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Analysis
(Type of Report)

REPORT NO.: MSL-5615
ESC-UAG-86-32

JOB/PROJECT NO.: 6216

DATE: April 22, 1986

TITLE: ANALYSIS OF TOTAL PCB CONGENERS IN KRUMMRICH SOIL
SAMPLES

AUTHORS: B.M. Hughes, D. McKenzie, and C. Trang

ABSTRACT: Krummrich Plant soil samples were analyzed for total
PCB congener concentrations using capillary GC/EC
analysis techniques. All samples contained total
PCB congeners at levels less than 50 µg/g.

TECHNICAL APPROVAL:

Paul L. Sherman
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Senior Research Group Leader

APPROVED BY:

William M. Kees
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Environmental Sciences Center
Quality Assurance Specialist

APPROVAL DATE:

April 25, 1986

MONS 023974

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REPT. NO.: ESC-UAG-86-32
AUTHORS: B.M. Hughes, D. McKenzie, and C. Trang
TITLE: ANALYSIS OF TOTAL PCB CONGENERS IN KRUMMRICH SOIL SAMPLES
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ANALYSIS REPORT

DATE: April 22, 1986

LOG NUMBER: U-86-04-08-01

PROJECT NO: 760.28-6216

TO: Jim Gloeckner/Krummrich Plant

FROM:

Environmental Sciences Center
Monsanto Company
800 N. Lindbergh Blvd.
St. Louis, Mo. 63167

Phone No: (314) 694-1464

Title: Analysis of Total PCB Congeners in Krummrich Soil Samples.

ABSTRACT

Krummrich Plant soil samples were analyzed for total PCB congener concentrations using capillary GC/EC analysis techniques. All samples contained total PCB congeners at levels less than 50 ug/g.

INTRODUCTION

Ten soil samples were analyzed for total PCB congener concentrations. These analyses are required as a part of the Krummrich Plant cleanup of an area where chemicals had previously been stored.

SUMMARY

Table 1 summarizes the results of capillary GC/EC analyses of 10 soil extracts. Note that all samples contained a total PCB concentration less than 50 ug/g. Table 1 also summarizes quality control experiments conducted on one blank soil sample. Table 2 summarizes results from quality control native spiking studies for blank soil and sample 1B-3. See PCX21 and PCX03 for the details of how the samples were extracted and cleaned up prior to GC/EC analysis. Refer to Standard Analysis Method ECA02 for the details of how the cleaned up extracts were analyzed. This information is included in a later section of this report.

The patterns of PCB congeners found in these soil samples were similar to the Aroclor 1262 pattern. However they were not identical. In a previous project for the Krummrich Plant (Project Number 760.28-6204), similar results were obtained for soil samples which were analyzed using GC/EC and GC/MS extract analysis methods. In that study, the PCB congener patterns were also not identical to Aroclor 1262. Therefore we have chosen to report data based upon this less expensive, yet technically equivalent, GC/EC analysis method for this present study.

(continued)

MONS 023976

QC RESULTS

Quality control results are shown in Tables 1 and 2. These tables show good recoveries of surrogate compounds which were added to each sample before extraction. This information indicates how well each soil sample was extracted. Duplicate analyses of IB-2 and IB-3 are also shown in Table 1. Table 2 summarizes recoveries of Aroclor 1254 which was added to Blank Soil samples, and IB-3. These data show that good PCB congener recoveries would be expected using these methods for the analysis of soils from the Krumerich Plant.

DETAILS

Standard Sampling Method No(s): Samples supplied by the Krumerich Plant.

Standard Extraction Method No(s): PCEX03 and PCEX21 (See PCFS03 and PCFS21 for flow sheets describing these methods.)

Standard Analysis Method No(s): ECAN02

Final Report File No(s): A21618

Table File No(s): B21618, C21618

Also included in this report are:

- 1) Summaries of Standard Extraction and Analysis Methods used for this project.
- 2) Chain-of-custody information.
- 3) Analysis Request Information recorded at sample login.
- 4) A copy of the ESC/UAC Good Laboratory Practices Manual.

ANALYST(S): B. Mason Hughes, David McKenzie and Chi Trang

ANALYST APPROVAL: B. Mason Hughes

TABLE 1. Summary of PCB Concentrations in Soil Samples Determined using PCEX21 and PCEX03 Extraction Methods and ECAN02 Analytical Method.

ESC Project No: 760.28 - 6216

Date: April 21, 1986

Sample I.D.	Surrogate Recoveries(%Recovery)		PCB(a)
	CL5-benzene 2,4,6-CL3-biphenyl		Concentration (ug/g)
			Total

Method Blank	34	56	ND(b)
IB-1	49	73	"
IB-2	58	65	8.9
" (DUP)	95	82	10.6
IB-3	80	86	3.1
" (DUP)	-	61	3.1
IB-4	49	68	ND(b)
IB-5	49	71	"
IB-6	72	103	"
IB-7	52	74	8.1
IB-8	72	97	ND(b)
OB-30	84	96	"
OB-32	60	97	"

Footnotes:

- a. Aroclor 1254 and Aroclor 1262 standard formulations were used to calibrate the capillary GC/EC response for PCB congeners. However the pattern of congeners in these soil samples were not identical to these formulations. See text for more details.
- b. ND- None Detected (<1.0 ug/g)

TABLE FILE NO: B21618:MH

MONS 023978

TABLE 2. Summary of Aroclor 1254 Spiking Concentrations and Recoveries in Quality Control Experiments for Soil Samples.

