USEPA REGION 4 SUPERFUND & EMERGENCY MANAGEMENT DIVISON QAPP CHECKLIST/ FUND-LEAD UNIFORM FEDERAL POLICY (UFP) PROJECT

QAPP Title: Project Location: Originating Organization: QAPP Date: Receipt Date: Review Date: Reviewer: EPA Regional Project Manager: EPA Project Officer/Contracting Officer's Representative:

Topic covered in accordance with requirements: \Box Yes \Box No

 \Box Yes - Indicates that the topic/element was covered in sufficient detail to meet EPA's requirements as specified in this checklist.

 \Box No - Indicates that the topic/element covered in the QAPP does not provide sufficient detail to meet EPA's requirements or the topic is entirely missing from the document.

Element	Meets Requirements	□ Yes	□ No
QAPP Worksheet #1 & 2: Title and Approval Page			
Project Identifying Information: Site name/project name, Site location/number, Contract/Work assignment number	□ Yes □ No		
Lead Organization (Federal department, agency, or program; usually the entity that owns the facility or installation), Lead Organization Project Manager (name/title/signature/date), Lead Organization Quality Manager (name/title/signature/date).	□ Yes □ No		
Federal Regulatory Agency (name/title/signature/date) – USEPA Region 4 Remedial Project Manager/Designated Approving Official -or- Remedial Project Manager and Quality Assurance Manager	□ Yes □ No		
State Regulatory Agency (name/title/signature/date)	□ Yes □ No		
Other stakeholders as needed, including the	\Box Yes \Box No		

organization preparing the QAPP		
Plans and reports from previous	□ Yes	🗆 No
investigations relevant to this project		
QAPP Worksheet #3 & 5: Project		
Organization and QAPP Distribution		
Organization chart provided: Depicts key	□ Yes	🗆 No
personnel, lines of authority, and lines of		
communication among the lead agency, prime		
contractor, subcontractors, and regulatory		
agencies		
Documents recipients of controlled copies of	□ Yes	🗆 No
the QAPP (use asterisks on chart to designate		
QAPP recipients)		
QAPP Worksheet #4, 7, and 8: Personnel		
Qualifications and Sign-off Sheet		
Identifies key project personnel for	□ Yes	🗆 No
organizations performing tasks (such as prime		
contractor and subcontractors): individuals'		
project titles or roles, qualifications, and any		
specialized/non-routine training, certifications		
or clearances required by the project,		
signatures/dates (signatures indicate personnel		
have read and agree to implement the QAPP		
as written)		
QAPP Worksheet #6: Communication		
Pathways		
Documents specific issues (communication	□ Yes	\Box No
drivers) that will trigger the need to		
communicate with other project personnel or		
stakeholders; ensures procedures are in place		
for providing the appropriate notifications and		
generating the appropriate documentation		
when handling important communications		
QAPP Worksheet #9: Project Planning		
Session Summary		
Provides a worksheet for each internal and	\Box Yes	\Box No
external project planning session (including		
phone, web-conferencing, and/or face-to-face)		
Each worksheet provides record of	\Box Yes	\Box No
participants, key decisions or agreements		
reached, and action items (may attach meeting		
minutes)		
Identifies all electronic data deliverables	□ Yes	\Box No
(EDDs) that will be submitted for the project		

explains in specific terms how all data will be		
used; identifies information inputs consistent		
with decisions made during project scoping		
consistent with QAPP Worksheet #9		
Specifies the target (statistical) populations	\Box Yes	\Box No
and characteristics of interest; defines		
spatial/temporal limits and the scale of		
inference - which (statistical) populations will		
be represented by which data; develops		
focused list of target analytes		
Defines the parameter(s) of interest, specify	\Box Yes	\Box No
the types of inference and which sample		
results will be used to support which		
decisions. Uses "ifthen" statements for		
decision problems and/or the estimator and		
estimation procedure for estimation problems		
Specifies probability limits for decision errors	□ Yes	□ No
for projects that involve hypothesis testing		
and/or specifies performance (new data) or		
acceptance (existing data) criteria for		
estimations or other analytic approaches		
Briefly explains the rationale for the sampling	□ Yes	🗆 No
design; refers to subsequent worksheets for		
sampling design details and analysis design		
requirements		
Assesses what analytical resources will meet		
the analytical needs (Regional laboratory,	□ Yes	🗆 No
CLP, direct contract, subcontract), including		
any special requests or modified analysis for		
the Regional laboratory or CLP.		
QAPP Worksheet #12: Measurement		
Performance Criteria		
Provides a worksheet for each type of field or	□ Yes	🗆 No
laboratory measurement; for analytical		
methods, criteria are determined for each		
matrix, analyte, and concentration level -or- If		
using Regional laboratory/Contract		
Laboratory Program (CLP), cites the		
current Laboratory Services Branch		
Laboratory Operations and Quality		
Assurance Manual and/or relevant CLP		
Statements of Work.		
Each worksheet provides quantitative	□ Yes	🗆 No
measurement performance criteria in terms of		

