

November 20, 2023

Mr. Josh Peters On-Scene Coordinator U.S. Environmental Protection Agency, Region 5 Superfund and Emergency Management Division 2565 Plymouth Road Ann Arbor, MI 48105 We are in the process of ensuring this document is accessible to all audiences. If you need assistance accessing this document, or any material on the EPA East Palestine, Ohio emergency response web pages, please contact the Region 5 Public Information Officer on-call at: R5 EastPalestine@epa.gov

Subject: Data Validation Report E Palestine Site - ER EPA Contract No.: 68HE0519D0005 Task Order/Task Order Line Item No.: 68HE0520F0032/0001EB201 Document Tracking No. 2199

Dear Mr. Peters:

Tetra Tech, Inc. (Tetra Tech) is submitting this data validation report for 79 air samples (including 5 field duplicate samples, 5 field blank samples, and 3 media blanks) collected at the E Palestine site. The samples were collected on July 13, July 14, and July 15, 2023, and were analyzed for acrylates by Eurofins Analytics, LLC. The final laboratory data package was received on July 18, 2023.

Analytical data were evaluated in general accordance with the Tetra Tech Quality Assurance Project Plan, East Palestine Train Derailment Site, East Palestine, Columbiana County, Ohio, Revision 3 (April 2023), the Tetra Tech Quality Assurance Project Plan, Superfund Technical Assessment and Response Team (START V), EPA Region 5, Revision 4 (August 2022), and the National Functional Guidelines (NFG) for Organic Superfund Methods Data Review (November 2020).

No qualification or rejection of results was required for these data packages. The results may be used as reported by the laboratory.

If you have any questions regarding this data validation report, please contact me via the project manager.

Sincerely,

Patrick JDigitally signed by
Patrick J GarrityGarrityDate: 2023.11.20
08:16:59 -05'00'

Environmental Chemist

Enclosure

 cc: Karl Schultz, Tetra Tech Program Manager Dustin Grams, Tetra Tech Project Manager Mayra Arroyo Ortiz, Tetra Tech Project Document Control Coordinator TO-TOLIN File

ATTACHMENT

DATA VALIDATION REPORT EUROFINS ANALYTICS, LLC REPORT NOS. B198-117, B198-118, B198-119, B198-120

Site Name	E Palestine Site - ER	TO/TOLIN No.	68HE0520F0032/0001EB201
Document Tracking No.	2199a	TO/TOLIN NO.	08HE0520F0032/0001EB201
Laboratory Report No.	B198-117	Laboratory	Eurofins Analytics, LLC – Ashland, VA
Analyses	n-Butyl acrylate by NIOSH Method 1450M		
Samples and Matrix	32 air samples including 2 field blanks, 2 me	edia blanks, and 3 field	l duplicate pairs
Collection Date(s)	07/13/2023		
	EPD-PB-OD-06-071323-2/EPD-PB-OD-066-0	71323-2	
Field Duplicate Pairs	EPD-PB-WA-04-071323-2/EPD-PB-WA-044-071323-2		
	EPD-PB-OD-03-071323-2/EPD-PB-OD-033-0	71323-2	
Field QC Blanks	EPD-PB-FB-02-071323-2, EPD-PB-FB-03-071	323-2, EPD-PB-MB-02	-071323-2, and EPD-PB-MB-03-071323-2

INTRODUCTION

This checklist summarizes the Stage 2A validation performed on the subject laboratory report, in accordance with the U.S. Environmental Protection Agency (EPA) *Guidance for Labeling Externally Validated Laboratory Analytical Data for Superfund Use* (January 2009). Analytical data were evaluated in general accordance with the Tetra Tech *Quality Assurance Project Plan East Palestine Train Derailment Site East Palestine, Columbiana County, Ohio, Revision 3* (April 2023), the Tetra Tech *Quality Assurance Project Plan, Superfund Technical Assessment and Response Team (START V), EPA Region 5, Revision 4* (August 2022), and the EPA *National Functional Guidelines (NFG) for Organic Superfund Methods Data Review* (November 2020).