ESC Project No: 760.28 - 6216

Date: April 21, 1986

Sample I.D.	Surrogate Recoveries (%Recovery)		Aroclor 1254	
	CL3-benzene	2,4,6-CL3-Biphenyl	Amount Added (ug/g)	Recovery (%)
Soil Blank(L. Spike)	34	48	1.00	110
" (H. Spike)	44	84	10.00	105
IB-3 (L. Spike)	22	77	1.00	109
" (H. Spike)	99	84	10.00	79

TABLE FILE NO: C20718:MM

MONS 023979

The following pages describe Standard Sampling, Extraction, and/or Analysis Methods used for this project.

MONS 023980

ESC Standard Extraction Method (SEM)
TITLE: PCBs in Non-EC Active Matrices

ESC Standard Extraction Method No:
PCEX21

MATRIX(X),(CES):

TYPICAL SAMPLE SIZE (g)	TYPICAL EXTRACT VOLUME (ml)
-------------------------------	-----------------------------------

Soils and other non-EC active
solids and solvents

0.5 - 5.0

10

TYPICAL ANALYTES: Aroclors 1232, 1242,
1248, 1254, 1260, 1262, trichloro-
biphenyl isomers through octachloro-
biphenyl isomers

ESC Standard Analysis Method Nos:
ECAN01, ECAN02

INTERFERENCES: Any electron capturing
species in the retention time region
of trichlorobiphenyl isomers through
octachlorobiphenyl isomers that is
non acid hydrolyzable

TYPICAL SURROGATES

TYPICAL RECOVERIES (%)

pentachlorobenzene
2,4,6-trichloro- biphenyl

50 - 120

50 - 120

APPARATUS AND MATERIALS:

Burdick and Jackson benzene or hexane for sample extraction or dilution.
Pierce vials.
Sulfuric acid at varying concentrations.
Ultrasonic bath.
Disposable glassware.

(See flow sheet for this extraction method in PCFS21.)

LIMITATIONS: Method applies only to matrices which do not contain interfering
electron capturing species in the trichlorobiphenyl through octachloro-
biphenyl isomer retention time region.

SAFETY PRECAUTIONS: See PCB handling protocols in laboratory safety manual.

DATES OF METHOD CREATION AND UPDATES: 12/17/84

MONS 023981

FLOW SHEET PCFS21
FOR ESC EXTRACTION METHOD PCX21

SAMPLE ID: MRC EXTRACT #

..... LOG NO:

EXTRACTION DATE and TIME:

SAMPLE COLOR AND DESCRIPTION

WEIGH OUT SAMPLE INTO
A 14 OR 40 mL PIERCE []
VIAL

SAMPLE WEIGHT-----

SURROGATE
SPIKE OF SAMPLE []

ADD 10.0 mL BENZENE
TO SAMPLE. []

ULTRASONICATE SAMPLE []
FOR 1/2 HOUR IN A
SOUND BATH.

PLACE 2 mL OF THE
EXTRACT INTO A 7 mL []
VIAL

ADD 2 mL []
SULFURIC ACID AND
SHAKE

ALLOW PHASES TO
SEPARATE REMOVE 1 mL []
FOR ANALYSIS.

10.0 mL
FINAL VOLUME

EXTRACT NO:

DATE/TIME COMPLETED

ANALYST:

ESC JOB NO:

MONS 023982

ESC Standard Extraction Method (SEM)
TITLE: PCBs in Non-EC Active Matrices

ESC Standard Extraction Method No:
PCEX03

MATRIX(X), (GES):

TYPICAL SAMPLE SIZE (g)	TYPICAL EXTRACT VOLUME (ml)
-------------------------------	-----------------------------------

Soils and other non-EC active
solids, solvents and oils.

0.5 - 5.0	10
-----------	----

TYPICAL ANALYTES: Aroclora 1232, 1242,
1248, 1254, 1260, 1262, trichloro-
biphenyl isomers through octachloro-
biphenyl isomers

ESC Standard Analysis Method Nos:
ECAN01, ECAN02

INTERFERENCES: Any electron capturing
species in the retention time region
of trichlorobiphenyl isomers through
octachlorobiphenyl isomers which are
not removed by the three-phase silica-
gel column.

TYPICAL SURROGATES	TYPICAL RECOVERIES (%)
pentachlorobenzene	50 - 120
2,4,6-trichloro- biphenyl	50 - 120

APPARATUS AND MATERIALS:

Burdick and Jackson hexane for sample extraction or dilution.

Sodium sulfate for drying the sample matrix.

Silica gel columns impregnated with potassium permanganate, sulfuric acid,
and potassium hydroxide.