precision, bias, and sensitivity -or- If using		
Regional laboratory/Contract Laboratory		
Program (CLP), cites the current		
Laboratory Services Branch Laboratory		
Operations and Quality Assurance Manual		
and/or relevant CLP Statements of Work.		
QAPP Worksheet #13: Secondary Data		
Uses and Limitations		
Identifies sources of secondary data (sampling	\Box Yes \Box] No
and testing data collected during previous		
investigations, historical data, background		
information, interviews, modeling data,		
photographs, aerial photographs, topographic		
maps, and published literature)		
Discusses the rationale for using this data and	\Box Yes] No
explains its relevance to the project		
Identifies factors affecting the reliability of	□ Yes □] No
data and limitations on data use, including		
how limitations will be communicated to all		
end data users and stakeholders		
QAPP Worksheet #14/16: Project Tasks &		
Schedule		
Provides a summary of key on-site and off-	□ Yes □	No
site activities, the person or group responsible		
for each activity, planned start and end dates,		
deliverables to be produced, and deliverable		
due dates (may be table or Gantt Chart)		
QAPP Worksheet #15: Project Action		
Limits and Laboratory-Specific		
Detection/Quantitation Limits		
Provides a worksheet for each type of field or	□ Yes □	No
laboratory measurement; criteria are		
determined for each matrix, analyte,		
analytical method and concentration level		
Identifies Project Action Limit (actual	□ Yes □	No
numerical criteria) for each analyte and the		
reference upon which it is based (such as		
MCLs or other ARARs, risk assessment		
screening levels, etc.); identifies Project		
Quantitation Limit Goal below the Project		
Action Limit; highlights the critical		
contaminants/analytes for project decision-		
making		
Provides laboratory-specific detection and	□ Yes □	No

quantitation limits for comparison to Project		
Quantitation Limit Goal. Laboratory provides		
documentation that demonstrates precision		
and bias at the laboratory-specific quantitation		
limit (at lowest calibration standard) -or- If		
using Regional laboratory/Contract		
Laboratory Program (CLP), cites the		
current Laboratory Services Branch		
Laboratory Operations and Quality		
Assurance Manual and/or relevant CLP		
Statements of Work for quantitation limit.		
(The laboratory-specific detection limit will		
not be available during planning.)		
QAPP Worksheet #17: Sampling Design		
and Rationale		
Provides design of the sampling/collection	□ Yes	□ No
network, including physical and temporal		
boundaries, basis for dividing the site into		
decision units, basis for number and		
placement of samples, sample location maps		
or diagrams, alternate locations, process for		
determining sample locations in the field (if		
applicable), and field condition contingencies		
Provides a discussion regarding the basis for	□ Yes	🗆 No
selection of probability-based designs vs.		
judgmental designs		
QAPP Worksheet #18: Sampling Locations		
and Methods		
Provides a table with type and number of	□ Yes	□No
samples required for collection such as		
surface soil, subsurface soil, or groundwater,		
preferably by individual Sample ID and		
collection frequency (if applicable), though		
sample groups may be listed in a single row		
Identifies each sample type using matrix	□ Yes	
codes and descriptions found in the Region 4		
Reference Values for EQuIS.		
Uses existing Station IDs where available in	□ Yes	□No
EQuIS for the planned location (matched by		
latitude/longitude).		
Provides the sample collection method for	\Box V	🗆 No
	YAC	
-	\Box Yes	
each sample or sample group and references	\Box Yes	
-		