OVERALL EVALUATION

No rejection or qualification of results was required for this data package. The results may be used as reported by the laboratory.



Data completeness:

Within Criteria	Exceedance/Notes
	Level II laboratory report does not include some required QC information; therefore, the Level IV laboratory report was used for this data validation effort.
	The Level II and Level IV laboratory reports did not contain quality control (QC) results for the laboratory reagent blank (LRB IHG230717C); however, the electronic data deliverable (EDD) results and the raw data results did contain the LRB results. No revisions or qualifications were applied since the LRB result is nondetect.
	The results for the field blank and media blank were reported in units of micrograms (µg) while the other field samples results were reported in units of µg, milligrams per cubic meter (mg/m3), and parts per million (ppm) (volume) in the laboratory report and only in units of ppm in the laboratory electronic data deliverable (EDD).
N	The site-specific QAPP specifies analysis of acrylates in air by Eurofins Analytics, LLC standard operating procedure (SOP) IHGC-001- v.22-3. The laboratory confirmed that NIOSH Method 1450M, which is mentioned in the laboratory deliverables, is equivalent to SOP IHGC-001-v.22-3; therefore, these method references may be used interchangeably.
	To facilitate sample reporting, large sample delivery groups may be logged by the laboratory separately by individual pages of the COC form. The ratio of field QC samples (field blanks, media blanks, field duplicates) to non-QC field samples is monitored independent of this validation and therefore the ratio of field QC samples to non-QC field samples was not verified during this validation. No qualifications were applied because all field sample results were non-detect.
	Note, the following fields in the laboratory EDD may be formatted as date only or date/time: Date_Collected, Date_Received, Date_Extracted, and Date_Analyzed. The time value was not required to be provided in the EDD. If no time value was provided, then the entered value may appear as date only or with a default value of 0:00, 00:00, or similar.



Sample preservation, receipt, and holding times:

Within Criteria	Exceedance/Notes
Y	

Method blanks:

Within Criteria	Exceedance/Notes
N	Nondetect results for laboratory method blank LMB IHG230717C and laboratory reagent blank LRB IHG230717C were reported as "0" in the laboratory EDD rather than at the reporting limit (RL). The laboratory was contacted on August 28, 2023, and agreed to report nondetect laboratory method blank and LRB results at the RL in future laboratory EDDs. No qualifications were applied.

Field blanks:

Within Criteria	Exceedance/Notes
Y	

Surrogates and labeled compounds:

Within Criteria	Exceedance/Notes
NA	

MS/MSDs:

Within Criteria	Exceedance/Notes
NA	



Laboratory duplicates:

Within Criteria	Exceedance/Notes
NA	

Field duplicates:

Within Criteria	Exceedance/Notes
Y	

LCSs/LCSDs:

Within Criteria	Exceedance/Notes
N	The site-specific QAPP requires a laboratory reagent blank (LRB), laboratory media blank (LMB), laboratory control sample (LCS), and LCS duplicate (LCSD) to be analyzed per batch of 20 samples. However, the laboratory analyzed 32 field samples in one sample preparation batch that included one LRB, laboratory media blank, LCS, and LCSD, when the batch should have included two LRBs, laboratory media blanks, LCSs, and LCSDs. The laboratory was contacted on August 8, 2023, regarding this deviation from the site-specific QAPP and agreed that they would follow the quality control (QC) sample frequency requirements in the site-specific QAPP in future reports. No qualifications were applied based on professional judgement because the QC sample results met the QAPP acceptance criteria, and the QC sample results from previous datasets for this project have met the QAPP acceptance criteria.

Sample dilutions:

Within Criteria	Exceedance/Notes
NA	



Re-extraction and reanalysis:

Within Criteria	Exceedance/Notes
NA	

MDLs/RLs:

Within Criteria	Exceedance/Notes
Y	Method detection limits were not reported. Non-detect sample results are reported as less than the RL in the laboratory report and at the RL (flagged U) in the validated EDD and attached analytical results summary.