Nitrogen for extract volume reduction.

Disposable glassware.

(See flow sheet for this extraction method in PCFS03.)

LIMITATIONS: Method applies only to matrices which do not contain interfering
electron capturing species in the trichlorobiphenyl through octachloro-
biphenyl isomer retention time region when electron capture analytical methods
are used. This limitation does not apply to mass spectrometric methods.

SAFETY PRECAUTIONS: See PCB handling protocols in laboratory safety manual.

DATES OF METHOD CREATION AND UPDATES: 2/13/84, 9/25/84

MONS 023983

FLOW SHEET PCFS03
FOR ESC EXTRACTION METHOD PCER03

SAMPLE ID: MRC EXTRACT #

..... LOG NO:

EXTRACTION DATE and TIME:

WEIGH OUT SAMPLE INTO
A 14 mL VIAL []

FINAL WEIGHT

INITIAL WEIGHT

SAMPLE WEIGHT

SURROGATE
SPIKE OF SAMPLE []

.....
.....

DRY SAMPLE IF NEEDED BY
ADDING ABOUT 0.1 g []
Na SO TO VIAL
2 4 []

RINSE COLUMN WITH 2 X 20 mL
HEXANE

DILUTE SAMPLE IN VIAL []
WITH HEXANE AND PLACE
ONTO CLEAN-UP COLUMN.

CLEAN-UP
COLUMN DESCRIPTION
25 mL DISPO PIPET

QUANTITATIVELY RINSE
VIAL ONTO COLUMN WITH []
4 X 1 mL RINSES.

ELUTE SAMPLE INTO 50 mL
CENTRIFUGE TUBE WITH []
40 mL HEXANE.

NITROGEN BLOW DOWN TO
..... mL []

	1 cm Na SO 2 4
	4 cm 1% POTASSIUM PERMANGANATE SILICA GEL
	1 cm SILICA GEL
	4 cm ACID SILICA GEL
	1 cm SILICA GEL
	4 cm BASE SILICA GEL
	GLASS WOOL

EXTRACT NO:

DATE/TIME COMPLETED

ANALYST:

ESC JOB NO:

MONS 023984

ESC Standard Analysis Method (SAM)
TITLE: Aroclor Formulations in Solvents

ESC Standard Analysis Method No:
EGAN02

ANALYTICAL INSTRUMENT AND MODE OF OPERATION: HP-5880 Capillary GC/EC interfaced to an HP-3357-LAS Laboratory Data System

TYPICAL ANALYTE	TYPICAL INSTRUMENT LEVEL OF DETECTION (LOD) (ug/ml)	TYPICAL PRECISION @ 10X LOD (%)	TYPICAL ACCURACY @ 10X LOD (%)
Aroclor formulations 1232, 1242, 1248, 1254, 1260, and 1262	0.1	30	30

INTERFERENCES AND LIMITATIONS: Any electron capturing species in the retention time region of trichlorobiphenyl through octachlorobiphenyl isomers are potential interferences. Exact retention times of most likely PCB isomers can be obtained from the analysis of Aroclor 1248, Aroclor 1254 or Aroclor 1260 formulations.

CHROMATOGRAPHIC COLUMN: 30-meter fused silica J&W DB-5 (0.25 μ m) with wide bore (0.32mm).

COLUMN TEMPERATURE PROGRAM: 130(1)/4/300

INJECTION TYPE: Splitless

CALIBRATION AND STANDARDIZATION: Use internal standard quantitation technique and calibrate instrument response factors from authentic Aroclor formulation standards. Use a minimum of 10 PCB isomers for the determination of formulation identity and quantity.

CALCULATIONS: Formulation concentration =
(isomer RRF)(isomer area/internal standard area)

where: RRF is the relative response factor for the appropriate PCB isomer in a given formulation. Formulation concentration is obtained from the average of at least 10 such calculations.

SAFETY PRECAUTIONS: See PCB handling protocols in laboratory safety manual.

DATE OF METHOD CREATION: 10/25/83, 2/3/84 Time Stamp - <860422.0936>

MONS 023985

The following pages show chain-of-custody information for this project.

MONS 023986

Monsanto

MONSANTO INDUSTRIAL CHEMICALS CO.
Sequel, Illinois 62201
Phone: (618) 271-6835

CHAIN OF CUSTODY

SAMPLING RECORD

Sampling Identification:	3-31-86 IB-1 0-12'	4-2-86 IB-6 78'-17 1/2'
Log Number:	4-1-86 IB-2 0-13 1/2'	4-2-86 IB-7 0-14 1/2'
	4-1-86 IB-3 0-17 1/2'	4-3-86 IB-1 78'-28'
Date:	4-7-86	4-1-86 IB-4 7 1/2'-19 1/2'
	4-2-86 IB-5 10'-19'	4-3-86 IB-30 22'-11 1/2'
Analyses Required:	PCB	4-3-86 IB-32 26'-22 1/2'

Sample Taken By: PAUL HARGRELL

Date:

LABORATORY RECORD

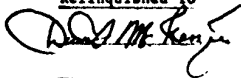
Received By: RON McENTIRE

Date: 4-4-86

Relinquished To

Date

Comments



4-8-86

REPORT OF ANALYSES

Date

Report No.

