Fort Ord OU CT Bio Pilot Study Data Validation Reports LDC# 11183

Volatiles



Laboratory Data Consultants, Inc. Data Validation Report

Project/Site Name:

Fort Ord OU CT Bio Pilot Study

Collection Date:

October 31, 2003

LDC Report Date:

December 2, 2003

Matrix:

Water

Parameters:

Volatiles

Validation Level:

EPA Level III

Laboratory:

Sequoia Analytical

Sample Delivery Group (SDG): P311018

Sample Identification

0344MOCT240F 0344MOCT262F 0344MOCT241F 0344MOCT264F 0344MOCT242F 0344MOCT265F 0344MOCT243F 0344MOCT266F 0344MOCT244F 0344MOCT267F 0344MOCT245F 0344MOCT268F 0344MOCT246F 0344MOCT269F 0344MOCT247F 0344MOCT272F 0344MOCT248F 0344MOCT273F 0344MOCT249F

0344MOCT249F 0344MOCT250F 0344MOCT251F 0344MOCT253F 0344MOCT255F 0344MOCT256F 0344MOCT257F 0344MOCT258F 0344MOCT258F 0344MOCT260F

0344MOCT261F

Introduction

This data review covers 29 water samples listed on the cover sheet including dilutions and reanalysis as applicable. The analyses were per EPA SW 846 Method 8260B for Volatiles.

The review follows a the USACE Environmental Data Quality Management Program Specifications, USACE Sacramento District (Version 1.08) and a modified outline of the USEPA Contract Laboratory Program National Functional Guidelines for Organic Data Review (October 1999) as there are no current guidelines for the methods stated above.

A table summarizing all data qualification is provided at the end of this report. Flags are classified as P (protocol) or A (advisory) to indicate whether the flag is due to a laboratory deviation from a specified protocol or is of technical advisory nature.

Blank results are summarized in Section V.

Field duplicates are summarized in Section XVI.

Raw data were not reviewed for this SDG. The review was based on QC data.

The following are definitions of the data qualifiers:

- J+ Data are qualified as estimated, with a high bias likely to occur. False positives or false negatives are unlikely to have been reported.
- J- Data are qualified as estimated, with a low bias likely to occur. False positives or false negatives are unlikely to have been reported.
- J Data are qualified as estimated; it is not possible to assess the direction of the potential bias. False positives or false negatives are unlikely to have been reported.
- P Data are qualified as rejected. There is a significant potential for the reporting of false negatives or false positives.
- U Data are qualified as non-detected, because the analyte was observed in an associated laboratory or field blank.
- A Indicates the finding is based upon technical validation criteria.
- P Indicates the finding is related to a protocol/contractual deviation.

None Indicates the data was not significantly impacted by the finding, therefore qualification was not required.

I. Technical Holding Times

All technical holding time requirements were met.

The chain-of-custodies were reviewed for documentation of cooler temperatures. All cooler temperatures met validation criteria.

II. GC/MS Instrument Performance Check

Instrument performance was checked at 12 hour intervals.

All ion abundance requirements were met.

III. Initial Calibration

Initial calibration was performed using required standard concentrations.

Percent relative standard deviations (%RSD) were less than or equal to 15.0% for each individual compound and less than or equal to 30.0% for calibration check compounds (CCCs).

In the case where %RSD was greater than 15.0%, the laboratory used a calibration curve to evaluate the compound. All coefficients of determination (r^2) were greater than or equal to 0.990.

For the purposes of technical evaluation, all compounds were evaluated against the 30.0% (%RSD) National Functional Guideline criteria. Unless noted above, all compounds were within the validation criteria.

Average relative response factors (RRF) for all volatile target compounds and system performance check compounds (SPCCs) were within method and validation criteria.

IV. Continuing Calibration

Continuing calibration was performed at the required frequencies.

Percent differences (%D) between the initial calibration RRF and the continuing calibration RRF were within the method criteria of less than or equal to 20.0% for calibration check compounds (CCCs).

For the purposes of technical evaluation, all compounds were evaluated against the 25.0% (%D) National Functional Guideline criteria. Unless noted above, all compounds were within the validation criteria.

