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## CHECKLIST FOR REVIEWING EPA QUALITY MANAGEMENT PLANS

This checklist will be used to review the Quality Management Plans (QMPs) that are submitted to the Quality Staff of the Office of Environmental Information (OEI) for Agency review under EPA Order 5360.1 A2. Items from this checklist are discussed in detail in Chapter 3 of EPA Manual 5360 A1 and in *EPA Requirements for Quality Management Plans (QA/R-2)*. Consult these resources for more information on the items below.

Note that all items below must be included in a QMP. If an item is not relevant, an explanation must be provided. Also note that process may either be described or <u>referenced</u> in the QMP; however, all references should be readily accessible within the organization and provided to the Quality Staff with the QMP.

		Page(s)	Comments			
MA	MANAGEMENT AND ORGANIZATION					
1.	Signed and dated by senior manager?					
2.	Signed and dated by senior line management?					
3.	Signed and dated QA manager?					
4.	Includes signature lines for Quality Staff approval?					
5.	Includes signature lines for OEI approval?					
6.	Includes statement of the organization's QA policy?					
	6a. QA policy statement includes general objectives/goals?					
	6b. QA policy statement includes allocation of intramural, extramural, and travel funds and personnel?					

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		Page(s)	Comments
7.	Includes organizational chart?		
	7a. Organizational chart identifies all components of organization?		
	7b. Organizational Chart identifies position of QA manager?		
	7c. Organizational Chart identifies lines of reporting of the QA manager?		
	7d. Organization Chart identifies any other QA staff?		
8.	Includes discussion of authorities of the QA manager and staff?		
9.	Documents the independence of QA manager?		
10.	Describes procedures to ensure QA staff have access to appropriate levels of management?		
11.	Discusses technical activities or programs that require quality management?		
12.	Discusses where oversight of delegated or extramural programs is needed?		

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	<b>D</b> ()	G
	Page(s)	Comments
13. Identifies where internal coordination of QA and QC activities among organizations is needed?		
14. Discusses how management assures understanding and implementation in all programs?		
15. Describes process for resolving disputes?		
QUALITY SYSTEM COMPONENTS		
16. Includes description of quality system?		
17. Describes principal quality system components (e.g., quality system documentation, annual reviews and planning, project-specific quality documentation? (Note, identify components in Column 3.)		
18. Description of components includes how they are implemented?		
19. Description of components includes responsibilities of management and staff?		
20. Lists tools for implementing each component (e.g., QMPs, Quality Systems Audits, Training Plans, QA Project Plans? (Note: list tools in Column 3.)		

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		Page(s)	Comments
21. Id	dentifies internal organizations that develop QMPs?		
	dentifies review and approval procedures for these internal QMPs?		
pe	ncludes assurance that QA responsibility is incorporated into performance standards (consistent with Agency personnel policy)?		
QUA	LIFICATIONS AND TRAINING		
24. S	states policy regarding QA training for management and staff?		
	Describes process for identifying, ensuring, and documenting that personnel have necessary quality-related qualifications?		
	Describes process for ensuring personnel maintain quality-related qualifications?		
	Describes process for identifying the need for quality-related etraining based on changing requirements?		
	ncludes roles, responsibilities, and authorities in description of bove processes?		

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	Page(s)	Comments
PROCUREMENT OF ITEMS AND SERVICES		
29. Describes process for reviewing and approving all extramural agreements (grants, cooperative agreements and contracts)?		
29a. Review process ensures documents are complete and accurate?		
29b. Review process ensures agreement clearly describes the item or service needed?		
29c. Review process ensures agreement describes the associated technical and quality requirements?		
29d. Review process ensures agreement describes the quality system elements for which the supplier is responsible?		
29e. Review process ensures that the supplier's conformance to the customer's requirements will be verified?		
30. Describes process for reviewing and approving applicable responses to solicitations to ensure that they satisfy all technical and quality requirements?		
30a. Review process ensures the review of evidence of the supplier's capability to satisfy EPA quality requirements?		
30b. Review process ensures procured items and services are acceptable?		
31. Describes process for review and approval of suppliers' quality-related documentation (e.g., QA Project Plans and QMPs)?		

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		Page(s)	Comments
	of any policy and criteria for delegations of oct Plans and QMPs?	Tage(3)	Comments
33. Describes process to satisfied?	ensure EPA extramural agreement policies		
34. Includes roles, responsible above processes?	onsibilities, and authorities in description of		
DOCUMENTS AND	RECORDS		
	or identifying quality-related documents and lectronic) requiring control?		
	or preparing, reviewing, approving, issuing, g, and revising documents and records?		
37. Describes process for accurately reflect co	or ensuring that records and documents ompleted work?		
including transmitta	or maintaining documents and records I, distribution, retention, access, preservation, I, removal of obsolete documentation, and		
_	or establishing and implementing appropriate d confidentiality procedures for evidentiary		

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		<b>D</b> ()	a .
		Page(s)	Comments
40.	Above processes comply with EPA Order 2160 and EPA Directive 2100, Chapter 10?		
41.	Includes roles, responsibilities, and authorities in description of above processes?		
CO	MPUTER HARDWARE AND SOFTWARE		
42.	Describes process for developing, installing, testing, using, maintaining, controlling, and documenting computer hardware and software?		
43.	Describes process for assessing and documenting the impact of changes to user requirements and/or the hardware and software on performance?		
44.	Describes process for evaluating purchased hardware and software?		
45.	Describes process for ensuring that data and information produced from or collected by computers meet applicable requirements and standards?		
46.	Includes roles, responsibilities, and authorities in description of above processes?		
47.	Are the requirements of EPA Directive 2100 are addressed in the above processes?		