Author

4/22/86

1151-5615

B. Mann Hargrell

INSTRUCTIONS

The person taking the sample completes the information in the Sampling Record area.

Each person to whom the sample is relinquished should sign the form and forward as necessary. Analyses performed should be listed with comments.

When all analyses are complete, the original should return to the "Received By" signature.

MONS 023987

The following pages describe Analytical Request Information generated for this project.

MONS 023988

Time Stamp - <860422.0920>

MONSANTO COMPANY/ENVIRONMENTAL SCIENCES CENTER

ANALYSIS REQUEST

DATE: 4-8-86

LOG NO: U-86-04-08-01

REQUESTER: Krummrich

PROJECT NO: 6216

REPORT RESULTS TO: FILES/MEES, Hughes, Sherman, McKenzie, Trong

ANALYSIS REQUESTS SENT TO: FILES, Meese, Sherman, Hughes, McKenzie

SAMPLE PICKUP BY: McKenzie

SAMPLE LOCATION: U-420

SAMPLE DESCRIPTION FOR 10 SAMPLES

SAMPLE IDENTIFICATION	NUMBER OF BOTTLES	ESC NUMBER(S)
Blank soil (METHOD BLANK)		PCB-1582
Blank soil + LS		PCB-1583-LS
Blank soil + HS		PCB-1584-HS
IB-1, 3-31-86, 0-12'	1 x 4 oz	PCB-1585
IB-2, 4-1-86, 0-13 1/2'	1 x 4 oz	PCB-1586, PCB-1587-DUP
		PCB-1599-NS
IB-3, 4-1-86, 0-17 1/2'	1 x 4 oz	PCB-1588, PCB-1589-DUP
		PCB-1590-LS, PCB-1591-HS
		PCB-1600
IB-4, 4-1-86, 7 1/2-19 1/2'	1 x 4 oz	PCB-1592
IB-5, 4-2-86, 10-19'	1 x 4 oz	PCB-1593
IB-6, 4-2-86, 7 1/2-17 1/2'	1 x 4 oz	PCB-1594
IB-7, 4-2-86, 0-14 1/2'	1 x 4 oz	PCB-1595
IB-8, 4-3-86, 7 1/2-20'	1 x 4 oz	PCB-1596
OB-30, 4-3-86, 7-14 1/2'	1 x 4 oz	PCB-1597
OB-32, 4-3-86, 20-22 1/2'	1 x 4 oz	PCB-1598

Note- DUP refers to a duplicate analysis, LS refers to a low level native spike, HS refers to a high level native spike, and NS refers to no simulated spike, which was done to determine if any of the surrogate compounds were in the samples since this was suggested by low recovery of the pentachlorobenzene surrogate. It was determined that the samples did contain measurable quantities of both pentachlorobenzene (surrogate spiking compound) and hexachlorobenzene (internal standard), so an external standard method was used for quantitation.

RECORDED BY: McKenzie

REQUIRED ANALYSES: PCBs

ANALYTICAL TECHNIQUES:

TECHNIQUE (EN)	DATE COMPLETED	TECHNIQUE(EN)	DATE COMPLETED
x GC/EC (18)	4/22/86		
QUALITATIVE: x	SEMIQUANTITATIVE:	QUANTITATIVE: x	

MONS 023989

REQUIRED QA/QC: SEE GOOD LABORATORY PRACTICES MANUAL

COMPLETION DATE REQUESTED:

COMMENTS: Verbal results reported to Gloeckner on April 21.

SAMPLE DISPOSITION: HOLD FOR 30 DAYS

SAFETY PRECAUTIONS: SEE SAFE LABORATORY PRACTICES MANUAL

ANALYSIS REQUEST FILE NAME: AA216:LI

MONS 023990

ANALYSIS REQUEST

DATE

LOG NO.

MONSANTO COMPANY
DAYTON LABORATORY

REQUESTER**

CHARGE NO.**

REPORT RESULTS TO:
FILES/MEES

SAMPLE DESCRIPTION

NUMBER OF BOTTLES:

NUMBER OF SAMPLES **

ANALYSIS REQUEST NO.
FILES

SAMPLES FOR:

SAMPLE LOCATION

RECORDED BY:

REQUIRED ANALYSIS**

PCB

KNOWN HEALTH HAZARD PRESENTED BY SAMPLE: **

ANALYTICAL TECHNIQUE **

- | | |
|---|--|
| ☐ GAS CHROMATOGRAPHY | ☐ NMR |
| ☐ ULTRAVIOLET | ☐ HPLC |
| ☐ VISIBLE | ☐ XRD |
| ☐ INFRARED | ☐ ICP |
| ☐ EMISSION | ☐ ATOMIC ABSORPTION |
| ☐ EDAX XRF | ☐ MASS SPEC |
| ☐ MICROSCOPY | ☐ GC/MS |
| ☐ OTHER _____ | ☐ AUTO ANALYZER |
| QUAL _____ | SEMIQUANT. _____ |
| QUANT. _____ | |

