX-1. The discharge rates are normalized on a production basis by relating amount of wastewater generated to the mass the of the intermediate product which is produced by the process associated with the waste stream in question. These production normalizing parameters, or PNPs, are also listed in Table IX-1.

Section V of this document further describes the discharge flow rates and presents the water use and discharge flow rates for each plant by subdivision in Tables V-1 through V-7 (pages 4788 - 4790).

REFINERY SUMP FILTRATE

The BPT wastewater discharge rate at proposal for refinery sump filtrate (formerly, refinery filtrate at proposal) was 34,800 l/kkg (8,340 gal/ton) of uranium trioxide produced. This rate was allocated for those plants that acid leach scrap uranium materials to recover the uranium. The BPT discharge rate was based on flow information provided by one plant. Post-proposal comments and information showed that the proposed flow was not accurate and that the production normalizing parameter was not appropriate. EPA studied the data and decided to promulgate a corrected (larger) flow allowance and, in addition, change the PNP for this subdivision.

The BPT wastewater discharge rate at promulgation for refinery sump filtrate is 73,340 l/kkg (17,580 gal/ton) of uranium processed in the refinery. This rate is allocated only for those plants that employ acid leaching and dissolution operations to recover uranium from secondary sources. The flow from the one plant in this subcategory having this operation was used to determine the promulgated BPT regulatory flow.

SLAG LEACH RESLURRY

The BPT wastewater discharge rate at proposal for slag leach reslurry (formerly, slag leach slurry at proposal) was 3,800 1/kkg (910 gal/ton) of uranium trioxide produced. This rate was allocated only for those plants which leach magnesium fluoride slag, recycled from the magnesium reduction operation, to recover the residual uranium in the slag. Post-proposal comments from the industry indicated a difficulty with the PNP chosen for this If a plant operates leaching on a campaign basis, subdivision. it must have a way to determine its regulatory allowances without waiting for the next process step to be completed. EPA difficulty and chose a acknowledged this new PNP for promulgation. However, the wastewater flow (1/yr) used to calculate the regulatory flow (1/kkg) was not altered.

The promulgated BPT wastewater discharge rate for slag leach reslurry is 4,566 l/kkg (l,094 gal/ton) of uranium processed in the refinery. This rate is allocated only for those plants which recover uranium by leaching magnesium fluoride slag. One plant in this subcategory has this operation, and the promulgated BPT discharge rate is based on the water use at this plant. Table V-2 (page 4688) presents water use and discharge rates for slag leach reslurry.

DIGESTION WET AIR POLLUTION CONTROL

The BPT wastewater discharge rate at proposal for digestion wet pollution control was 30 l/kkg (7.2 gal/ton) of uranium air trioxide produced based on partial recycle of scrubber liquor. This rate was allocated only for those plants that incorporate a water scrubber on the acid leaching and dissolution operation. Post-proposal comments from the industry clarified the Agency's understanding of the digestion scrubber's operation. The comments indicated that scrubber blowdown is reused in the leaching and dissolution operation because it contains nitric Thus no discharge occurs from this scrubber, and EPA acid. decided to change the discharge rate for this subdivision at promulgation.

The BPT wastewater discharge rate at promulgation for digestion wet air pollution control is zero. This rate is allocated to those plants that operate a water scrubber to control fumes from acid leaching and dissolution. The promulgated BPT discharge rate is based upon water use information supplied by a plant in this subcategory having a digestion operation scrubber. Existing practice at this plant is such that 100 percent of the scrubber liquor is recycled or reused. Therefore, it is appropriate that the BPT regulatory flow should be zero.

SOLVENT EXTRACTION RAFFINATE FILTRATE

The BPT wastewater discharge rate used at proposal for solvent extraction raffinate filtrate (formerly, solvent extraction raffinate at proposal) was 5,300 l/kkg (1,270 gal/ton) of uranium This rate was allocated for those plants trioxide produced. which purify the acid-dissolved uranium compound by extracting the uranium compound into an organic solvent, leaving behind all impurities that were leached along with the uranium. Postproposal comments from the industry indicated a difficulty with the PNP chosen for this subdivision. If a plant operates solvent extraction on a campaign basis, it must have a method to determine its regulatory allowance without waiting for the next process step to be completed. EPA acknowledged this difficulty and chose a new PNP for promulgation. However, the wastewater flow (1/yr) used to calculate the regulatory flow (1/kkg) was not altered.

The BPT wastewater discharge rate at promulgation for action raffinate filtrate is 6,369 l/kkg (1,526 gal/ton) of uranium those plants using solvent extraction procedures to purify uranium compounds dissolved in solution as a result of acid leaching and dissolution. The BPT discharge rate at promulgation is based on the water use at one plant in the subcategory having this operation. Production normalized water use and discharge rates are presented in Table V-4 (page 4689).

EVAPORATION AND DENITRATION WET AIR POLLUTION CONTROL

Zero discharge is used at promulgation for evaporation and denitration wet air pollution control (formerly, evaporation and calcination wet air pollution control at proposal). This requirement is applicable to those plants that use evaporators and calcinators to respectively concentrate an intermediate uranium compound and then calcine it to produce uranium trioxide. intermediate BPT discharge rate is promulgated as zero because the one The discharging plant in this subcategory that uses these operations recycles all scrubber liquor to the digestion operation. This production normalized discharge rate is also presented in Table V-5 (page 4689).

HYDROFLUORINATION WATER SCRUBBER

Zero discharge is used at promulgation for hydrofluorination water scrubber (formerly, hydrofluorination wet air pollution control at proposal) wastewater. This requirement is applicable only to those plants which use a water scrubber to control acid fumes from the hydrofluorination unit. The BPT discharge rate is promulgated as zero because the one plant in this subcategory

that operates such a scrubber recycles the scrubber liquor to absorb the hydrofluoric acid fumes until a desired concentration of hydrofluoric acid is attained. Then the scrubber solution is drawn off and sold for industrial use. Since this recycle technology is demonstrated within this subcategory, the BPT discharge rate reflects that capability. Table V-6 (page 4689) also presents the water use and discharge rate for the hydrofluorination water scrubber system.

HYDROFLUORINATION ALKALINE SCRUBBER

The BPT wastewater discharge rate at proposal and promulgation hydrofluorination alkaline scrubber (formerly, for hydrogen reduction and hydrofluorination KOH wet air pollution control at proposal) wastewater is 20 l/kkg (4.8 gal/ton) of uranium tetrafluoride produced based on partial recycle. This rate is allocated only for those plants that hydrofluorinate uranium dioxide to produce uranium tetrafluoride, and scrub the gases from this operation with a KOH scrubber. The BPT discharge rate reflects the demonstrated performance of this scrubber operating at a high rate of recycle. Table V-7 (page 4690) also presents the water use and discharge rates for this wastewater stream.

MAGNESIUM REDUCTION AND CASTING FLOOR WASH WATER

The BPT wastewater discharge rate used at promulgation for magnesium reduction and casting floor wash water is 30.1 l/kkg of uranium produced by magnesium reduction. This rate is allocated to all plants which produce uranium metal by magnesium reduction of uranium tetrafluoride. This rate is based on the production normalized flow reported by plant 1066. This plant uses a floor washing machine, thereby achieving a lower wastewater discharge rate than facilities which do not use a floor washing machine. This rate is used as the basis for the BPT flow allowance because this water use and discharge rate is demonstrated and is achievable through the use of a floor washing machine.

There was no wastewater discharge allowance at proposal for this subdivision because the Agency did not have sufficient data at the time to quantify production normalized water use and discharge rates. These rates, which are based on data collected by the Agency since proposal, are presented in Table V-8 (page 4690).

LAUNDRY WASTEWATER

The BPT wastewater discharge rate at promulgation for laundry wastewater is 192 l/kkg of uranium produced by magnesium reduction. This rate is allocated to all plants which produce uranium metal by magnesium reduction of uranium tetrafluoride. This rate is based on one facility for which the Agency obtained sufficient data to calculate a production normalized flow.