QAPP	
Provides the analytes or analyte groups for	\Box Yes \Box No
each sample or sample group	
QAPP Worksheet #19 and 30: Sample	
Containers, Preservation, and Hold Times	
Provides a worksheet for each laboratory used	□ Yes □ No □ Not Applicable –
and lists any required	using only Regional
accreditations/certifications for the laboratory;	laboratory/CLP
attaches accreditations/certifications to the	
QAPP	
For each analyte/analyte group and matrix	\Box Yes \Box No \Box Not Applicable –
pair, provides the analytical method reference,	using only Regional
accreditation expiration date for the	laboratory/CLP
laboratory for that analyte/matrix/method	
combination (if global expiration date, this	
may be in the header	
For each analyte/analyte group, matrix, and	\Box Yes \Box No
analytical method, provides container(s)	
(Number, size, and type per sample),	
preservation requirements, preparation	
holding time, analytical holding time, and	
data package turnaround QAPP Worksheet #20: Field Quality	
Control Summary	
For each matrix and analyte/analytical group	\Box Yes \Box No
pair, provides a summary of the number of	
field samples, the number and types of field	
QC samples to be collected, and the total	
number of analyses (field and field QC	
samples combined).	
QAPP Worksheet #21: Field SOPs	
Lists SOPs (including title, revision, date, and	\Box Yes \Box No
originating organization) containing detailed	
procedures for all field activities, including	
sample collection, sample preservation,	
equipment cleaning and decontamination,	
equipment testing, maintenance, and	
inspection, and sampling handling and	
custody and notes any project-specific options	
or modifications, if applicable	
Referenced field SOPs are attached to the	\Box Yes \Box No
QAPP	
QAPP Worksheet #22: Field Equipment	
Calibration, Maintenance, Testing and	

Inspection	
Provides a list of all in-situ testing	\Box Yes \Box No
instruments and field equipment	
Documents the procedures for calibrating,	\Box Yes \Box No
maintaining, testing, and/or inspecting all	
field equipment	
Identifies the individual(s) responsible for	\Box Yes \Box No
field equipment	
Includes frequency, acceptance criteria, and	\Box Yes \Box No
corrective action or references and attaches	
the relevant SOP or manufacturer's	
instructions	
QAPP Worksheet #23: Analytical SOPs	
List SOPs (including title, revision, and date)	\Box Yes \Box No
containing the specific sample preparation	
and analytical procedures to be used to	
perform on-site or fixed laboratory analysis	
for each matrix/analytical group; indicate	
whether the procedure produces screening or	
definitive data; note any project-specific	
options or modifications, if applicable -or- If	
using Regional laboratory/Contract	
Laboratory Program (CLP), cites the	
current Laboratory Services Branch	
Laboratory Operations and Quality	
Assurance Manual and/or relevant CLP	
Statements of Work	
Referenced analytical SOPs are attached to	\Box Yes \Box No \Box Not Applicable –
the QAPP	using only Regional
	laboratory/CLP
QAPP Worksheet #24: Analytical	
Instrument Calibration	
Identifies all analytical instruments, whether	\Box Yes \Box No
used in the field or the laboratory -or- If	
using Regional laboratory/Contract	
Laboratory Program (CLP), cites the	
current Laboratory Services Branch	
Laboratory Operations and Quality	
Assurance Manual and/or relevant CLP	
Statements of Work	
For each instrument, identifies the calibration	□ Yes □ No
procedure and title/position responsible for	
corrective action; references and attaches the	
SOP or identifies the calibration range,	

frequency, and acceptance criteria, and			
corrective action in the table; calibration			
process should link the calibration to a			
specific instrument identification number -or-			
If using Regional laboratory/Contract			
Laboratory Program (CLP), cites the			
current Laboratory Services Branch			
Laboratory Operations and Quality Assurance Manual and/or relevant CLP			
Statements of Work			
QAPP Worksheet #25: Analytical			
Instrument and Equipment Maintenance,			
Testing, and Inspection			
For a laboratory with a quality system that	\Box Yes	\Box No	
conforms to ISO 17025:2005, the laboratory's			
quality manual may be referenced for this			
work sheet; otherwise or if project-specific			
modifications apply, lists each analytical			
instrument/equipment that requires			
maintenance, testing, and inspection			
activities, list those activities, and provides			
the frequency, acceptance criteria, corrective			
action, title/position responsible for corrective			
action, and reference for those activities -or-			
If using Regional laboratory/Contract			
Laboratory Program (CLP), cites the			
current Laboratory Services Branch			
Laboratory Operations and Quality			
Assurance Manual and/or relevant CLP			
Statements of Work			
QAPP Worksheet #26 & 27: Sample			
Handling, Custody, and Disposal			
Lists all activities from sample labeling	□ Yes	\Box No	
through sample disposal, indicating the			
organization and title/position responsible for			
each activity and the SOP reference -or- If			
using Regional laboratory/Contract			
Laboratory Program (CLP), lists all field			
activities and cites the current Laboratory			
Services Branch Laboratory Operations			
and Quality Assurance Manual and/or			
relevant CLP Statements of Work for all			
laboratory activities			
Referenced SOPs are attached to the QAPP -	□ Yes	□ No	