REFERENCE TO PREVIOUS WORK (REQUEST #, JOURNAL, ETC.) **

COMPLETION DATE

TIME ALLOTTED FOR ANALYSES

ANALYST

HOURS

INST. TIME

REQUIRED QA/QC

**SAMPLE DISPOSAL:
AFTER REPORT COMPLETE☐ DISCARD☐ RETURNHOLD FOR ☐ 30 DAYS☐ 60 DAYS

OTHER

** THIS INFORMATION MUST BE SUPPLIED BY ANALYSIS REQUEST ORIGINATOR

MONS 023991

The following is a copy of the ESC/UAG Good Laboratory Practices
Manual.

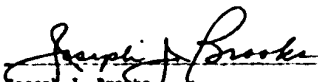
MONS 023992

GOOD LABORATORY PRACTICES MANUAL

A Guide to Quality Control/Quality Assurance

Ultratrace Analysis Group

March 1984


Joseph J. Brooks
Group Leader, Ultratrace Analysis


H. A. Woltermann
Manager, Environmental Sciences Center

Issued to: _____

Date: _____

Environmental Sciences Center
Dayton Laboratory
Dayton, Ohio 45407

JR3/JOB A

MONS 023993

TABLE OF CONTENTS

	<u>Page</u>
I Objectives	1
II Scope	1
III Analysis	1
A. Methods	1
B. QA/QC Program	2
1. Sample Preparation QA/QC	2
a. Method Blanks	2
b. Replicates	2
c. Spikes	2
2. Instrumental Analysis QA/QC	3
a. Instrumentation	3
b. Blanks	4
c. Quantitation Internal Standard	4
d. Standards	4
C. Degrees of Quantitation	4
1. Qualitative Analysis	7
a. Tentative	7
b. Confirmed	7
2. Semi-Quantitative Analysis	7
3. Quantitative Analysis	7
D. Quality Criteria and Remedial Action	8
1. Quality Criteria	8
a. Blanks	8
b. Recoveries	8
c. Precision	8
d. Limit of Detection	9
e. Accuracy	9
2. Remedial Action	9
IV Data	9
V Sample Disposal	10

FIGURES

<u>Figure</u>	<u>Page</u>
1 Example of GC/MS Logbook page	5
2 Example of GC/EC Logbook page	6
3 Example of Disc/Tape Logbook page	11

I. Objectives

The main objectives of the Ultratrace Analysis Group quality assurance/quality control program are: 1) to assure that our laboratory generates high quality results; and 2) to maintain the necessary records that document laboratory performance.

II. Scope

The scope of this manual includes areas of GLP not specifically addressed in the ESC section manual and expands upon certain areas where QA/QC practices differ from those described in the section manual. It is intended to be used along with the section manual to provide a complete picture of the GLP within the Ultratrace Analysis Group.

III. Analysis

The generation of analytical results within the Ultratrace Analysis Group routinely involves two processes: sample preparation and instrumental analysis. Sample preparation involves one or more of a variety of techniques including extraction, dilution, acid/base washes, column chromatography, concentration and numerous other techniques to render the sample in a suitable condition for instrumental analysis. Instrumental analysis takes the suitably prepared sample and introduces it into an analytical instrument to measure the desired parameter(s). Both of these processes can be described by clearly defined and separable methods called sample preparation methods (SPMs) and instrumental analysis methods (IAMS). SPMs and IAMS may be combined in numerous ways (depending upon the nature of the samples and the needs of the customer) to produce a successful analytical protocol. For example, several different SPMs may be used to produce samples that are suitable for analysis by a single IAM. The measures incorporated into each of these experimental processes to ensure that an assessment of the quality of the results can be made constitute the QA/QC program for a project. The following subsections describe the elements of the analysis process:

A. Methods

Methods are the written documentation that describe the analytical process. They contain sufficient detail such that a competent scientist or technician can readily perform the required analyses. The usual practice is to have methods that describe the sample preparation process (SPMs) and methods that describe the instrumental analysis process (IAMS). Each method developed within the Ultratrace Analysis Groups is assigned a method number as described in the ESC Section Manual.

Methods are obtained from various sources including in-house development, modification/adaptation of literature methods, or specific protocol methods dictated by government agencies. Methods may be formally validated by a series of experiments that describe a classical validation protocol or may be less vigorously proven owing to factors such as frequency or use, customer needs and

cost/timing restrictions. In any case a QA/QC program is applied to the analysis process to provide a measure of the quality of the results. Unless a QA/QC program is prescribed by a required protocol (e.g., EPA Dioxin in Soil protocol), certain minimum QA/QC measures (depending upon the analysis type) are incorporated into all Ultratrace Analysis projects. The following subsection describes the Ultratrace QA/QC program:

B. QA/QC Program

An effective QA/QC program incorporates experiments that provide information relative to each of the following questions:

- (1) Are there any impurities, background or contamination that will influence the results?
- (2) How reproducible are the results?
- (3) How well can the analyte be recovered from the sample matrix?
- (4) What is the limit of detection for the analysis?
- (5) What is the dynamic range of the analysis?
- (6) What is the accuracy of the analysis?