There was no wastewater discharge allowance at proposal for this subdivision because the Agency did not have sufficient data at

the time to calculate production normalized water use and discharge rates. These rates, which are based on information collected by the Agency since proposal, are presented in Table V-9 (page 4690).

REGULATED POLLUTANT PARAMETERS

The raw wastewater concentrations from individual operations and the subcategory as a whole were examined to select certain pollutant parameters for limitation. This examination and evaluation was presented in Section VI. A total of seven pollutants or pollutant parameters are selected for limitation under BPT and are listed below:

119. chromium 120. copper 124. nickel fluoride TSS pH

EFFLUENT LIMITATIONS

The treatable concentrations achievable by application of the promulgated BPT are discussed in Section VII of Vol. I and summarized there in Table VII-21 (page 248). These treatable concentrations (both one day maximum and monthly average values) are multiplied by the BPT normalized discharge flows summarized in Table IX-1 (page 4755) to calculate the mass of pollutants allowed to be discharged per mass of product. The results of these calculations in milligrams of pollutant per kilogram of product represent the BPT effluent limitations and are presented in Table IX-2 (page 4756) for each individual waste stream.

Table IX-1

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BPT WASTEWATER DISCHARGE RATES FOR THE SECONDARY URANIUM SUBCATEGORY

Wastewater Stream		malized ge Rate gal/ton	Production Normalizing Parameter
Refinery sump filtrate	73,340	17,580	Uranium processed in the refinery
Slag leach reslurry	4,566	1,094	Uranium processed in the refinery
Digestion wet air pollution control	0	0	Uranium processed in the refinery
Solvent extraction raffinate filtrate	6,369	1,526	Uranium processed in the refinery
Evaporation and denitration wet air pollution control	0	0	Uranium trioxide produced
Hydrofluorination water scrubber	. 0	0	Uranium tetrafluoride produced
Hydrofluorination alkaline scrubber	20	4.8	Uranium tetrafluoride produced
Magnesium reduction and cast- ing floor wash	30.1	7.2	Uranium produced by mag- nesium reduction
Laundry wastewater	192	46.1	Uranium produced by mag- nesium reduction

SECONDARY URANIUM SUBCATEGORY

SECT - IX

TABLE IX-2

BPT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(a) <u>Refinery</u> <u>Sump</u> <u>Filtrate</u> BPT

Pollutant	or	Maximum	for	Maximum		
pollutant		any one	day	monthly	average	

mg/kg (lb/million lb	os) of uranium pro	cessed in the refinery
Antimony	210.500	93.880
Arsenic	153.300	68.210
Cadmium	24.940	11.000
*Chromium	32.270	13.200
*Copper	139.300	73.340
	30.800	14.670
Lead	140.800	93.140
*Nickel	90.210	40.340
Selenium	30.070	12.470
Silver	107.100	44.740
Zinc	2,567.000	1,459.000
*Fluoride		346.900
Uranium	476.700	1,430.000
*TSS	3,007.000	
*pH Within the range	e of 7.5 to 10.0 a	C ALL LIMES

(b) Slag Leach Reslurry BPT

Pollutant pollutant			imum for one day		imum thly	for average	
mg/kg	(lb/milli	on lbs)	of urani	um proce	essed	in the	refinery
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium *TSS			13.100 9.543 1.552 2.009 8.675 1.918 8.767 5.616 1.872 6.666 159.800 29.680 187.200		• .	5.844 4.246 0.685 0.822 4.566 0.913 5.799 2.511 0.776 2.785 90.860 21.600 89.040	
*pH Wi	thin the	range of	7.5 to	10.0 at	all t	imes	

Table IX-2 (Continued)

BPT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(c) Solvent Extraction Raffinate Filtrate BPT

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average
mg/kg (lb/million lbs	s) of uranium pr	ocessed in solvent extraction
Antimony	18.280	8.152
Arsenic	13.310	5.923
Cadmium	2.165	0.955
*Chromium	2.802	1.146
*Copper	12.100	6.369
Lead	2.675	1.274
*Nickel	12.230	8.089
Selenium	7.834	3.503
Silver	2.611	1.083
Zinc	9.299	3.885
*Fluoride	222.900	126.700
Uranium	41.400	30.130
*TSS	261.100	124.200
*pH Within the ran	nge of 7.5 to 10	.0 at all times
		······································
(d) Digestion Operat:	ion Wet Air Poll	ution Control BPT
Pollutant or	Maximum for	Maximum for
pollutant property		
porrulance property	any one day	monenty average
ma/ka (lb/million	lbs) of uranium	processed in the refinery
	100) of ardina	
Antimony	0.000	0.000
Arsenic	0.000	0.000
Cadmium	0.000	0.000
*Chromium	0.000	0.000
*Copper	0.000	0.000
Lead	0.000	0.000
*Nickel	0.000	0.000
Selenium	0.000	0.000
Silver	0.000	0.000
······································		

 Zinc
 0.000
 0.000

 *Fluoride
 0.000
 0.000

 Uranium
 0.000
 0.000

 *TSS
 0.000
 0.000

 *pH
 Within the range of 7.5 to 10.0 at all times

*Regulated Pollutant

0.000

0.000

0.000

Table IX-2 (Continued)

BPT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(e) Evaporation and Denitration Wet Air Pollution Control BPT

Pollutant or	Maximum for	Maximum	average
pollutant property	any one day	monthly	
mg/kg (lb/milli	on lbs) of uran	ium trioxi	de reduced

mg/	Ng (±2/-		,				
Antimony			0.0	00		0.000	
Arsenic			0.0	00		0.000	
Cadmium			0.0	00		0.000	•
*Chromium			0.0			0.000	
			0.0			0.000	
*Copper		i.	0.0			0.000	
Lead *Nickel			0.0			0.000	
			0.0			0.000	
Selenium			0.0			0.000	
Silver			0.0			0.000	
Zinc			0.0	<i></i>		0.000	
*Fluoride			0.0			0.000	
Uranium						0.000	
*TSS			0.0				
*pH Wit	thin the	range of	: 7.5 t	0 TO.0	at all	times	

(f) Hydrofluorination KOH Scrubber BPT

pollutant property an	ximum for y one day	Maximum for monthly average
mg/kg (1b/million 1bs) of uranium	tetrafluoride produced
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium *TSS *pH Within the range C	$\begin{array}{c} 0.057\\ 0.042\\ 0.007\\ 0.009\\ 0.038\\ 0.008\\ 0.025\\ 0.008\\ 0.025\\ 0.008\\ 0.029\\ 0.700\\ 0.130\\ 0.820\\ 0.75\ to\ 10. \end{array}$	0.026 0.019 0.003 0.004 0.020 0.004 0.025 0.011 0.003 0.012 0.398 0.095 0.390 0 at all times

TABLE IX-2 (Continued)

BPT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(g) Hydrofluorination Water Scrubber BPT

Pollutant	or	Maximum for	Maximum for
pollutant	property	any one day	monthly average
mg/kg	(lb/million	lbs) of uranium	tetrafluoride produced
Antimony		0.000	0.000
Arsenic		0.000	0.000
Cadmium		0.000	0.000
*Chromium		0.000	0.000
*Copper		0.000	0.000
Lead	¥.	0.000	0.000
*Nickel		0.000	0.000
Selenium		0.000	0.000
Silver		0.000	0.000
Zinc		0.000	0.000
*Fluoride		0.000	0.000
Uranium		0.000	0.000
*TSS		0.000	0.000
*pH Wit	thin the rang	ge of 7.5 to 10.0	

(h) <u>Reduction</u> and <u>Casting</u> <u>Floor</u> <u>Wash</u> BPT

Pollutant or	Maximum for Maximu	um for
pollutant property	any one day monthl	y average
mg/kg (lb/million lbs)	of uranium produced by	magnesium reduction
Antimony	0.086	0.039
Arsenic	0.063	0.028
Cadmium	0.010	0.005
*Chromium	0.013	0.005
*Copper	0.057	0.030
Lead	0.013	0.006
*Nickel	0.058	0.038
Selenium	0.037	0.017
Silver	0.012	0.005
Zinc	0.044	0.018
*Fluoride	1.054	0.599
Uranium	0.196	0.142
*TSS	1.234	0.587
	e of 7.5 to 10.0 at all	