Tentatively identified compounds:

Within Criteria	Exceedance/Notes
NA	

Other [None]:

Within Criteria	Exceedance/Notes
NA	



Overall Qualifications:

See results summary pages attached for changes to the laboratory qualifiers based upon this validation. The following is a list of qualifiers and definitions that may be used for the validation of this data package:

 be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate. 	 _	
J+ biased high. J- The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be biased low. NJ The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).	J	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample.
J- biased low. NJ The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).	J+	
biased low. NJ The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).	1-	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be
 NJ concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). 	J-	biased low.
 concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). 	NU	The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate
 be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate. 	INJ	concentration of the analyte in the sample.
be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).	D	The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not
The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	n	be present in the sample.
The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	U	The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).
1111 I the analyte has analyte has not detected at or above the above the above the above the above the boothing initial, which is considered approximate		The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate
due to deficiencies in one or more quality control criteria.	UJ	due to deficiencies in one or more quality control criteria.



E PALESTINE SITE - ER AIR ANALYTICAL RESULTS SUMMARY EUROFINS ANALYTICS, LLC REPORT NO. B198-117

Sample_ID	Method	CAS#	Analyte	Lab_Result Lab_Qual	RL	Units	VAL_Result VAL_Qual
EPD-PB-BKBA-01-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-BKBA-02-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-06-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-07-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-08-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-09-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-10-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-11-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-12-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-CM-14-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-DW-B-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-FB-02-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	2 U	2	ug	2 U
EPD-PB-FB-03-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	2 U	2	ug	2 U
EPD-PB-MB-02-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	2 U	2	ug	2 U
EPD-PB-MB-03-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	2 U	2	ug	2 U
EPD-PB-OD-01-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-02-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-03-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-033-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-04-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-05-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-06-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-066-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-OD-07-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-UW-F-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-WA-01-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-WA-02-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-WA-03-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-WA-04-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-WA-044-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-WA-05-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U
EPD-PB-WA-06-071323-2	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 U	0.0091	ppm	0.0091 U

Site Name	E Palestine Site - ER	TO/TOLIN No.	68HE0520F0032/0001EB201		
Document Tracking No.	2199b	TO/TOLIN NO.	08HE0320F0032/0001EB201		
Laboratory Report No.	B198-118	Laboratory	Eurofins Analytics, LLC – Ashland, VA		
Analyses	n-Butyl acrylate by NIOSH Method 1450M				
Samples and Matrix	29 air samples including 1 field blank, 1 media blank, and 2 field duplicate pairs				
Collection Date(s)	07/15/2023				
Field Duplicate Dairs	EPD-PB-CM-06-071523-1/EPD-PB-CM-066-	071523-1			
Field Duplicate Pairs	EPD-PB-WA-04-071523-1/EPD-PB-WA-044-071523-1				
Field QC Blanks	EPD-PB-FB-01-071523-1 and EPD-PB-MB-01-071523-1				

INTRODUCTION

This checklist summarizes the Stage 2A validation performed on the subject laboratory report, in accordance with the U.S. Environmental Protection Agency (EPA) *Guidance for Labeling Externally Validated Laboratory Analytical Data for Superfund Use* (January 2009). Analytical data were evaluated in general accordance with the Tetra Tech *Quality Assurance Project Plan East Palestine Train Derailment Site East Palestine, Columbiana County, Ohio, Revision 3* (April 2023), the Tetra Tech *Quality Assurance Project Plan, Superfund Technical Assessment and Response Team (START V), EPA Region 5, Revision 4* (August 2022), and the EPA *National Functional Guidelines (NFG) for Organic Superfund Methods Data Review* (November 2020).

OVERALL EVALUATION

No rejection or qualification of results was required for this data package. The results may be used as reported by the laboratory.