All of these questions are addressed by properly selecting/applying the use of blanks, replicates, spikes and standards in a QA/QC program. The discussion of the use of these QA/QC techniques as they apply to Ultratrace Analysis Group activities is best done by considering the sample preparation and instrumental analysis processes separately.

1. Sample Preparation QA/QC

- a. Method blanks - Representative glassware, solvents and reagents are used with sample matrices which are similar to those for which results are being reported, yet do not contain the analytes of interest. Method blanks are analyzed at a frequency equal to 10% of the analyzed samples for sample sets which are multiples of 10. At least one method blank is analyzed for sample sets containing between 1 and 9 samples.
- b. Replicates - If sufficient sample amounts have been submitted for analysis, replicate frequency is the same as the method blank frequency. If at all possible, replicates should be obtained from the same sample bottle in order to minimize sample inhomogeneity problems due to samples being taken at slightly different times.
- c. Spikes - If sufficient sample is available for each sample type, the sample is split and spiked at two levels with

the native (unlabeled) analyte(s). The levels are typically 2x (low) and 20x (high) the anticipated detection limit and serve as a technique for estimating the limit of detection for each matrix. These low and high spikes are conducted at a frequency equal to the Method Blank frequency. Obviously spiking with the native analyte cannot be done for widescan analysis since this presupposes knowledge of the identity of the analyte(s) in the sample.

Other spiking techniques are used to evaluate the extraction/recovery efficiencies for the analyte(s) of interest. Ideally extraction/recovery efficiencies are determined for each sample, since, in principle, each sample may represent a different matrix. This can be done very easily when the detection technique is mass spectrometry if an isotopically labeled version of the analyte(s) is available. Documentation of extraction/recovery efficiencies is not required for each sample if the sample matrix for a set of samples is constant. However, for a sample set for which the similarity of samples is not known, and for which adequate sample is available, each sample must be split and spiked with a known amount of the analyte(s) of interest to determine extraction efficiency. If adequate sample size is not available or the identity of the analyte(s) is not known before the fact (i.e., widescan analysis), surrogate spiking compounds may be used to assess extraction efficiencies in each sample without actually spiking each sample with the analyte(s) of interest. The choice of the surrogate spiking compound is tailored for each analyte. This surrogate must be chosen so that it is either chemically identical to the analyte of interest (for example using isotopically labeled analogues of the analytes), or is closely similar in terms of liquid/liquid partition properties, chemical reaction properties, column elution properties or other matrix isolation properties which are used to isolate the analyte(s) from the sample matrix. Another property of the surrogate spiking compound is that it not be present in the sample being analyzed.

2. Instrumental Analysis QA/QC

The final step in obtaining analytical results involves the use of an appropriate instrumental technique to produce the analytical data. Several QA/QC practices must be incorporated in this process to ensure validity of the data.

a. Instrumentation

Careful documentation of instrument usage, calibration, and maintenance is essential for the generation of high quality results. Each major instrument within the Ultratrace Analysis Group will be the assigned responsibility of a professional employee designated as the Instrument Steward. The Instrument Steward is responsible for making sure that logbooks are available and properly completed by instrument users. It is the

responsibility of each individual using the equipment to record all the pertinent data required in the logbook. An example of a page from a GC/MS logbook is given in Figure 1 and that of a GC/EC logbook is given in Figure 2.

Instruments will be calibrated according to the instrument manufacturers' specifications or according to the method being used with the instrument. A log of instrument calibration will be maintained with the instrument.

The Instrument Steward will be responsible for ensuring that preventative and corrective maintenance programs are carried out on all instruments under his/her responsibility. If a specific problem is noted by an instrument user, it should be noted in the logbook and the Instrument Steward notified so that appropriate corrective action can be taken.

b. Blanks

A solvent blank will be analyzed at a frequency equal to the method blank. The method blank can serve as the solvent blank provided it is clean in the area of the analyte(s) of interest. This practice eliminates the possibility of background instrumental contamination.

c. Quantitation Internal Standard

A quantitation internal standard will be added to each sample following sample preparation but prior to instrumental analysis. This compound will serve for determining relative response factors and for correcting for variations in sample injection volumes and evaporation of solvents.

d. Standards

Standards are solutions of the analyte(s) of interest used to determine the response of the detector over the range of analyte concentrations found in the samples. For Quantitative analyses, sufficient standards and replication of standards will be run to demonstrate linearity and precision of the detector response over the range of analyte concentrations. Should sample concentrations fall outside the demonstrated linear range, either additional standards are prepared to expand the range, or appropriate dilution/concentration of the sample is done to place it in the proper range.

C. Degrees of Quantitation

The information provided for any particular analysis is tailored to the needs of the customer. In actual practice these needs vary across the broad continuum from the grossly qualitative to the precisely quantitative. It is informative to examine what is involved in providing the varying degrees of quantitation required by our customers.