TABLE IX-2 (Continued)

BPT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(i) Laundry Washwater BPT

Pollutant or	Maximum for	Maximum for	
pollutant property	any one day	monthly average	

mg/kg (lb/million lbs) of uranium produced by magnesium reduction

Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium *TSS	0.551 0.401 0.065 0.085 0.365 0.081 0.369 0.236 0.079 0.280 6.720 1.248 7.872	0.246 0.179 0.029 0.035 0.192 0.038 0.244 0.106 0.033 0.117 3.821 0.908 3.744 at all times
*pH Within	the range of 7.5 to 10.0	at all times

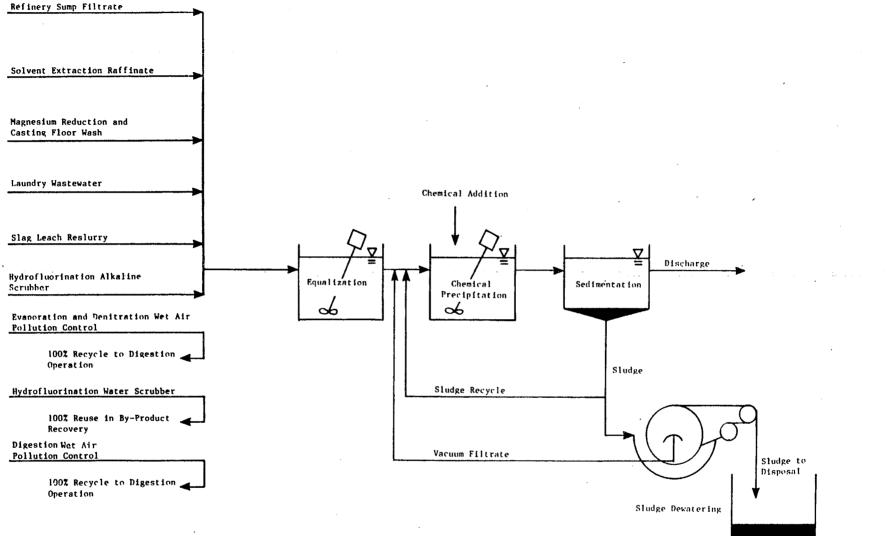


Figure IX-1

BPT TREATMENT SCHEME FOR THE SECONDARY URANIUM SUBCATEGORY

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SECT - IX

SECONDARY URANIUM SUBCATEGORY

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SECTION X

BEST AVAILABLE TECHNOLOGY ECONOMICALLY ACHIEVABLE

These effluent limitations are based on the best control and treatment technology used by a specific point source within the industrial category or subcategory, or by another industry from which it is transferable. Emphasis is placed on additional treatment techniques applied at the end of the treatment systems currently used, as well as reduction of the amount of water used and discharged, process control, and treatment technology optimization.

The factors considered in assessing best available technology economically achievable (BAT) include the age of equipment and facilities involved, the process used, process changes, nonwater quality environmental impacts (including energy requirements), and the costs of application of such technology. BAT represents the best available technology economically achievable at plants of various ages, sizes, processes, or other characteristics. BAT may be transferred from a different subcategory or category. BAT may include feasible process changes or internal controls, even when not in common industry practice.

The statutory assessment of BAT considers costs, but does not require a balancing of costs against pollutant removals. However, in assessing the proposed and promulgated BAT, the Agency has given substantial weight to the economic achievability of the technology.

TECHNICAL APPROACH TO BAT

The Agency reviewed a wide range of technology options and evaluated the available possibilities to ensure that the most effective and beneficial technologies were used as the basis of BAT. To accomplish this, the Agency elected to examine two technology options which could be applied to the secondary uranium subcategory as alternatives for the basis of BAT effluent limitations.

For the development of BAT effluent limitations, mass loadings were calculated for each wastewater source or subdivision in the subcategory using the same technical approach as described in Section IX for BPT limitations development. The differences in the mass loadings for BPT and BAT are due to increased treatment effectiveness achievable with the more sophisticated BAT treatment technology.

The treatment technologies considered for BAT are summarized below:

Option A (Figure X-1, page 4777):

o Chemical precipitation and sedimentation

Option C (Figure X-2, page 4778):

o Chemical precipitation and sedimentation o Multimedia filtration

The two options examined for BAT are discussed in greater detail below. The first option considered (Option A) is the same as the BPT treatment and control technology which was presented in the previous section. The second option represents substantial progress toward the reduction of pollutant discharges above and beyond the progress achievable by BPT.

OPTION A

Option A for the secondary uranium subcategory is equivalent to the control and treatment technologies which were analyzed for BPT in Section IX (see Figures IX-1 or X-1, pages 4761 or 4777). The BPT end-of-pipe treatment scheme includes chemical precipitation and sedimentation. The discharge rates for Option A are equal to the discharge rates allocated to each stream as a BPT discharge flow.

OPTION C

Option C for the secondary uranium subcategory consists of all control and treatment requirements of Option A (chemical precipitation and sedimentation plus multimedia filtration technology added at the end of the Option A treatment scheme (see Figure X-2). Multimedia filtration is used to remove suspended solids, including precipitates of toxic metals, beyond the concentrations attainable by gravity sedimentation. The filter suggested is of the gravity, mixed media type, although other forms of filters, such as rapid sand filters or pressure filters, would perform satisfactorily.

INDUSTRY COST AND POLLUTANT REMOVAL ESTIMATES

As one means of evaluating each technology option, EPA developed estimates of the pollutant removals and the compliance costs associated with each option. The methodologies are described below.

POLLUTANT REMOVAL ESTIMATES

A complete description of the methodology used to calculate the estimated pollutant removal, or benefit, achieved by the application of the various treatment options is presented in Section X of the General Development Document. The pollutant removal estimates have been revised from proposal based on industry comments and new information; however, the methodology for calculating pollutant removals was not changed. The data used for estimating removals are the same as those used to revise compliance costs.

Sampling data collected during the field sampling program were used to characterize the major waste streams considered for At each sampled facility, the sampling data were regulation. production normalized for each unit operation (i.e., mass of pollutant generated per mass of product manufactured). This value, referred to as the raw waste, was used to estimate the mass of pollutants generated within the secondary uranium subcategory. The pollutant removal estimates were calculated for each plant by first estimating the total mass of each pollutant in the untreated wastewater. This was calculated by first multiplying the raw waste values by the corresponding production value for that stream and then summing these values for each pollutant for every stream generated by the plant.

Next, the volume of wastewater discharged after the application of each treatment option was estimated for each operation at each plant by comparing the actual discharge to the regulatory flow. smaller of the two values was selected and summed with the The other plant flows. The mass of pollutant discharged was then estimated by multiplying the achievable concentration values attainable with the option (mg/l) by the estimated volume of process wastewater discharged by the subcategory. The mass of pollutant removed is the difference between the estimated mass of pollutant generated by each plant in the subcategory and the mass of pollutant discharged after application of the treatment The pollutant removal estimates for direct dischargers option. in the secondary uranium subcategory are presented in Table X-1 (page 4769).

COMPLIANCE COSTS

In estimating subcategory-wide compliance costs, the first step was to develop a cost estimation model, relating the total costs with installation and operation of associated wastewater treatment technologies to plant process wastewater discharge. EPA applied the model to each plant. The plant's investment and operating costs are determined by what treatment it has in place and by its individual process wastewater discharge flow. As discussed above, this flow is either the actual or the BAT regulatory flow, whichever is lesser. The final step was to annualized the capital costs, and to sum the annualized capital costs, and the operating and maintenance costs for each plant, the cost of compliance for the subcategory. yielding Α comparison of the costs developed for proposal and the revised costs for promulgation is presented in Table X-2 (page 4778) for direct discharges in the secondary uranium subcategory. These costs were used in assessing economic achievability.

