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| Project: | ANIMIDA III | | | | | | |
| Parameters: | PAH and Biomarkers | | | | | | |
| Laboratory: | Battelle, Norwell, MA | | | | | | |
| Matrix: | Tissue | | | | | | |
| Data Set: | DP-14-0585 | | | | | | |
| Analytical SOP: | 5-157 | | | | | | |
| Method Reference: | Modified EPA Method 8270D | | | | | | |
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| Sample Custody | Receipt Date | | | Temp (°C) | | | |
| 8/14/2014 | | | 4.0 | | | |
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| Corrective Actions | None. | | | | | | |
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| Sample Storage | The samples were stored in an access-limited freezer until sample preparation could begin. | | | | | | |
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|  | METHOD SUMMARIES |  | | |  |  | |
| Sample Preparation | Tissue samples were homogenized with titanium blades and split for metals analysis at Sequim and FIT.    The tissue samples were extracted following a modified EPA Method 3510C. Samples were prepared for analysis by weighing approximately 20 grams of sample material into a pre-cleaned extraction vessel and dried using sodium sulfate.  Each sample was spiked with PAH, Biomarker and SHC surrogates and extracted 3 times using methylene chloride by tissuemizer.  The combined extracts were dried over sodium sulfate and concentrated by Kuderna-Danish (KD) and nitrogen evaporation techniques. Sample clean-up was performed on the extracts using alumina columns. Extracts were further cleaned up and fractionated using silica gel columns. The F1 fraction was collected and split for TPH/SHC and biomarker analyses. The F2 fraction was collected for PAH and alkyated PAH analysis. The extracts were concentrated and spiked with IS for analysis. | | | | | | |
| Prep comments | Sample M5883 was noted to contain water after the post column cleanup. Sodium sulfate was added and the prep continued with the rest of the batch.  Also, the GC/MS fraction went dry. 250uL of hexane was added to the vial before re-combining for the FID dilutions.  M5901 had a low sample amount. Dry weight was not performed on this sample. | | | | | | |
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| Analysis | PAH, alkylated PAH (F2 fraction) and Biomarkers (F1 fraction) were measured by gas chromatography-mass spectrometry (GC/MS) in the selected ion mode (SIM). An initial calibration consisting of target analytes was analyzed prior to analysis to demonstrate the linear range of analysis. Calibration verification was performed every 24 hours in which samples were analyzed. Concentrations of target compounds were calculated versus internal standards. Target PAH were quantified using the average response factors (RF) generated from the initial calibration. The alkyl homologue PAH series were assigned the RF of the parent PAH. Biomarkers used RFs from the single individual biomarkers within the calibration standard curve. All reported data (except NSC) is corrected based on surrogate recoveries.  All data is reported on dry weight basis except the SRM (wet weight) and NSC and CO (oil weight). | | | | | | |
| Analysis comments | None. | | | | | |  |
| Holding Times | Extraction Date(s) |  | Analysis Date(s) | | | |  |
|  | 10/8/2014 & 10/15/2014 | 11/4-6, 12-15/2014 | | | | | |

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| Procedural Blank (PB) | A PB was prepared with this analytical batch to ensure the sample extraction and analysis methods are free of contamination. |
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| PB <5 X MDL  Samples must be >5x PB | Four exceedences noted. |
| Comments: There were four exceedences (three Naphthalene and one Phenanthrene) for analytes detected in samples at less than five times the blank concentration. Reanalysis of the PB on another instrument confirmed results. |
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| Laboratory Control Spike (LCS) | A LCS was prepared with this analytical batch. The percent recoveries of target analytes were calculated to measure accuracy. |
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| Recovery of 70-130% | Eighteen exceedences noted. |
| Comments: There were 18 exceedences for LCS recovery. Results were verified by reanalysis on another instrument. It was determined that the LCS standard vial used to spike this batch had a low volume remaining. The next batch spiked with a different vial passed all MQO criteria. No further action was taken. |
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| Surrogate Recovery | Surrogate compounds were added prior to extraction. The surrogate recoveries are calculated to measure extraction efficiency. |
| Recovery of 40-120% | Two exceedences noted. |
| Comments: Surrogate recoveries for 5B(H)-Cholane were high in the NSC and in the CO, at 130% and 126% respectively. Reanalysis on another instrument confirmed the results. Due to this anomaly, the 5B(H)-Cholane standard vial that was used to spike this batch was  inspected, and was determined to have a low volume remaining. Recoveries of all other surrogates, for which a different spiking  standard is utilized, were acceptable. The NSC and CO are not surrogate corrected, so there was no impact on data quality. |
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| Sample Duplicate (QADUP) | A QADUP was prepared with this analytical batch. The RPD of target analytes were calculated to measure data quality in terms of accuracy. |  |
| Relative Percent Difference (RPD) < 30% | No exceedences noted. |  |
| Comments: None. |  |
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| Standard Reference Material (SRM) | An SRM was prepared with this batch to assess accuracy of the analytical procedures. |
| < 30 PD from target concentration and the 95% confidence level analyte concentration must be > 5x the MDL. Concentration must be certified and >5x the MDL for MQO to apply | One exceedence noted. |
| Comments: Benzo(a)anthracene was recovered low in the SRM. Results were confirmed by reanalysis on another instrument. Prep records and integrations were verified. Recoveries for this analyte were acceptable in all CCVs for this batch. No further action was taken. |
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| North Slope Crude (NSC) | A NSC Reference Oil was prepared with this batch to evaluate the instrumental accuracy and also provide petroleum pattern information, aiding in the qualitative identification of target analytes.  The control oil also run in this batch has no associated target values and is not evaluated. |
| < 30% RPD for 90% of analytes | No exceedences noted. |
| Comments: None. |
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| Initial Calibration (ICAL) | The GC/MS is calibrated with a minimum 5 level curve for all compounds. |
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| Individual RSD ≤25%; Mean RSD ≤15% | No exceedences noted. |
| Comments: None. |
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| Independent Calibration Check (ICC) | The independent check was run after each initial calibration to verify the calibration. This standard is from a different source than the ICAL. |
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| Individual and Mean PD <25% | No exceedences noted. |
| Comments: None. |
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| Continuing Calibration Verification (CCV) | Continuing calibration standards were run every 24 hours to ensure that initial calibration is still valid. |
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| Individual RSD ≤25%; Mean RSD ≤15% | Two exceedences noted. |
| Comments: Indeno(1,2,3-cd)pyrene and Dibenz(a,h)anthracene were recovered low in CCV B2478, with %Ds of -43.8 and -28.9 respectively. These analytes were not detected in the samples bracketed by this CCV. The samples were reanalyzed outside of holding time with acceptable CCVs on another instrument to confirm the absence of these analytes. The in-hold analyses are reported with documentation. |
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