HP 5985A

No. 11004

[illegible]

FIGURE 1. Example of GC/MS Logbook page.

MONS 023999

[illegible]

FIGURE 2. Example of CC/EC Logbook page.

1. Qualitative Analysis

Qualitative analysis implies no quantitation at all but rather seeks only the identity of the compound(s) present. GC/MS is particularly applicable to this type of analysis problem. Two types of identifications can be made:

a. Tentative

A tentative identification is made on the basis of the analyst's judgement of a match between the mass spectrum from the sample and a mass spectrum from a reference library. This is often done for the large number of compounds found in a videscan organic analysis.

b. Confirmed

A confirmed identification requires that the chromatographic retention time and the mass spectrum of the unknown compound match those of a known standard of the tentatively identified compound. This is often done in videscan analyses when standards can be obtained for tentatively identified compounds that are of concern.

The normal QA/QC requirements for qualitative analysis are (1) analysis of a blank, (2) analysis of duplicate samples and (3) use of a calibration compound (usually Decafluorotriphenylphosphine, DFIPP) to assure that the mass spectrometer will produce mass spectra that can be compared with library spectra.

2. Semi-Quantitative Analysis

Semi-Quantitative Analysis extends the information provided by a qualitative analysis to include some estimate of the amount of analyte(s) present. The method of semi-quantitation involves the comparison of detector response of the analyte(s) with that of an internal standard. The estimated concentration is obtained by assuming a unit response factor between the analyte(s) and the internal standard(s). An indication of recovery from the sample matrix is obtained from the recovery of the surrogate internal standard(s).

3. Quantitative Analysis

Quantitative Analysis provides for the accurate assessment of the amount of analyte(s) present in a sample. To be able to do quantitative analysis one must know the identity of the analyte so that standards can be prepared for spiking and for the generation of calibration curves. The method of quantitation is based on a three-point calibration curve using standards of the analyte(s) of interest over the range of concentrations in the samples. A one-point calibration method can be used (as with the laboratory data

system) provided that linearity has been demonstrated over the concentration range of the samples using standards of the analyte(s) at three concentrations in the range. This calibration curve is then used to provide concentrations of the analyte(s) in the samples via an external standards method or more preferably via an internal standards method. For the internal standards method, the quantitation internal standard is added to each of the standards (just as with the samples) prior to instrumental analysis and relative response factors are determined.

For quantitative analysis the mass spectrometer is operated as a gas chromatographic detector. The MS is therefore tuned and operated in a manner specified in the approved method rather than using a calibration compound as required for qualitative analyses.

D. Quality Criteria and Remedial Action

The inclusion of QA/QC measures into an analysis program has little meaning unless the results are interpreted and appropriate action is taken when necessary. The ability to do this assumes some set of criteria that will trigger a response or remedial action. The following subsections describe the criteria that are used in the Ultratrace Analysis Group to evaluate when quality control data indicate a need for corrective measures and what remedial action will be taken.

1. Quality Criteria

The overriding quality criterion above all others is: Do the results meet the needs of the customer. This determination will be made jointly between the customer and the analyst. The analyst must exercise caution in not allowing the customer to overstate, misinterpret or unintentionally misrepresent the data. In addition to this prime quality criterion, the following conditions must be met:

- a. Blanks - In the event that a blank produces a positive value, it may not be used as a valid blank if it exceeds 10% of any reported value for a sample.
- b. Recoveries - Recoveries for all surrogate and native spikes must lie in the range of 50%<Recovery<150%.
- c. Precision - Precision will be calculated based on the recoveries of surrogate internal standards. Average recoveries will be calculated with $\pm$ standard deviation for each sample set. Replicate analyses and low and high native spike recoveries may also be used to provide additional information/insight into the precision of the analysis.

MONS 024002

- d. Limit of Detection - The limit of detection (LOD) will normally be stated as $\frac{1}{2}$ the value of the low level native spike if it is recovered and quantifiable. Should the low level native spike not be recovered, the LOD will be stated as $\frac{1}{2}$ the value of the high level native spike if it is recovered and quantifiable. Should neither spike be recovered, a decision must be made regarding the advisability of reprocessing the sample. Should all samples contain such large quantities of the analyte(s) that the levels of the low and high spikes are insignificant ($<10\%$) by comparison, the value of the LOD becomes less important and is simply stated as the LOD for the instrumental analysis process determined by the analysis of standards and the ability to recover the surrogate spikes.
- e. Accuracy - The accuracy of the analyses is best determined by analyzing some sample which contains the analyte at some predetermined level (e.g., an NBS standardized sample). If this type of standard is not available then the low level and high level spikes can be used as a measure of the accuracy of the analyses. Results must fall within a range of 75% to 120% of the expected value for the analyses to be considered valid.

If the above quality criteria are not met the analyst will immediately assess the need for and extent of remedial action that may be necessary.