BAT OPTION SELECTION - PROPOSAL

EPA selected Option C for the proposed BAT which included pretreatment with ammonia steam stripping for selected waste

streams, followed by chemical precipitation, sedimentation, and multimedia filtration. The estimated capital cost of proposed BAT was \$54,300, and the annual cost was \$86,500 (1982 dollars). Implementation of the proposed BAT technology was estimated to remove 1,304 kilograms of priority pollutants and 1,951 kilograms of TSS annually.

BAT OPTION SELECTION - PROMULGATION

EPA is promulgating BAT limitations for the secondary uranium subcategory based on chemical precipitation, sedimentation, and multimedia filtration. The end-of-pipe technology basis for BAT limitations being promulgated is the same as that for the proposed limitations. In addition, the treatment performance concentrations, upon which the mass limitations are based, are equal to values used to calculate the proposed mass limitations. Ammonia steam stripping is no longer part of the model technology for BAT. Data collected after proposal indicated that ammonia is no longer used in secondary uranium processing. The treatment configuration for Option C is presented in Figure X-2 (page 4778).

EPA is promulgating multimedia filtration as part of the BAT technology because this technology is demonstrated by 25 plants in the nonferrous metals manufacturing category, and results in additional removal of priority metals. In addition, filtration adds reliability to the treatment system by making it less susceptible to operator error and to sudden changes in raw wastewater flow and concentrations.

Implementation of the control and treatment technologies of Option C will remove annually an estimated 126 kilograms of priority metal pollutants and 4,350 kilograms of nonconventional pollutants, which is 26 kilograms of priority metal pollutants over the estimated BPT removal. The estimated capital cost for achieving promulgated BAT is \$88,000 and the estimated annual cost is \$106.700 (1982 dollars).

WASTEWATER DISCHARGE RATES

A BAT discharge rate was calculated for each subdivision based upon the flows of the existing plants, as determined from analysis of the data collection portfolios. The discharge rate is used with the achievable treatment concentrations to determine BAT effluent limitations. Since the discharge rate may be different for each wastewater source, separate production normalized discharge rates for each of the seven wastewater determined and are summarized in Table X-3 sources were (page 4771). The discharge rates are normalized on a production basis by relating the amount of wastewater generated to the mass of the intermediate product which is produced by the process associated with the waste stream in question. These production normalizing parameters, or PNPs, are also listed in Table X-3.

The BAT wastewater discharge rates are the same as the BPT

discharge rates with the exception of the laundry wastewater flow. The BAT flow rate for laundry wastewater is 96 l/kkg of uranium produced by magnesium reduction. based on 50 percent recycle of the BPT flow using a holding tank and recycle equipment.

REGULATED POLLUTANT PARAMETERS

The raw wastewater concentrations from individual operations and the subcategory as a whole were examined to select certain pollutants and pollutant parameters for limitation. This examination and evaluation was presented in Section VI. The Agency, however, has chosen not to regulate all eight toxic pollutants selected in this analysis.

The high cost associated with analysis for toxic metal pollutants has prompted EPA to develop an alternative method for regulating and monitoring toxic pollutant discharges from the nonferrous metals manufacturing category. Rather than developing specific effluent mass limitations and standards for each of the toxic metals found in treatable concentrations in the raw wastewater from a given subcategory, the Agency is promulgating effluent mass limitations only for those pollutants generated in the greatest quantities as shown by the pollutant removal estimate analysis. The pollutants selected for specific limitation are listed below:

119. chromium
120. copper
124. nickel
fluoride

By establishing limitations and standards for certain priority metal pollutants, dischargers will attain the same degree of control over toxic metal pollutants as they would have been required to achieve had all the toxic metal pollutants been directly limited.

This approach is technically justified since the treatable concentrations used for chemical precipitation and sedimentation technology are based on optimized treatment for concomitant multiple metals removal. Thus, even though metals have somewhat different theoretical solubilities, they will be removed at very precipitation the same rate in a chemical nearlv and sedimentation treatment system operated for multiple metals removal. Filtration as part of the technology basis is likewise justified because this technology removes metals nonpreferentially.

The priority metal pollutants selected for specific limitation in the secondary uranium subcategory to control the discharges of toxic metal pollutants are chromium, copper, and nickel. The following toxic metal pollutants are excluded from limitation on the basis that they are effectively controlled by the limitations developed for chromium, copper, and nickel: 114. antimony
115. arsenic
118. cadmium
122. lead
125. selenium
126. silver
128. zinc

EFFLUENT LIMITATIONS

The concentrations achievable by application of BAT are discussed in Section VII of Vol. I and are summarized there in Table VII-21 (page 248). The achievable concentrations both one day maximum and monthly average values are multiplied by the BAT normalized discharge flows summarized in Table X-3 to calculate the mass of pollutants allowed to be discharged per mass of product. The results of these calculations in milligrams of pollutant per kilogram of product represent the promulgated BAT effluent limitations and are presented in Table X-4 (page 4772) for each waste stream.

Table X-1

POLLUTANT REMOVAL ESTIMATES FOR DIRECT DISCHARGERS

Pollutant	Total Raw Discharge (kg/yr)	Option A Discharge (kg/yr)	Option A Removed (kg/yr)	Option C Discharge (kg/yr)	Option C Removed (kg/yr)
Antimony	0.00	0.00	0.00	0.00	0.00
Arsenic	24.23	15.18	9.04	10.08	14.14
Cadmium	0.47	0.47	0.00	0.47	0.00
Chromium (Total)	7.41	2.50	4.91	2.07	5.33
Copper	34.91	17.26	17.65	11.57	23.34
Cyanide (Total)	0.00	0.00	0.00	0.00	0.00
Lead /	10.80	3.57	7.23	2.37	8.43
Mercury	2.91	1.78	1.13	1.06	1.84
Nickel	13.57	13.57	0.00	6.52	7.04
Selenium	19.40	8.93	10.47	5.93	13.47
Silver	0.00	0.00	0.00	0.00	0.00
Thallium	0.00	0.00	0.00	0.00	0.00
Zinc	59.15	9,82	49.32	6.82	52.32
TOTAL PRIORITY POLLUTANTS	172.89	73.11	99.78	46.94	125.95
Aluminum	1,273.29	66.68	1,206.60	44.20	1,229.08
Ammonia	0.00	0.00	0.00	0.00	0.00
Cobalt	0.00	0.00	0.00	0.00	0.00
Uranium	314.87	119.07	195.80	78.32	236,55
Iron	2,700.78	12.20	2,688.57	8.30	2,692.47
Manganese	196.38	4.76	191.62	4.15	192.22
Phosphorus	0.00	0.00	0.00	0.00	0.00
TOTAL NONCONVENTIONALS	4,485.33	202.72	4,282.61	135.00	4,350.33
TSS	1,008.55	357.22	651.33	77 1/	000 / /
Oil and Grease	0.00	0.00		77.14	931.41
-	0.00	0.00	0.00	0.00	0.00
TOTAL CONVENTIONALS	1,088.55	357.22	651.33	77.14	931.41
TOTAL POLLUTANTS	5,666.79	633.05	5,033.73	259.08	5,407.71

SECONDARY URANIUM SUBCATEGORY SECT -

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TABLE X-2

COST OF COMPLIANCE FOR THE SECONDARY URANIUM SUBCATEGORY DIRECT DISCHARGERS

(March 1982 Dollars)

<u>Option</u>	Proposal Capital Cost	Cost Annual Cost	Promulgati <u>Capital Cost</u>	on Cost <u>Annual</u> Cost
A	28,600	73,600	54,800	90,400
С	54,300	86,500	88,000	106,700

Table X-3

BAT WASTEWATER DISCHARGE RATES FOR THE SECONDARY URANIUM SUBCATEGORY

		malized ge Rate	Production Normalizing
Wastewater Stream	1/kkg	gal/ton	Parameter
Refinery sump filtrate	73,340	17,580	Uranium processed in the refinery
Slag leach reslurry	4,566	1,094	Uranium processed in the refinery
Digestion wet air pollution control	0	0	Uranium processed in the refinery
Solvent extraction raffinate filtrate	6,369	1,526	Uranium processed in the refinery
Evaporation and denitration wet air pollution control	0	0	Uranium trioxide produced
Hydrofluorination water scrubber	0	0	Uranium tetrafluoride produced
Hydrofluorination alkaline scrubber	20	4.8	Uranium tetrafluoride produced
Magnesium reduction and cast- ing floor wash	30.1	7.2	Uranium produced by magnesium reduction
Laundry wastewater	96	23	Uranium produced by magnesium reduction