Data completeness:

Within Criteria	Exceedance/Notes
	Level II laboratory report does not include some required QC information; therefore, the Level IV laboratory report was used for this data validation effort.
	The Level II and Level IV laboratory reports did not contain quality control (QC) results for the laboratory reagent blank (LRB IHG230717D); however, the electronic data deliverable (EDD) results and the raw data results did contain the LRB results. No revisions or qualifications were applied since the LRB result is nondetect.
	The results for the field blank and media blank were reported in units of micrograms (µg) while the other field samples results were reported in units of µg, milligrams per cubic meter (mg/m3), and parts per million (ppm) (volume) in the laboratory report and only in units of ppm in the laboratory electronic data deliverable (EDD).
N	The site-specific QAPP specifies analysis of acrylates in air by Eurofins Analytics, LLC standard operating procedure (SOP) IHGC-001- v.22-3. The laboratory confirmed that NIOSH Method 1450M, which is mentioned in the laboratory deliverables, is equivalent to SOP IHGC-001-v.22-3; therefore, these method references may be used interchangeably.
	The ratio of field QC samples (field blanks, media blanks, field duplicates) to non-QC field samples is monitored independent of this validation and therefore the ratio of field QC samples to non-QC field samples was not verified during this validation. No qualifications were applied because all field sample results were non-detect.
	Note, the following fields in the laboratory EDD may be formatted as date only or date/time: Date_Collected, Date_Received, Date_Extracted, and Date_Analyzed. The time value was not required to be provided in the EDD. If no time value was provided, then the entered value may appear as date only or with a default value of 0:00, 00:00, or similar.



Sample preservation, receipt, and holding times:

Within Criteria	Exceedance/Notes
Y	

Method blanks:

Within Criteria	Exceedance/Notes
N	Nondetect results for laboratory method blank LMB IHG230717D and laboratory reagent blank LRB IHG230717D were reported as "0" in the laboratory EDD rather than at the reporting limit (RL). The laboratory was contacted on August 28, 2023, and agreed to report non-detect laboratory method blank and LRB results at the RL in future laboratory EDDs. No qualifications were applied.

Field blanks:

Within Criteria	Exceedance/Notes
N	Only one field blank and one media blank were included in this data package although the site-specific QAPP specifies the collection of one field blank and one media blank per 20 field samples. No qualifications were applied because all sample results were non-detect.

Surrogates and labeled compounds:

Within Criteria	Exceedance/Notes
NA	

MS/MSDs:

Within Criteria	Exceedance/Notes
NA	



Laboratory duplicates:

Within Criteria	Exceedance/Notes
NA	

Field duplicates:

Within Criteria	Exceedance/Notes
Y	

LCSs/LCSDs:

Within Criteria	Exceedance/Notes
N	The site-specific QAPP requires a laboratory reagent blank (LRB), laboratory media blank (LMB), laboratory control sample (LCS), and LCS duplicate (LCSD) to be analyzed per batch of 20 samples. However, the laboratory analyzed 29 field samples in one sample preparation batch that included one LRB, laboratory media blank, LCS, and LCSD, when the batch should have included two LRBs, laboratory media blanks, LCSs, and LCSDs. The laboratory was contacted on August 8, 2023, regarding this deviation from the site-specific QAPP and agreed that they would follow the quality control (QC) sample frequency requirements in the site-specific QAPP in future reports. No qualifications were applied based on professional judgement because the QC sample results met the QAPP acceptance criteria, and the QC sample results from previous datasets for this project have met the QAPP acceptance criteria.

Sample dilutions:

Within Criteria	Exceedance/Notes
NA	

Re-extraction and reanalysis:

Within Criteria	Exceedance/Notes
NA	



MDLs/RLs:

Within Criteria	Exceedance/Notes
Y	Method detection limits were not reported. Non-detect sample results are reported as less than the RL in the laboratory report and at the RL (flagged U) in the validated EDD and attached analytical results summary.