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			Page(s)
PI.	ANNING		
	Includes a de	escription of the systematic planning process for tal data operations?	
	-	process include identification and involvement of all mers and suppliers?	
		process include description of the project goal, ives, and questions and issues to be addressed?	
		process include identification of project schedule, rces, milestones, and any applicable requirements?	
	of data	process include identification of the type and quantity a needed and how the data will be used to support the et's objectives?	
		process include specification of performance criteria easuring quality?	
		process include specification of needed QA and QC ies to assess the quality performance criteria?	
	the dat	process include description of how, when, and where ta will be obtained (including existing data) and fication of any constraints on data collection?	
	will be	process include description of how the acquired data e analyzed, evaluated, and assessed against its ed use and the quality performance criteria?	

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		Page(s)	Comments
49.	Describes process for developing, reviewing, approving, implementing, and revising QA Project Plans?		
50.	Describes process for evaluating and qualifying data collected for other purposes or from other sources?		
51.	Includes roles, responsibilities, and authorities in description of above processes?		
IM	PLEMENTATION OF WORK PROCESSES		
52.	Describes process for ensuring that work is performed according to planning and technical documents?		
53.	Describes process for identifying operations needing procedures?		
54.	Describes process for preparation, review, approval, revision, and withdrawal of these procedures?		
55.	Describes policy for use of these procedures?		
56.	Describes process for controlling and documenting the release, change, and use of planned procedures?		

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		Page(s)	Comments
	56a. Process includes description of necessary approvals?		
	56b. Process includes removal of obsolete documentation from work areas?		
	56c. Process includes verification that the changes are made as prescribed?		
57.	Includes roles, responsibilities, and authorities in description of above process?		
AS	SESSMENT AND RESPONSE		
58.	Describes the process for assessing the adequacy of the quality system at least annually?		
59.	Describes the process for planning, implementing and documenting assessments and reporting results to management?		
	59a. Process includes selecting an assessment tool, the expected frequency, and the roles and responsibilities of assessors?		
	59b. Process includes determining the level of competence, experience and training needed for assessment personnel?		
	59c. Process includes ensuring that personnel have no real or perceived conflict of interest, and have no direct involvement or responsibility for the work being assessed?		

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	Page(s)	Comments
59d. Process includes ensuring that personnel conducting assessments have sufficient authority, access to programs and managers, access to documents and records, and organizational freedom?		
60. Describes process for management's review of, and response to, findings?		
61. Describes process for identifying how and when corrective actions are to be taken in response to the findings of the assessment?		
61a. Process includes ensuring corrective actions are made promptly?		
61b. Process includes confirming the implementation and effectiveness of any corrective action?		
61c. Process includes documenting actions?		
62. Describes process for addressing disputes encountered as a result of assessments?		
63. Includes roles, responsibilities, and authorities in description of above processes?		

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	Page(s)	Comments
QUALITY IMPROVEMENT		
64. Describes process for ensuring that conditions adverse to quality are prevented, identified promptly, corrected promptly and that actions are taken toward prevention, documented and actions tracked to closure?		
65. Describes process for encouraging staff to establish communications between customers and suppliers, identify process improvement opportunities, and identify and propose solutions for problems?		
66. Includes roles, responsibilities, and authorities in description of above processes?		
OTHER REVIEW CRITERIA	•	
67. Are regulatory or other citations accurate?		
68. Are there any inconsistencies in the text?		
69. Is the writing clear?		
70. Are organizational units identified consistent with the most recent reorganization?		
71. Are past QS management assessment findings resolved? (Put date of Final Report in Column 3.)		

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72. Are activities described in the QMP consistent with QA Annual Report and Work Plans?		
73. Are tasks proposed for other organizations not covered solely by this QMP documented elsewhere (e.g., in another organization's QMP)?		

## ATTACHMENT 2 EPA-SPECIFIC QUALITY MANAGEMENT PLAN REQUIREMENTS

The following items from the *Checklist for Reviewing EPA Quality Management Plans* are only applicable for EPA QMPs required under EPA Order 5360.1 CHG 1 (July 1998):

- 4. Signature line for Quality Staff
- 5. Signature lines for OEI approval<sup>1</sup>
- 15. Dispute resolution process
- 23. Performance standards
- 30a. Review and approval of responses to solicitations to ensure they satisfy EPA quality requirements
- 31. Review and approval of quality-related documentation from suppliers
- 32. Policy and criteria for delegating approval of quality-related documentation
- 33. Process to ensure EPA contracting policies satisfied
- 40. Conformance to EPA Order 2160 and EPA Directive 2100, Chapter 10

<sup>&</sup>lt;sup>1</sup>For non-EPA organizations, if EPA approval of a QMP is required, the approval page must include a section for the signature of the responsible EPA official.