SECONDARY URANIUM SUBCATEGORY SECT -×

TABLE X-4

BAT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(a) <u>Refinery</u> <u>Sump</u> <u>Filtrate</u> BAT

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average

mg/kg	(lb/million	lbs)	of	uranium	processed	in	the	refinery
mg/kg Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride	l l			41.500 01.900 14.670 27.140 93.880 20.540 40.340 50.140 21.270 74.810 57.000		6: 4! 1: 4: 2: 3:	3.07(5.47(5.86) 1.00(4.74(9.534 7.14(7.14(3.80) 0.80(9.00(D D D D D 4 D D D D D D D
Uranium			33	14.600		228	8.80	D

(b) <u>Slag Leach Reslurry</u> BAT

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average
mg/kg (lb/million	lbs) of uranium	processed in the refinery
Antimony	8.812	3.927
Arsenic	6.347	2.831
Cadmium	.913	.365
*Chromium	1.689	.685
*Copper	5.844	2.785
Lead	1.278	.594
*Nickel	2.511	1.689
Selenium	3.744	1.689
Silver	1.324	.548
Zinc	4.657	1.918
*Fluoride	159.800	90.860
Uranium	19.590	14.250

TABLE X-4 (Continued)

BAT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(c) Solvent Extraction Raffinate Filtrate BAT

Pollutant	or	Maximum	for	Maximum	for
pollutant	property	any one	day	monthly	average

mg/kg (lb/million lbs) of uranium processed in solvent extraction

7 m h i m a m r	12 200	E 477
Antimony	12.290	5.477
Arsenic	8.853	3.949
Cadmium	1.274	0.510
*Chromium	2.357	0.955
*Copper	8.152	3.885
Lead	1.783	0.828
*Nickel	3.503	2.357
Selenium	5.223	2.357
Silver	1.847	0.764
Zinc	6.496	2.675
*Fluoride	222.900	126.700
Uranium	27.320	19.870

(d) Digestion Operation Wet Air Pollution Control BAT

Pollutant or pollutant property	Maximum for any one day	Maximum for monthly average
mg/kg (lb/million	lbs) of uranium	processed in the refinery
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium	$\begin{array}{c} 0.000\\ 0.$	$\begin{array}{c} 0.000\\ 0.$

TABLE- X-4 (Continued)

BAT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(e) Evaporation and Denitration Wet Air Pollution Control BAT

Pollutant or	<u>.</u>	Maximum for		
pollutant p	coperty	any one day	monthly	average
mg/kg	g (lb/millio	n lbs) of u	iranium triox	ide produced
Antimony		0.000)	0.000
Arsenic		0.000)	0.000
Cadmium		0.000)	0.000
*Chromium		0.000)	0.000
*Copper		0.00)	0.000
Lead		0.00)	0.000
*Nickel		0.00)	0.000
Selenium		0.00)	0.000
Silver		0.00)	0.000
Zinc		0.00)	0.000
*Fluoride		0.00)	0.000
Uranium		0.00)	0.000

(f) Hydrofluorination KOH Scrubber BAT

Pollutant pollutant		Maximum for any one day	Maximum for monthly average
mg/kg	(lb/million	lbs) of uranium	tetrafluoride produced
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium		0.039 0.028 0.004 0.007 0.026 0.006 0.011 0.016 0.006 0.020 0.700 0.086	0.017 0.012 0.002 0.003 0.012 0.003 0.007 0.007 0.007 0.002 0.008 0.398 0.062

Table X-4 (Continued)

BAT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(g) Hydrofluorination Water Scrubber BAT

Pollutant	or	Maximum for	Maximum for
pollutant	property	any one day	monthly average

mg/kg (lb/million lbs) of uranium tetrafluoride produced

Antimony	0.000	0.000
Arsenic	0.000	0.000
Cadmium	0.000	0.000
*Chromium	0.000	0.000
*Copper	0.000	0.000
Lead	0.000	0.000
*Nickel	0.000	0.000
Selenium	0.000	0.000
Silver	0.000	0.000
Zinc	0.000	0.000
*Fluoride	0.000	0.000
Uranium	0.000	0.000
,		

(h) <u>Reduction</u> and <u>Casting</u> Floor Wash BAT

Pollutant or pollutant property		ximum for nthly average
<pre>mg/kg (lb/million lbs)</pre>	of uranium produce	d by magnesium reduction
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride	0.058 0.042 0.006 0.011 0.039 0.008 0.017 0.025 0.009 0.031 1.054	0.026 0.019 0.002 0.005 0.018 0.004 0.011 0.011 0.001 0.004 0.013 0.599
Uranium	0.129	0.094

TABLE-4 (Continued)

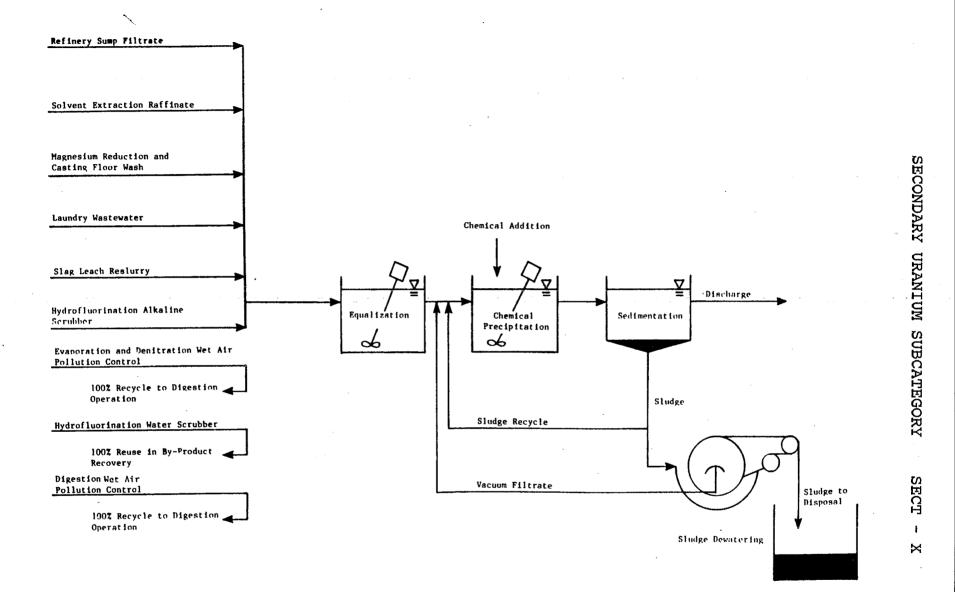
BAT MASS LIMITATIONS FOR THE SECONDARY URANIUM SUBCATEGORY

(i) Laundry Washwater BAT

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average

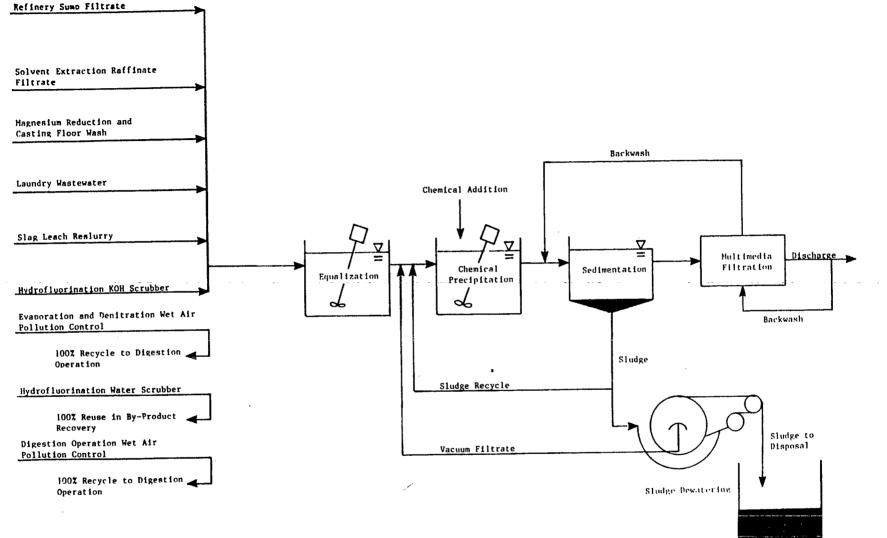
mg/kg (lb/million lbs) of uranium produced by magnesium reduction