Tentatively identified compounds:

Within Criteria	Exceedance/Notes
NA	

Other [None]:

Within Criteria	Exceedance/Notes
NA	

Overall Qualifications:

See results summary pages attached for changes to the laboratory qualifiers based upon this validation. The following is a list of qualifiers and definitions that may be used for the validation of this data package:

U	The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).
R	The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample.
NJ	The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample.
J-	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be biased low.
J+	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be biased high.
J	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample.



UJ The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate due to deficiencies in one or more quality control criteria.



E PALESTINE SITE - ER AIR ANALYTICAL RESULTS SUMMARY EUROFINS ANALYTICS, LLC REPORT NO. B198-118

Sample_ID	Method	CAS#	Analyte	Lab_Result L	Lab_Qual	RL	Units	VAL_Result	VAL_Qual
EPD-PB-BKBA-01-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-BKBA-02-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-06-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-066-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-07-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-08-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-09-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-10-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-11-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-12-071523-1	NIOSH Method 1450M		n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-CM-14-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	J	0.0091	ppm	0.0091	U
EPD-PB-DW-C-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	J	0.0091	ppm	0.0091	U
EPD-PB-FB-01-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	2 l	J	2	ug	2	U
EPD-PB-MB-01-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	2 נ	J	2	ug	2	U
EPD-PB-OD-01-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	J	0.0091	ppm	0.0091	U
EPD-PB-OD-02-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	J	0.0091	ppm	0.0091	U
EPD-PB-OD-03-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-OD-04-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-OD-05-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-OD-06-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-OD-07-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-UW-G-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-WA-01-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-WA-02-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-WA-03-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-WA-04-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-WA-044-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-WA-05-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U
EPD-PB-WA-06-071523-1	NIOSH Method 1450M	141-32-2	n-Butyl acrylate	0.0091 l	U	0.0091	ppm	0.0091	U

Site Name E Palestine Site - ER			TO/TOLIN No.	68HE0520F0032/0001EB201			
Document Tracking No.	2199c		TO/TOLIN NO.	08HE0520F0052/0001EB201			
Laboratory Report No.	B198-119		Laboratory	Eurofins Analytics, LLC – Ashland, VA			
Analyses	2-Ethylhexyl acrylate and n-butyl acrylate by laboratory standard operating procedure - (SOP) IHGC-P029						
Samples and Matrix	Nine air samples including one field blank						
Collection Date(s)	07/14/2023						
Field Duplicate Pairs	None						
Field QC Blanks	EPD-ST-FB-071423-1						

INTRODUCTION

This checklist summarizes the Stage 2A validation performed on the subject laboratory report, in accordance with the U.S. Environmental Protection Agency (EPA) *Guidance for Labeling Externally Validated Laboratory Analytical Data for Superfund Use* (January 2009). Analytical data were evaluated in general accordance with the Tetra Tech *Quality Assurance Project Plan East Palestine Train Derailment Site East Palestine, Columbiana County, Ohio, Revision 3* (April 2023), the Tetra Tech *Quality Assurance Project Plan, Superfund Technical Assessment and Response Team (START V), EPA Region 5, Revision 4* (August 2022), and the EPA *National Functional Guidelines (NFG) for Organic Superfund Methods Data Review* (November 2020).

OVERALL EVALUATION

No rejection or qualification of results was required for this data package. The results may be used as reported by the laboratory.



Data completeness:

Within Criteria	Exceedance/Notes
	The results for the field blank and media blank were reported in units of micrograms (µg) while the other field samples results were reported in units of µg, milligrams per cubic meter (mg/m3), and parts per million (ppm) (volume) in the laboratory report and only in units of ppm in the laboratory electronic data deliverable (EDD).
N	Rohm & Haas IH9805 is cited in the AIHA certification as "IHGC-P029) and may be cited by the abbreviation "Rohm & Hass IH9805" or "IHGC-P029" interchangeably throughout the laboratory report. Note, the following fields in the laboratory EDD may be formatted as date only or date/time: Date_Collected, Date_Received, Date_Extracted, and Date_Analyzed. The time value was not required to be provided in the EDD. If no time value was provided, then the entered value may appear as date only or with a default value of 0:00, 00:00, or similar.