or- If using Regional laboratory/Contract		
Laboratory Program (CLP), attaches		
referenced SOPs for field activities and		
cites the current Laboratory Services		
Branch Laboratory Operations and		
Quality Assurance Manual and/or relevant		
CLP Statements of Work for all laboratory		
activities		
Example forms, sample labels, and chain-of-	□ Yes	\Box No
custody documentation are attached to the		
QAPP		
QAPP Worksheet #28: Analytical Quality		
Control and Corrective Action		
Provides a separate worksheet for each	\Box Yes	\Box No
analytical method/SOP, matrix, and analytical		
-or- If using Regional laboratory/Contract		
Laboratory Program (CLP), cites the		
current Laboratory Services Branch		
Laboratory Operations and Quality		
Assurance Manual and/or relevant CLP		
Statements of Work		
Identifies the type, number and frequency of	\Box Yes	\Box No
QC sample collection (field) or QC sample		
analysis procedure (laboratory) along with the		
required QC statistically derived limits/		
acceptance criteria for each analyte; includes		
corrective action and title/position responsible		
for corrective action-or- If using Regional		
laboratory/Contract Laboratory Program		
(CLP), provides this information for QC		
sample collection (field) and cites the		
current Laboratory Services Branch		
Laboratory Operations and Quality		
Assurance Manual and/or relevant CLP		
Statements of Work for this information		
for QC sample analysis procedure		
(laboratory)		
QAPP Worksheet #29: Project Documents		
and Records	- T Z	
Provides a comprehensive list of the	\Box Yes	□No
documents and records required for this		
-	\Box res	
project Describes the generation, verification, and storage location/archival of hard-copy and	□ Yes	□ No

electronic information produced during the		
course of the project for sample collection and		
field records		
Describes the generation, verification, and	□ Yes	\Box No
storage location/archival of hard-copy and		
electronic information produced during the		
course of the project for project assessments;		
attaches assessment checklists or other		
standardized forms to the QAPP		
Describes the generation, verification, and	\Box Yes	\Box No
storage location/archival of hard-copy and		
electronic information produced during the		
course of the project for laboratory records-		
or- If using Regional laboratory/Contract		
Laboratory Program (CLP), cites the		
current Laboratory Services Branch		
Laboratory Operations and Quality		
Assurance Manual and/or relevant CLP		
Statements of Work plus discusses		
SEMS/Federal Records and EQuIS	- 37	
Provides requirements for laboratory data	\Box Yes	□ No
deliverable contents consistent with the		
expected stages selected for data validation		
(see EPA 540-R-08-005) -or- If using		
Regional laboratory/Contract Laboratory Program (CLP), cites the current		
Laboratory Services Branch Laboratory		
Operations and Quality Assurance Manual		
and/or relevant CLP Statements of Work		
Describes data handling equipment and	□ Yes	
procedures used to process, compile and		
analyze data; provides a complete list of		
computer hardware and software needs;		
specifies requirements such as information		
security controls for ensuring quality of		
electronic information (utility, objectivity,		
and integrity) If using Regional		
laboratory/Contract Laboratory Program		
(CLP), includes discussion of the current		
Laboratory Services Branch Laboratory		
Operations and Quality Assurance Manual		
and/or relevant CLP Statements of Work		
plus relevant Region 4 Data Validation		
Standard Operating Procedures for		
Contract Laboratory Program data		

Provides electronic data deliverable	□ Yes	
requirements for analytical deliverables and		
field documentation according to the Region		
4 Format for EQuIS Data Processor (EDP);		
describes process for assuring that Region 4		
Format for EQuIS Data Processor (EDP)		
electronic data deliverables (EDDs) are		
provided to EPA Region 4 and identifies		
individual(s) responsible for EDD submittals		
QAPP Worksheet #31, 32, & 33:		
Assessments and Corrective Action		
Lists the required number, frequency and type	□ Yes	🗆 No
of assessments with approximate dates and		
title/position and organization of each		
individual(s) responsible for performing these		
assessments. Note: If using Regional		
laboratory and/or CLP, laboratory		
assessments will be conducted at the		
program level		
Discusses one or more of the following types	□ Yes	\Box No
of assessments: peer reviews, technical		
audits, surveillance, management system		
reviews, readiness reviews, quality system		
audits, performance evaluations, data quality		
assessments.		
Discusses the authority and independence of	□ Yes	\Box No
the individual(s) performing the assessments		
in relation to those being assessed.		
Discusses where assessment findings will be	□ Yes	□ No
documented and how the assessment findings		
will be communicated to all key project staff,		
state and EPA personnel responsible for the		
study oversight and the deliverable due dates.		
For each assessment listed, provides the	□ Yes	\Box No
title/position and organization of the		
individual(s) responsible for responding to		
assessment findings, assessment response		
documentation, and timeframe for response		
For each assessment listed, provides the	□ Yes	🗆 No
title/position and organization of the		
individual(s) responsible for implementing		
the corrective action and for monitoring		
corrective action implementation		
QAPP Worksheet #34: Data Verification		

