# The Analysis of Chlorinated Dioxins, Difurans in Pet Food



## Introduction

Screening for Chlorinated Dioxins and Difurans in human supplies is a wellestablished practice. Both the EU and US have extensive protocols for the testing of food for human consumption as well as additives to feed for commercial livestock. With recent contamination issues in pet food like the 2007 melamine incident, there is a growing concern for pet safety.

With increasing efforts to screen both finished pet foods and pet food additives, the development of reliable analytical practices is of high priority. For Persistent Organic Pollutants, the high lipid content of canned and dried pet foods make them an ideal source of dietary contamination for household pets.

Using the FMS Total Prep solution (TRP) for the extraction and extract clean-up of pet food, labs can deliver extracts for same day analysis. The following outlines the procedures for processing canned dog food for Chlorinated Dioxin and Difuran analysis via EPA 1613.

### Instrumentation

- FMS, Inc. PLE®
- FMS, Inc. SuperVap® Concentrator
- FMS, Inc. SuperVap® Vial Concentrator
- FMS, Inc. 200ml concentrator tubes
  FMS, Inc. 40ml extraction cells w/disposable end caps
- FMS, Inc. Power Prep®
- Thermo Trace Ultra GC with DFS HRMS
- Restek Dioxin 2, 60m GC column

### Consumables

• FMS, Inc. High Capacity ABN Silica columns

- FMS, Inc. Basic Alumina columns
- FMS, Inc. Carbon columns
- Fisher Optima\* Toluene
- Fisher Optima\* n-Hexane
- Fisher Optima\* Methylene Chloride
- Acros n-Nonane
- Fisher dodecane

• Agilent Hrdromatrix (diatomaceous earth)

- · Restek Ottawa Sand
- Cambridge Isotopes EDF-9999, EPA
- 1613 calibration Standards
- Cambridge Isotopes EDF-8999, EPA
- 1613 Labeled Surrogate
- Cambridge Isotopes EDF-5999, EPA
- 1613 Recovery Standard
- Cambridge Isotopes EDF-6999, Labeled Clean-up Standard



	Run #1		Run #2		Run #3		MB	
Analyte	<u>13C</u> <u>Rec</u>	<u>Native</u>	<u>13C</u> <u>Rec</u>	<u>Native</u>	<u>13C</u> <u>Rec</u>	<u>Native</u>	<u>13C</u> <u>Rec</u>	<u>Native</u>
2,3,7,8-TCDF	82	<.0178	104	0.0456	78	0.0075	90	ND
2,3,7,8-TCDD 1,2,3,7,8-	84	0.0113	105	0.1394	77	0.0331	92	ND
PeCDF 2,3,4,7,8-	67	<.0291	79	0.037	65	0.0119	74	ND
PeCDF 1,2,3,7,8-	69	0.0162	84	0.0317	70	<.0088	78	ND
PeCDD 1,2,3,4,7,8-	72	0.0373	85	0.7676	72	<.0174	79	ND
HxCDF 1,2,3,6,7,8-	84	0.0192	112	<.0216	89	0.0148	96	ND
HxCDF 2,3,4,6,7,8-	80	0.0148	108	0.0244	83	0.0108	91	ND
HxCDF 1,2,3,7,8,9-	82	0.0174	105	0.0161	85	0.0116	94	ND
HxCDF 1,2,3,4,7,8-	90	0.0088	114	0.0262	91	0.0068	99	ND
HxCDD 1,2,3,6,7,8-	84	0.033	109	0.9383	87	<.0089	93	ND
HxCDD 1,2,3,7,8,9-	80	0.0246	105	1.3878	85	<.0188	92	ND
HxCDD 1,2,3,4,6,7,8-		0.0191		0.9943		0.0119		ND
HpCDF 1,2,3,4,7,8,9-	67	<.0754	85	<.0754	76	0.0701	79	ND
HpCDF 1,2,3,4,6,7,8-	76	ND	97	<.0142	85	0.0114	90	ND
HpCDD	73	0.2057	90	12.2956	81	0.0775	85	ND
OCDD	66	1.2475	75	115.7943	78	0.2291	77	0.2309
OCDF		0.0355		0.0393		<.0556		ND

Analytical results, native compounds reported in pg/g



Figure #1; examples of various canned dog food matrices.

# Procedure

# Sample Prep

- 3 brands of canned dog food were obtained.
- Aliquots of 10 grams were spiked with 13C labeled surrogate standards
- Samples were mixed with diatomaceous earth till all moisture was absorbed
- Dried samples are transferred to extraction cells.
- Remaining cell volume topped off with Ottawa Sand.

# PLE

- 1. Extraction cells loaded onto PLE
- 2. Cells filled sequentially with 50% DCM in Hexane
- 3. Extraction cells pressurized to 1500PSI
- 4. Cells heated to 120 degrees and held for 20 minutes
- 5. Cells cooled to approximately ambient temperature
- 6. Cells flushed with extraction solvent
- 7. Solvent purged from cells direct to Super Vap with Nitrogen

# SuperVap

- 1. Preheat temp: 20 minutes at 60 °C
- 2. Evap mode w/Sensor temp: 60 °C
- 3. Nitrogen Pressure: 10 PSI
- 4. Evaporator tubes rinsed with hexane several times to ensure no sample sticks to glass walls

# PowerPrep

- 1. Columns are conditioned
- 2. Samples are loaded onto PowerPrep
- 3. Samples are loaded across ABN silica columns in n-Hexane and eluted onto Alumina columns
- 4. Alumina columns are eluted onto carbon columns using Methylene Chloride.
- 5. Carbon column back eluted with Toluene and collected in SuperVap as Sample Fraction



# **Procedure Continued**

#### **SuperVap**

- 1. Preheat temp: 20 minutes at 60 °C
- 2. Evap mode w/Sensor temp: 60 °C
- 3. Nitrogen Pressure: 10 PSI

After extracts reach sensor termination point, extracts transferred to GC vials. 5 μl dodecane added to vials and vials transferred to SuperVap vial Concentrator

#### SuperVap Vial Concentrator

- 1. Temp: 25 °C
- 2. Nitrogen Pressure: 1.5 PSI
- 3. Vials evaporated till volume of dodecane reached.
- 5 μl recovery standard added to extract and vials transferred to Trace GC for analysis

### Conclusions

Review of the 3 pet food matrices shows labeled recoveries ranging 65-109%, well within EPA1613 limits. Analysis of the Method Blank showed no background levels above the CS .1 calibration standard level except OCDD. Combining the clean background with good recoveries demonstrates the TRP process' ability to handle wet pet food of various types.

With a total process time of ~5 hours from start to finish, the TRP process enables sample turnaround for same day analysis. When factoring in the need to qualify batches for release and product delays, the value of rapid testing becomes critical.



The Total Rapid Prep is expandable from 1 to 6 Modules and runs in parallel.

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