Sample preservation, receipt, and holding times:

Within Criteria	Exceedance/Notes
Y	

Method blanks:

Within Criteria	Exceedance/Notes
N	Nondetect results for laboratory method blank LMB IHG230717G and laboratory reagent blank LRB IHG230717G were reported as "0" in the laboratory EDD rather than at the reporting limit (RL). The laboratory was contacted on August 28, 2023, and agreed to report nondetect laboratory method blank and LRB results at the RL in future laboratory EDDs.

Field blanks:

Within Criteria	Exceedance/Notes
Y	



Surrogates and labeled compounds:

Within Criteria	Exceedance/Notes
NA	

MS/MSDs:

Within Criteria	Exceedance/Notes
NA	

Laboratory duplicates:

Within Criteria	Exceedance/Notes
NA	

Field duplicates:

Within Criteria	Exceedance/Notes
N	Per the site-specific QAPP, 1 field duplicate sample is required per 20 samples collected. However, fewer than 1 field duplicate sample per 20 samples were collected with this sample group. Based on professional judgement, no qualifications were applied.

LCSs/LCSDs:

Within Criteria	Exceedance/Notes
N	The laboratory report(s) and the laboratory EDD have one or more minor discrepancies in the LCS/LCSD results (+/- 1 ug), RPDs (+/- 2%) and/or percent recoveries (+/- 1%) that were verified with the laboratory to be a significant figures issue. No qualifications were applied.



Sample dilutions:

Within Criteria	Exceedance/Notes
NA	

Re-extraction and reanalysis:

Within Criteria	Exceedance/Notes
NA	

MDLs/RLs:

Within Criteria	Exceedance/Notes
Y	Method detection limits were not reported. Non-detect sample results are reported as less than the RL in the laboratory report and at the RL (flagged U) in the validated EDD and attached analytical results summary.

Tentatively identified compounds:

Within Criteria	Exceedance/Notes
NA	

Other [None]:

Within Criteria	Exceedance/Notes
NA	



Overall Qualifications:

See results summary pages attached for changes to the laboratory qualifiers based upon this validation. The following is a list of qualifiers and definitions that may be used for the validation of this data package:

J The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample. J+ The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be biased high. J- The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be biased low. NJ The analyte indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample and may be biased low. NJ The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte is the approximate concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). UJ The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate due to deficiencies in one or more quality control criteria.		
J+ biased high. J- The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be biased low. NJ The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). III The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	J	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample.
J- biased low. NJ The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). III The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	J+	
biased low. NJ The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). III The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	1-	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be
NJ concentration of the analyte in the sample. R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). III The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	J -	biased low.
R The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). III The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	NI	The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate
 be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate 	INJ	concentration of the analyte in the sample.
be present in the sample. U The analyte was analyzed for, but was not detected at or above the associated value (reporting limit). III The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	D	The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not
The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate	n	be present in the sample.
	U	The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).
UJ due to deficiencies in one or more quality control criteria		The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate
	01	due to deficiencies in one or more quality control criteria.