The initial calibration verification (ICV) percent differences (%D) were less within the QC limits.

All of the continuing calibration RRF values were within method and validation criteria.

V. Blanks

Method blanks were reviewed for each matrix as applicable. No volatile contaminants were found in the method blanks.

VI. Surrogate Spikes

Surrogates were added to all samples and blanks as required by the method. All surrogate recoveries (%R) were within QC limits.

VII. Matrix Spike/Matrix Spike Duplicates

The laboratory has indicated that there were no matrix spike (MS) and matrix spike duplicate (MSD) analyses specified for the samples in this SDG, and therefore matrix spike and matrix spike duplicate analyses were not performed for this SDG.

VIII. Laboratory Control Samples (LCS)

Laboratory control samples were reviewed for each matrix as applicable. Percent recoveries (%R) and relative percent differences (RPD) were within QC limits.

IX. Regional Quality Assurance and Quality Control

Not applicable.

X. Internal Standards

All internal standard areas and retention times were within QC limits.

XI. Target Compound Identifications

Raw data were not reviewed for this SDG.

XII. Compound Quantitation and CRQLs

Raw data were not reviewed for this SDG.

XIII. Tentatively Identified Compounds (TICs)

Raw data were not reviewed for this SDG.

XIV. System Performance

Raw data were not reviewed for this SDG.

XV. Overall Assessment of Data

Data flags have been summarized at the end of the report.

XVI. Field Duplicates

No field duplicates were identified in this SDG.

XVII. Field Blanks

No field blanks were identified in this SDG.

Fort Ord OU CT Bio Pilot Study Volatiles - Data Qualification Summary - SDG P311018

No Sample Data Qualified in this SDG

Fort Ord OU CT Bio Pilot Study Volatiles - Laboratory Blank Data Qualification Summary - SDG P311018

No Sample Data Qualified in this SDG

LDC #:	11183B1	VALIDATION COMPLETENESS WORKSHEET
SDG #:	P311018	Level III
Laborator	ry: Sequoia Analytical	

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Reviewer:	
2nd Reviewer:	20

METHOD: GC/MS Volatiles (EPA SW 846 Method 8260B)

The samples listed below were reviewed for each of the following validation areas. Validation findings are noted in attached validation findings worksheets.

	Validation Area		Comments
I.	Technical holding times	4	Sampling dates: 10/31/03
II.	GC/MS Instrument performance check	N	
III.	Initial calibration	\forall	70 RSD. Y2
IV.	Continuing calibration	A	70 D& 1CV
V.	Blanks	A	
VI.	Surrogate spikes	A	
VII.	Matrix spike/Matrix spike duplicates	N	client selfied
VIII.	Laboratory control samples	4	20=10
IX.	Regional Quality Assurance and Quality Control	N	
X.	Internal standards	N	
XI.	Target compound identification	N	
XII.	Compound quantitation/CRQLs	N	
XIII.	Tentatively identified compounds (TICs)	N	
XIV.	System performance	N	
XV.	Overall assessment of data	A	
XVI.	Field duplicates	N	
XVII.	Field blanks	N	

Note:

A = Acceptable

N = Not provided/applicable SW = See worksheet Samples: MA + D

ND = No compounds detected

R = Rinsate

FB = Field blank

D = Duplicate

TB = Trip blank

EB = Equipment blank

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Ţ	0344MOCT241F	12 0344MOCT251F	22 0344MOCT264F	32 35 0290 - BK
l	0344MOCT242F	13 O344MOCT253F	23 2 0344MOCT265F	33
1	0344MOCT243F	14 0344MOCT254F	24 ² 0344MOCT266F	34
١	0344MOCT244F	15 0344MOCT255F	25 0344MOCT267F	35
1	0344MOCT245F	16 0344MOCT256F	26 → 0344MOCT268F	36
1	0344MOCT246F	17 0344MOCT257F	27 0344MOCT269F	37
1	0344MOCT247F	18 0344MOCT258F	28 0344MOCT272F	38
1	0344MOCT248F	19 0344MOCT260F	29 0344MOCT273F	39
07	0344MOCT249F	20 ² 0344MOCT261F	30	40