2. Remedial Action

The first course of action will be for the analyst and/or group leader to contact the customer and discuss the results pointing out the limitations of the data indicated by the quality control results. At this point the overriding quality criterion of customer needs will be addressed first. It could be that the customer's needs are met without meeting one or more of the other criteria. In such a case no remedial action is necessary. It could also be that the customer has such a demand for tightness in the results that his needs are not met even though the other quality criteria are. This would trigger further analyses with tighter quality control designed around the results obtained (e.g., replications of positives to better establish precision or spikes of non-detected samples to better establish the limit of detection).

In consultation with the customer additional experiments will be designed including the appropriate quality control to ultimately provide the results of sufficient quality to meet the customer's needs.

IV. Data

This section deals with the subject of data and how they are obtained, recorded and reduced so as to be useable in a report and yet readily available for referral and inspection.

The samples will be labeled with a log number at sample login according to the procedure described in the ESC Section GLP Manual. When delivered to the responsible technical person, the samples may then receive additional labeling as deemed necessary. This additional labeling will be documented in a laboratory logbook and/or notebook. These logbooks and notebooks will be used by the analysts to record all pertinent work and results obtained from the samples. The laboratory notebooks will be filled out in accordance with Monsanto Uniform Notebook Procedures.

All raw data generated for a given project, which cannot be conveniently placed in the laboratory notebook (e.g., chromatograms, computer output, etc.), will be filed under some prescribed order (i.e., log number) in a central file under the supervision of the Project Leader and/or QA Specialist.

Most of the data generated by the Ultratrace Analysis Group are computer data stored on magnetic discs. In order to archive these data the specific file name and disc number associated with the analysis will be written down in the laboratory notebook and/or instrument logbook. When the disc is completely full of data an entire unedited disc image or copy of the files on the disc will be transferred to magnetic tape. A logbook will be kept for recording the disc number, name and prefix associated with it, along with the associated tape and file number to which the disc was transferred. An example of such a logbook is given in Figure 3. When the data are transferred a verification program is used to check all transferred records for correctness. When tapes are filled (they typically hold seven to ten disc images) they should be stored in a fireproof vault.

All calculations of the results of the data will be specified in the method used. These calculations will be entered in the analysts laboratory notebook unless they are computer generated. In this case, a copy of the computer output will be filed with the raw data. The Project Leader will be responsible for checking the correctness of the data.

V. Sample Disposal

Samples will be disposed of in accordance with the instructions on the Analysis Request form. The only exception to this would result when the analysis reveals a problem which requires sample disposal in a manner other than that originally specified. If the Project Leader notes from the results of the analysis that a potentially toxic product was present in the samples then the samples will be disposed of in a manner deemed safe by the Safety Department. The date of sample disposal and the manner should be noted on the request form stored with the raw data completing the chain of custody of the sample.

MONS 024004

CAN	(Multi-Drive Designation)	Person Opened	Date Opened	Person Taped	Date Taped	Tape # (File #)
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130	(AR0)	BMT	1/1/83	BMT	3/14/83	GD69(1)
131	(AR4)(AR)	BMT	4/6/83			GD69(5)
132	(AR2)		3/1/83			GD69(6)
133	(AR3)					GD69(7)
134	(AR4)					GD69(8)
250	(2E) ISD Program	Fin	3/1/83	BMT	3/1/83	GD69(8)
135	(AR05) [AR]	BMT	3/1/83	BMT	3/14/83	GD69(9)
136	(AR06) [AR]	BMT	3/1/83	LSK	5/12/83	GD73(1)
137	(AR07) [AR]	SM	3-11-83	LSK		GD73(2)
138	(AR08) [AR]	SM	3-11-83	LSK		GD73(3)
1	(AR00) [AR]			BMT	3/14/83	GD69(10)
250	(AR11) [AR] - R	Fin	3/1/83	BMT	3/15/83	GD71(1)
113	(AR13) [AR]	BMT	4/1/83			GD71(5)
114	(AR14) [AR]	"				GD71(4)
139	(AR09) [AR]	SM	3/1/83	LSK	5/1/83	GD73(4)
140	(AR10) [AR]	SM	3/1/83	LSK	5/1/83	GD73(5)
141	(AR01) [AR]	BMT	3/1/83	SB	5/27/83	GD73(6)
142	(AR02)			SB	5/1/83	GD73(7)
143	(AR03)			SB	5/1/83	GD73(8)
144	(AR04)			SB	5/27/83	GD73(9)
145	(AR05)			SB	5/27/83	GD73(10)
146	(AR06)			SB	6/1/83	GD72(2)
147	(AR07)			SB	6/27/83	GD72(3)
148	(AR08)			SB	6/1/83	GD72(4)
149	(AR09)			SB	6/27/83	GD72(5)
150	(AR10)			SB	6/27/83	GD72(6)
250	(2E) ISD Program	Fin	3/1/83	LSK	4-2-83	GD72(7)
151	(AR01) [AR]	BMT	5/27/83	LSK	7/1/83	GD72(7)
152	(AR02)		5/27/83	LSK		(1)
153	(AR03)		"	LSK		(7)

FIGURE 3. Example of DISC/Tape Logbook page.

MONS 024005