E PALESTINE SITE - ER AIR ANALYTICAL RESULTS SUMMARY EUROFINS ANALYTICS, LLC REPORT B198-119

Sample_ID	Method	CAS#	Analyte	Lab_Result Lab_Qual	RL	Units VAL_Result	VAL_Qual
EPD-ST-8H-DW-B-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm 0.015	U
EPD-ST-8H-DW-B-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-8H-UW-F-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm 0.015	U
EPD-ST-8H-UW-F-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-8H-WA-01-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm 0.015	U
EPD-ST-8H-WA-01-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-8H-WA-02-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm 0.015	U
EPD-ST-8H-WA-02-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-8H-WA-03-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.014 U	0.014	ppm 0.014	U
EPD-ST-8H-WA-03-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-8H-WA-04-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm 0.015	U
EPD-ST-8H-WA-04-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-8H-WA-05-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm 0.015	U
EPD-ST-8H-WA-05-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-8H-WA-06-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm 0.015	U
EPD-ST-8H-WA-06-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm 0.01	U
EPD-ST-FB-071423-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	2.8 U	2.8	ug 2.8	U
EPD-ST-FB-071423-1	IHGC-P029	141-32-2	n-Butyl acrylate	1.3 U	1.3	ug 1.3	U

Site Name	E Palestine Site - ER		
Document Tracking No.	2199d	TO/TOLIN No.	68HE0520F0032/0001EB201
Laboratory Report No.	B198-120	Laboratory	Eurofins Analytics, LLC – Ashland, VA
Analyses	2-Ethylhexyl acrylate and n-butyl acrylate b	y laboratory standard	operating procedure (SOP) IHGC-P029
Samples and Matrix	Nine air samples including one field blank		
Collection Date(s)	07/15/2023		
Field Duplicate Pairs	None		
Field QC Blanks	ield QC Blanks EPD-ST-FB-071523-1		

INTRODUCTION

This checklist summarizes the Stage 2A validation performed on the subject laboratory report, in accordance with the U.S. Environmental Protection Agency (EPA) *Guidance for Labeling Externally Validated Laboratory Analytical Data for Superfund Use* (January 2009). Analytical data were evaluated in general accordance with the Tetra Tech *Quality Assurance Project Plan East Palestine Train Derailment Site East Palestine, Columbiana County, Ohio, Revision 3* (April 2023), the Tetra Tech *Quality Assurance Project Plan, Superfund Technical Assessment and Response Team (START V), EPA Region 5, Revision 4* (August 2022), and the EPA *National Functional Guidelines (NFG) for Organic Superfund Methods Data Review* (November 2020).

OVERALL EVALUATION

No rejection or qualification of results was required for this data package. The results may be used as reported by the laboratory.



Data completeness:

Within Criteria	Exceedance/Notes
	The results for the field blank and media blank were reported in units of micrograms (µg) while the other field samples results were reported in units of µg, milligrams per cubic meter (mg/m3), and parts per million (ppm) (volume) in the laboratory report and only in units of ppm in the laboratory electronic data deliverable (EDD).
N	Rohm & Haas IH9805 is cited in the AIHA certification as "IHGC-P029) and may be cited by the abbreviation "Rohm & Hass IH9805" or "IHGC-P029" interchangeably throughout the laboratory report. Note, the following fields in the laboratory EDD may be formatted as date only or date/time: Date_Collected, Date_Received, Date_Extracted, and Date_Analyzed. The time value was not required to be provided in the EDD. If no time value was provided, then the entered value may appear as date only or with a default value of 0:00, 00:00, or similar.

Sample preservation, receipt, and holding times:

Within Criteria	Exceedance/Notes
Y	

Method blanks:

Within Criteria	Exceedance/Notes
N	Nondetect results for laboratory method blank LMB IHG230628C and laboratory reagent blank LRB IHG230628C were reported as "0" in the laboratory EDD rather than at the reporting limit (RL). The laboratory was contacted on August 28, 2023, and agreed to report nondetect laboratory method blank and LRB results at the RL in future laboratory EDDs.

Field blanks:

Within Criteria	Exceedance/Notes
Y	



Surrogates and labeled compounds:

Within Criteria	Exceedance/Notes
NA	

MS/MSDs:

Within Criteria	Exceedance/Notes
NA	

Laboratory duplicates:

Within Criteria	Exceedance/Notes
NA	

Field duplicates:

Within Criteria	Exceedance/Notes
N	Per the site-specific QAPP, 1 field duplicate sample is required per 20 samples collected. However, fewer than 1 field duplicate sample per 20 samples were collected with this sample group. Based on professional judgement, no qualifications were applied.