Antimony	0.185	0.083
Arsenic	0.133	0.060
Cadmium	0.019	0.008
*Chromium	0.036	0.014
*Copper	0.123	0.059
Lead	0.027	0.013
*Nickel	0.053	0.036
Selenium	0.079	0.036
Silver	0.028	0.012
Zinc	0.098	0.040
*Fluoride	3.360	1.910
Uranium	0.412	0.300



BAT TREATMENT SCHEME FOR OPTION A

4777





BAT TREATMENT SCHEME FOR OPTION C

4778

SECT

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SECONDARY URANIUM SUBCATEGORY

SECTION XI

NEW SOURCE PERFORMANCE STANDARDS

This section describes the technologies for treatment of wastewater from new sources and presents mass discharge standards for regulatory pollutants for NSPS in the secondary uranium subcategory, based on the selected treatment technology. New plants have the opportunity to design the best and most efficient production processes and wastewater treatment technologies without facing the added costs and restrictions encountered in retrofitting an existing plant. Therefore, EPA has considered the best demonstrated process changes, in-plant controls, and end-of-pipe treatment technologies which reduce pollution to the maximum extent feasible.

TECHNICAL APPROACH TO NSPS

New source performance standards are equivalent to the best available technology (BAT) selected for currently existing secondary uranium plants. This result is a consequence of careful review by the Agency of a wide range of technical options for new source treatment systems which is discussed in Section XI of the General Development Document. Additionally, there was found to indicate that the wastewater flows nothing and characteristics of new plants would not be similar to those from existing plants, since the processes used by new sources are not expected to differ from those used at existing sources. Consequently, BAT production normalized discharge rates, which are based on the best existing practices of the subcategory, can also be applied to new sources. These rates are presented in Table X-3 (page 4771).

Treatment technologies considered for the NSPS options are identical to the treatment technologies considered for the BAT options. These options are:

OPTION A

o Chemical precipitation and sedimentation

OPTION C

o Chemical precipitation and sedimentation

o Multimedia filtration

NSPS OPTION SELECTION - PROPOSAL

EPA proposed that the best available demonstrated technology for the secondary uranium subcategory be equivalent to Option C (chemical precipitation, sedimentation, and multimedia filtration with ammonia steam stripping for selected streams).

The wastewater flow rates proposed for NSPS were the same as those proposed for BAT. Flow reduction measures for NSPS and BAT were not considered feasible because EPA believed that no new demonstrated technologies existed within the subcategory that improved on present water use practices. Therefore, EPA concluded that NSPS flow rates should be equal to those of BPT and BAT.

NSPS OPTION SELECTION - PROMULGATION

EPA is promulgating best available demonstrated technology for the secondary uranium subcategory equivalent to Option C (chemical precipitation, sedimentation, and multimedia filtration). Filtration technology is demonstrated in 25 plants in the nonferrous metals manufacturing category.

The promulgated wastewater flow rates for NSPS are the same as the promulgated BAT flow rates. Flow reduction measures for NSPS are not considered feasible. EPA does not believe that new plants could achieve any additional flow reduction beyond the 90 to 100 percent scrubber effluent recycle presently practiced in the industry.

REGULATED POLLUTANT PARAMETERS

The Agency has no reason to believe that the pollutants that will be found in treatable concentrations in processes within new sources will be any different than with existing sources. Accordingly, pollutants and pollutant parameters selected for limitation under NSPS, in accordance with the rationale of Sections VI and X, are identical to those selected for BAT. The conventional pollutant parameters TSS and pH are also selected for limitation.

NEW SOURCE PERFORMANCE STANDARDS

The NSPS discharge flows for each wastewater source are the same as the discharge rates for BAT and are shown in Table XI-1 (page 4781). The mass of pollutant allowed to be discharged per mass of product is based on the product of the appropriate achievable concentration (mg/l) and the production normalized wastewater discharge flows (l/kkg). The results of these calculations are the production-based new source performance standards. These standards are presented in Table XI-2 (page 4782).

Table XI-1

NSPS WASTEWATER DISCHARGE RATES FOR THE SECONDARY URANIUM SUBCATEGORY

NSPS Normalized <u>Discharge Rate</u> Production Normalizing					
<u>Wastewater Stream</u>	1/kkg	gal/ton	Parameter		
Refinery sump filtrate	73,340	17,580	Uranium processed in the refinery		
Slag leach reslurry	4,566	1,094	Uranium processed in the refinery		
Digestion wet air pollution control	0	0	Uranium processed in the refinery		
Solvent extraction raffinate filtrate	6,369	1,526	Uranium processed in the refinery		
Evaporation and denitration wet air pollution control	0	0	Uranium trioxide produced		
Hydrofluorination water scrubber	0	0	Uranium tetrafluoride produced		
Hydrofluorination alkaline scrubber	20	4.8	Uranium tetrafluoride produced		
Magnesium reduction and cast- ing floor wash	30.1	7.2	Uranium produced by magnesium reduction		
Laundry wastewater	96	23	Uranium produced by magnesium reduction		

SECONDARY URANIUM SUBCATEGORY SECT -

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TABLE XI-2

NSPS FOR THE SECONDARY URANIUM SUBCATEGORY

(a) <u>Refinery</u> <u>Sump</u> <u>Filtrate</u> NSPS

			the second s	
Pollutant or	Maximum	for	Maximum	for
pollutant property	any one	day	monthly	average

mg/kg (lb/million lbs) of uranium processed in the refinery

Antimony 141.50 Arsenic 101.90 Cadmium 14.67 *Chromium 27.14 *Copper 93.88 *Lead 20.54 *Nickel 40.34 Selenium 60.14 Silver 21.27 Zinc 74.81 *Fluoride 2,567.00 Uranium 314.60	$\begin{array}{cccccccccccccccccccccccccccccccccccc$
*TSS 1,100.00 *pH Within the range of 7.5 to	

(b) <u>Slag Leach</u> <u>Reslurry</u> NSPS

Pollutant or pollutant pro		aximum ny one		Maximum monthly		
mg/kg (lb/	million lbs) of u	ranium	processed	in the	refinery
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium *TSS *pH Withir	n the range	6 1 5 1 2 3 1 4 159 19 68	.812 .347 .913 .689 .844 .278 .511 .744 .324 .657 .800 .590 .490 to 10	.0 at all	3.927 2.831 .365 .685 2.785 .594 1.689 1.689 1.689 .548 1.918 90.860 14.250 54.790 times	

TABLE XI-2 (Continued)

NSPS FOR THE SECONDARY URANIUM SUBCATEGORY

(c) Solvent Extraction Raffinate Filtrate NSPS

Pollutant	or	Maximum	for	Maximum	for
pollutant	property	any one	day	monthly	average

mg/kg (lb/million lbs) of uranium processed in solvent extraction

Antimony	12.290	5.477
Arsenic	8.853	3.949
Cadmium	1.274	0.510
*Chromium	2.357	0.955
*Copper	8.152	3.885
Lead	1.783	0.828
*Nickel	3.503	2.357
Selenium	5.223	2.357
Silver	1.847	0.764
Zinc	6.496	2.675
,*Fluoride	222.900	126.700
Uranium	27.320	19.870
*TSS	95.540	76.430
*pH Within t	he range of 7.5 to 10.0 at	all times