and Validation Inputs		
Identifies the planning documents (such as	□ Yes	□No
QAPP, contract, field SOPs, laboratory		
SOPs), field records, and laboratory records		
that will be used during data verification and		
validation; indicates whether each item will		
be used for verification (completeness),		
validation (conformance to specifications), or		
both If using Regional laboratory/Contract		
Laboratory Program (CLP), includes		
discussion of the current Laboratory		
Services Branch Laboratory Operations		
and Quality Assurance Manual and/or		
relevant CLP Statements of Work plus		
relevant Region 4 Data Validation		
Standard Operating Procedures for		
Contract Laboratory Program data		
QAPP Worksheet #35: Data Verification		
Procedures		
Data verification is a completeness check to	\Box Yes	\Box No
confirm that all required activities were		
conducted, all specific records are present,		
and the contents of the records are complete.		
Documents procedures that will be used to		
verify project data. For each field record,		
references the document containing the		
requirements, process description, and		
responsible person/organization.		
For each laboratory record, references the	\Box Yes	\Box No
document containing the requirements,		
process description, and responsible		
person/organization. If using Regional		
laboratory/Contract Laboratory Program		
(CLP), includes discussion of the current		
Laboratory Services Branch Laboratory		
Operations and Quality Assurance Manual		
and/or relevant CLP Statements of Work		
plus relevant Region 4 Data Validation		
Standard Operating Procedures for		
Contract Laboratory Program data		
For each audit and corrective action record,	\Box Yes	□ No
references the document containing the		
requirements, process description, and		
responsible person and organization.		

QAPP Worksheet #36: Data Validation		
Procedures		
Documents procedures that will be used to	\Box Yes	\Box No
validate project data. Data validation is an		
analyte and sample-specific process for		
evaluating compliance with contract		
requirements, methods/SOPs, and		
measurement performance criteria.		
Procedures should be summarized in the		
worksheet, including specific SOP references,		
if applicable. If using Regional		
laboratory/Contract Laboratory Program		
(CLP), includes discussion of the current		
Laboratory Services Branch Laboratory		
Operations and Quality Assurance Manual		
and/or relevant Region 4 Data Validation		
Standard Operating Procedures for		
Contract Laboratory Program data		
Referenced data validation SOPs are attached	□ Yes	🗆 No
to the QAPP, if applicableor- If using		
Regional laboratory/Contract Laboratory		
Program (CLP), cites the current		
Laboratory Services Branch Laboratory		
Operations and Quality Assurance Manual		
and/or relevant Region 4 Data Validation		
Standard Operating Procedures for		
Contract Laboratory Program data		
Validation procedures define validation stage	□ Yes	□No
code and define any data qualifiers to be		
applied by the data validator – or – If using		
Regional laboratory, external validation is		
Not Applicable; if using CLP, cites the		
relevant Region 4 Data Validation		
Standard Operating Procedures		
Validation procedures include checklists to be	□ Yes	□ No
used by the data validatoror- If using		
Regional laboratory/Contract Laboratory		
Program (CLP), cites the current		
Laboratory Services Branch Laboratory		
Operations and Quality Assurance Manual		
and/or relevant Region 4 Data Validation		
Standard Operating Procedures for		
Contract Laboratory Program data		
QAPP Worksheet #37: Data Usability		

Assessment		
Identifies the individual(s) responsible for	□ Yes	\square No
reconciling the data to the project-specific		
requirements		
Describes data usability assessment process	\Box Yes	\Box No
including statistics, equations, and computer		
algorithms to be used to analyze the data and		
reconcile it to project-specific requirements		
Discusses how limitations in the final data set	\Box Yes	\Box No
will be documented and communicated to all		
end data users and stakeholders		
Describes the circumstances under which data	\Box Yes	\square No
would be rejected and removed from the final		
data set and addresses resolution of potential		
data gaps		
Describes the data usability assessment	\Box Yes	\Box No
process to confirm that the useable data are		
adequate to make the site decision		

<u>Final QAPP Disposition</u>:

_____ *Approved, no comments*

Signature of Designated Approval Official (DAO)

Signature of Section Chief of the DAO _____

Not Approved, Address Comments, Submit Revised QAPP to the EPA Designated Approval Official