LCSs/LCSDs:

Within Criteria	Exceedance/Notes					
N	The laboratory report(s) and the laboratory EDD have one or more minor discrepancies in the LCS/LCSD results (+/- 1 ug), RPDs (+/- 2%) and/or percent recoveries (+/- 1%) that were verified with the laboratory to be a significant figures issue. No qualifications were applied.					



Sample dilutions:

Within Criteria	Exceedance/Notes
NA	

Re-extraction and reanalysis:

Within Criteria	Exceedance/Notes
NA	

MDLs/RLs:

Within Criteria	Exceedance/Notes					
Y	Method detection limits were not reported. Non-detect sample results are reported as less than the RL in the laboratory report and at the RL (flagged U) in the validated EDD and attached analytical results summary.					

Tentatively identified compounds:

Within Criteria	Exceedance/Notes
NA	

Other [None]:

Within Criteria	Exceedance/Notes
NA	



Overall Qualifications:

See results summary pages attached for changes to the laboratory qualifiers based upon this validation. The following is a list of qualifiers and definitions that may be used for the validation of this data package:

J	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample.						
J+	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be						
	biased high.						
	The analyte was positively identified; the associated value is the approximate concentration of the analyte in the sample and may be						
J-	biased low.						
NJ	The analysis indicates the presence of an analyte that has been "tentatively identified" and the associated value is the approximate						
INJ	concentration of the analyte in the sample.						
R	The sample result is rejected as unusable due to serious deficiencies in one or more quality control criteria. The analyte may or may not						
ĸ	be present in the sample.						
U	The analyte was analyzed for, but was not detected at or above the associated value (reporting limit).						
	The analyte was analyzed for, but was not detected at or above the associated value (reporting limit), which is considered approximate						
UJ	due to deficiencies in one or more quality control criteria.						



E PALESTINE SITE - ER AIR ANALYTICAL RESULTS SUMMARY EUROFINS ANALYTICS, LLC REPORT NO. B198-120

Sample_ID	Method	CAS#	Analyte	Lab_Result Lab_Qual	RL	Units	VAL_Result VAL_Qual
EPD-ST-8H-DW-C-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.013 U	0.013	ppm	0.013 U
EPD-ST-8H-DW-C-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.009 U	0.009	ppm	0.009 U
EPD-ST-8H-UW-G-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.013 U	0.013	ppm	0.013 U
EPD-ST-8H-UW-G-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.009 U	0.009	ppm	0.009 U
EPD-ST-8H-WA-01-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.013 U	0.013	ppm	0.013 U
EPD-ST-8H-WA-01-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.009 U	0.009	ppm	0.009 U
EPD-ST-8H-WA-02-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.013 U	0.013	ppm	0.013 U
EPD-ST-8H-WA-02-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.009 U	0.009	ppm	0.009 U
EPD-ST-8H-WA-03-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.015 U	0.015	ppm	0.015 U
EPD-ST-8H-WA-03-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.01 U	0.01	ppm	0.01 U
EPD-ST-8H-WA-04-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.013 U	0.013	ppm	0.013 U
EPD-ST-8H-WA-04-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.009 U	0.009	ppm	0.009 U
EPD-ST-8H-WA-05-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.013 U	0.013	ppm	0.013 U
EPD-ST-8H-WA-05-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.009 U	0.009	ppm	0.009 U
EPD-ST-8H-WA-06-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	0.014 U	0.014	ppm	0.014 U
EPD-ST-8H-WA-06-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	0.009 U	0.009	ppm	0.009 U
EPD-ST-FB-071523-1	IHGC-P029	103-11-7	2-Ethylhexyl acrylate	2.8 U	2.8	ug	2.8 U
EPD-ST-FB-071523-1	IHGC-P029	141-32-2	n-Butyl acrylate	1.3 U	1.3	ug	1.3 U