(d) Digestion Operation Wet Air Pollution Control NSPS

Pollutant pollutant	-	Maximu any or		Maximum monthly	for average
mg/kg	(lb/million	lbs) of	uranium	processed	in the refinery
Antimony			0.000		0.000
Arsenic Cadmium			0.000		0.000
*Chromium *Copper			0.000		0.000
Lead *Nickel			0.000		0.000
Selenium			0.000		0.000
Silver Zinc			0.000		0.000 0.000
*Fluoride Uranium			0.000		0.000 0.000
*TSS		_	0.000		0.000
*pH Wit	chin the ra	nge of 7.	5 to 10	.0 at all t	imes

TABLE XI-2 (Continued)

NSPS FOR THE SECONDARY URANIUM SUBCATEGORY

(e) Evaporation and Denitration Wet Air Pollution Control NSPS

Pollutant or pollutant prop	erty a	Maximum for any one day	Maximum monthly	average	
mg/kg (lb/millior	n lbs) of un	anium triox:	ide produ	ced
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium *TSS *pH Within	the range	0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000	10.0 at all	0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 0.000 times	

(f) Hydrofluorination KOH Scrubber NSPS

mg/kg (lb/million lbs) of uranium tetrafluoride produced Antimony 0.039 0.017 Arsenic 0.028 0.012 Cadmium 0.004 0.002 *Chromium 0.007 0.003 *Copper 0.026 0.012 Lead 0.006 0.007 Nickel 0.011 0.007 Selenium 0.016 0.007 Silver 0.006 0.002 Zinc 0.020 0.008 *Fluoride 0.700 0.398 Uranium 0.086 0.062 *TSS 0.300 0.240 *pH Within the range of 7.5 to 10.0 at all times	Pollutant pollutant	property	Maximum for any one day	Maximum for monthly average
Antimony 0.028 0.012 Arsenic 0.004 0.002 Cadmium 0.007 0.003 *Chromium 0.026 0.012 *Copper 0.026 0.012 Lead 0.006 0.003 *Nickel 0.011 0.007 Selenium 0.006 0.002 Silver 0.006 0.002 Zinc 0.020 0.398 Vranium 0.086 0.062 Uranium 0.300 0.240	mg/kg	(lb/million	lbs) of uranium	tetrafluoride produced
• ·	Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium		0.028 0.004 0.007 0.026 0.006 0.011 0.016 0.006 0.020 0.700 0.086 0.300	0.012 0.002 0.003 0.012 0.003 0.007 0.007 0.007 0.002 0.008 0.398 0.062 0.240

TABLE XI-2 (Continued)

NSPS FOR THE SECONDARY URANIUM SUBCATEGORY

(g) Hydrofluorination Water Scrubber NSPS

Pollutant	or	Maximum for	Maximum for
pollutant	property	any one day	monthly average
ma/ka	(lb/million	lbg) of uranium	tetrafluoride produced
ilig/ kg	(10/ 11111011	105) Of diamidm	cectariationiae produced
Antimony		0.000	0.000
Arsenic		0.000	0.000
Cadmium		0.000	0.000
*Chromium		0.000	0.000
*Copper		0.000	0.000
Lead		0.000	0.000
*Nickel		0.000	0.000
Selenium		0.000	0.000
Silver		0.000	0.000
Zinc		0.000	0.000
*Fluoride		0.000	0.000
Uranium		0.000	0.000
*TSS		0.000	0.000
*pH Wi	thin the rang	ge of 7.5 to 10.0	at all times

(h) <u>Reduction</u> and <u>Casting Floor</u> Wash NSPS

Polluta	ant or	Maximum for	Maximum	for	
polluta	ant property	any one day	monthly	average	
-			-	5	
mg/kg ((lb/million lbs)	of uranium	produced by	magnesium	reduction
				·	
Antimo	-	0.058		0.026	
Arseni	ic	0.042		0.019	
Cadmiu	um	0.006		0.002	
*Chromi	ium	0.011		0.005	
*Copper		0.039		0.018	
Lead		0.008		0.004	
*Nicke]	1	0.017		0.011	
Seleni	ium	0.025		0.011	
Silver		0.009		0.004	
Zinc	_	0.031		0.013	
*Fluori	ide	1.054		0.599	
Uraniı		0.129		0.094	
*TSS	a ***	0.452		0.361	
*pH	Within the rang			+ +	
Pu	Michilli che lang	e or /.J to	ivev at all	CTICO	

TABLE XI- (Continued)

NSPS FOR THE SECONDARY URANIUM SUBCATEGORY

(i) Laundry Washwater NSPS

Pollutant or		imum for
pollutant property	any one day mon	thly average
mg/kg (lb/million lbs)	of uranium produced	by magnesium reduction
Antimony	0.185	0.083
Arsenic	0.133	0.060
Cadmium	0.019	0.008
*Chromium	0.036	0.014
*Copper	0.123	0.059
Lead	0.027	0.013
*Nickel	0.053	0.036
Selenium	0.079	0.036
Silver	0.028	0.012
Zinc	0.098	0.040
*Fluoride	3.360	1.910
Uranium	0.412	0.300
*TSS	1.440	1.152
	e of 7.5 to 10.0 at	all times

SECTION XII

PRETREATMENT STANDARDS

This section describes the control and treatment technologies for pretreatment of process wastewaters from new sources in the secondary uranium subcategory. PSES are designed to prevent the discharge of pollutants which pass through, interfere with, or are otherwise incompatible with the operation of publicly owned (POTW). treatment works The Clean Water Act requires pretreatment for pollutants, such as toxic metals, that limit POTW sludge management alternatives. New indirect discharge facilities, like new direct discharge facilities, have the opportunity to incorporate the best available demonstrated technologies, including process changes, in-plant controls, and end-of-pipe treatment technologies, and to use plant site selection to ensure adequate treatment system installation. Pretreatment standards are to be technology based, analogous to the best available or best demonstrated technology for removal of toxic pollutants.

Pretreatment standards for existing sources (PSES) will not be promulgated for the secondary uranium subcategory because there are no existing indirect dischargers in this subcategory. However, pretreatment standards for new sources (PSNS) will be promulgated. Pretreatment standards for regulated pollutants are presented based on the selected control and treatment technology.

TECHNICAL APPROACH TO PRETREATMENT

Before proposing and promulgating pretreatment standards, the Agency examines whether the pollutants discharged by the industry pass through the POTW or interfere with the POTW operation or its sludge disposal practices. In determining whether chosen pollutants pass through a well-operated POTW achieving secondary treatment, the Agency compares the percentage of a pollutant removed by POTW with the percentage removed by direct dischargers applying the best available technology economically achievable. A pollutant is deemed to pass through the POTW when the average percentage removed nationwide by well-operated POTW meeting secondary treatment requirements, is less than the percentage removed by direct dischargers complying with BAT effluent limitations guidelines for that pollutant.

This definition of pass through satisfies the two competing objectives set by Congress that standards for indirect dischargers be equivalent to standards for direct dischargers while at the same time the treatment capability and performance of the POTW be recognized and taken into account in regulating the discharge of pollutants from indirect dischargers.

The Agency compares percentage removal rather than the mass or concentration of pollutants discharged because the latter would

not take into account the mass of pollutants discharged to the POTW from non-industrial sources or the dilution of the pollutants in the POTW effluent to lower concentrations due to the addition of large amounts of non-industrial wastewater.

PRETREATMENT STANDARDS FOR NEW SOURCES

Options for pretreatment of wastewaters from new sources are based on increasing the effectiveness of end-of-pipe treatment technologies. All in-plant changes and applicable end-of-pipe treatment processes have been discussed previously in Sections X and XI. The options for PSNS, therefore, are the same as the BAT options discussed in Section X. A description of each option is presented in Section X.

Treatment technologies considered for the PSNS options are:

OPTION A

o Chemical precipitation and sedimentation

OPTION C

- o Chemical precipitation and sedimentation
- o Multimedia filtration

PSNS OPTION SELECTION

Option C (chemical precipitation, sedimentation, and multimedia filtration) has been selected as the regulatory approach for pretreatment standards for new sources. The basis of this selection is in accordance with the rationale for selection of the BAT option in Section X. Option C prevents pass-through and equivalent to BAT treatment for direct dischargers. is In addition, Option C achieves effective removal of priority pollutants by incorporating filtration which is demonstrated by 25 plants throughout the nonferrous metals manufacturing category.

The wastewater discharge rates for promulgated PSNS are identical to the promulgated BAT discharge rates for each waste stream. The PSNS discharge rates are shown in Table XII-1 (page 4790). No additional flow reduction measures for PSNS are feasible. EPA does not believe that new plants should achieve flow reduction beyond the 90 to 100 percent scrubber effluent recycle presently practiced in the industry.

REGULATED POLLUTANT PARAMETERS

Pollutants selected for limitation, in accordance with the rationale of Sections VI and X, are identical to those selected for limitation for BAT. It is necessary to promulgate PSNS to prevent the pass-through of chromium, copper, nickel, and fluoride, which are the limited pollutants.

PRETREATMENT STANDARDS FOR NEW SOURCES

Pretreatment standards for new sources are based on the achievable concentrations from the selected treatment technology, (Option C), and the discharge rates determined in Section X for BAT. A mass of pollutant per mass of product (mg/kg) allocation is given for each subdivision within the subcategory. This pollutant allocation is based on the product of the treatable concentration from the promulgated treatment (mg/l) and the production normalized wastewater discharge rate (l/kkg). The achievable treatment concentrations for BAT are identical to those for PSNS. PSNS are presented in Table XII-2 (page 4791).

Table XII-1

PSNS WASTEWATER DISCHARGE RATES FOR THE SECONDARY URANIUM SUBCATEGORY

Wastewater Stream	PSNS Nor Dischar 1/kkg	rmalized <u>ge Rate</u> gal/ton	Production Normalizing Parameter
Refinery sump filtrate	73,340	17,580	Uranium processed in the refinery
Slag leach reslurry	4,566	1,094	Uranium processed in the refinery
Digestion wet air pollution control	0	0	Uranium processed in the refinery
Solvent extraction raffinate filtrate	6,369	1,526	Uranium processed in the refinery
Evaporation and denitration wet air pollution control	0	0	Uranium trioxide produced
Hydrofluorination water scrubber	0	0	Uranium tetrafluoride produced
Hydrofluorination alkaline scrubber	20	4.8	Uranium tetrafluoride produced
Magnesium reduction and cast- ing floor wash	30.1	7.2	Uranium produced by magnesium reduction
Laundry wastewater	96	23	Uranium produced by magnesium reduction

SECONDARY URANIUM SUBCATEGORY

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SECT - XII

TABLE XII-2

PSNS FOR THE SECONDARY URANIUM SUBCATEGORY

(a) <u>Refinery</u> <u>Sump</u> <u>Filtrate</u> PSNS

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average
	lbs) of uranium	processed in the refinery
Antimony	141.500	63.070
Arsenic	101.900	45.470
Cadmium	14.670	5.867
*Chromium	27.140	11.000
*Copper	93.880	44.740
Lead	20.540	9.534
*Nickel	40.340	27.140
Selenium	60.140	27.140
Silver	21.270	8.801
Zinc	74.810	30.800
*Fluoride	2,567.000	1,459.000
Uranium	314.600	228.800

(b) <u>Slag Leach Reslurry</u> PSNS

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average
mg/kg (lb/million	lbs) of uranium	processed in the refinery
Antimony	8.812	3.927
Arsenic	6.347	2.831
Cadmium	0.913	0.365
*Chromium	1.689	0.685
*Copper	5.844	2.785
Lead	1.278	0.594
*Nickel	2.511	1.689
Selenium	3.744	1.689
Silver	1.324	0.548
Zinc	4.657	1.918
*Fluoride	159.800	90.860
Uranium	19.590	14.250

TABLE XII-2 (Continued)

PSNS FOR THE SECONDARY URANIUM SUBCATEGORY

(c) Solvent Extraction Raffinate Filtrate PSNS

Pollutant or pollutant property	Maximum for any one day	Maximum monthly		
mg/kg (lb/million lbs)	of uranium p	rocessed in	solvent	extraction
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium	12.290 8.853 1.274 2.357 8.152 1.783 3.503 5.223 1.847 6.496 222.900 27.320		5.477 3.949 0.510 0.955 3.885 0.828 2.357 2.357 0.764 2.675 126.700 19.870	

(d) Digestion Operation Wet Air Pollution Control PSNS

Pollutant or pollutant property	Maximum for any one day	Maximum for monthly average
mg/kg (lb/million	lbs) of uranium	processed in the refinery
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium	0.000 0.000	$\begin{array}{c} 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\end{array}$

TABLE XII-2 (Continued)

PSNS FOR THE SECONDARY URANIUM SUBCATEGORY

(e) Evaporation and Denitration Wet Air Pollution Control PSNS

Pollutant or	Maximum for	Maximum for
pollutant property	any one day	monthly average
ng/kg (lb/million lbs)	of uranium tri	oxide produced
Antimony	0.000	0.000
Arsenic	0.000	0.000
Cadmium	0.000	0.000
*Chromium	0.000	0.000
*Copper	0.000	0.000
Lead	0.000	0.000
*Nickel	0.000	0.000
Selenium	0.000	0.000
Silver	0.000	0.000
Zinc	0.000	0.000
*Fluoride	0.000	0.000
Uranium	0.000	0.000

(f) Hydrofluorination KOH Scrubber PSNS

Pollutant or	Maximum for	Maximum for
··· •· ··· ·· ·· ·· ···		
pollutant property	any one day	monthly average
(he (lb /million lbe)	ali	afluggide produced
mg/kg (lb/million lbs)	of uranium tetr	arruoride produced
Antimony	0.039	0.017
Arsenic	0.028	0.012
Cadmium	0.004	0.002
*Chromium	0.007	0.003
*Copper	0.026	0.012
Lead	0.006	0.003
*Nickel	0.011	0.007
Selenium	0.016	0.007
Silver	0.006	0.002
Zinc	0.020	0.008
*Fluoride	0.700	0.398
Uranium	0.086	0.062

Table XII-2 (Continued)

PSNS FOR THE SECONDARY URANIUM SUBCATEGORY

(g) Hydrofluorination Water Scrubber PSNS

Pollutant or pollutant property	Maximum for any one day	Maximum for monthly average
<pre>mg/kg (lb/million lbs)</pre>	of uranium tetr	afluoride produced
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium	$\begin{array}{c} 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\end{array}$	$\begin{array}{c} 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\\ 0.000\end{array}$

(h) Reduction and Casting Floor Wash PSNS

Pollutant or pollutant property		mum for hly average
mg/kg (lb/million lbs)	of uranium produced	by magnesium reduction
Antimony Arsenic Cadmium *Chromium *Copper Lead *Nickel Selenium Silver Zinc *Fluoride Uranium	0.058 0.042 0.006 0.011 0.039 0.008 0.017 0.025 0.009 0.031 1.054 0.129	0.026 0.019 0.002 0.005 0.018 0.004 0.011 0.011 0.011 0.004 0.013 0.599 0.094

Table XII-2 (Continued)

PSNS FOR THE SECONDARY URANIUM SUBCATEGORY

(i) Laundry Washwater PSNS

Pollutant or	Maximum for	Maximum for	
pollutant property	any one day	monthly average	
<pre>mg/kg (lb/million lbs)</pre>	of uranium produ	aced by magnesium reduction	L
Antimony	0.185	0.083	
Arsenic	0.133	0.060	
Cadmium	0.019	0.008	
*Chromium	0.036	0.014	
*Copper	0.123	0.059	
Lead	0.027	0.013	
*Nickel	0.053	0.036	
Selenium	0.079	0.036	
Silver	0.028	0.012	
Zinc	0.098	0.040	
*Fluoride	3.360	1.910	
Uranium	0.412	0.300	

SECTION XIII

BEST CONVENTIONAL POLLUTANT CONTROL TECHNOLOGY

EPA is not promulgating best conventional pollutant control technology (BCT) for the secondary uranium subcategory at this time.

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