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APPENDIX

ศูนย์วิทยทรัพยากร จหาลงกรณ์มหาวิทยาลัย

APPENDIX A

PREPARATION OF REAGENTS

Nitric Acid (HNO3) Conc.

AR grade nitric acid (${\rm HNO_3}$) was purified by sub-boiling distillation in a teflon distillation apparatus and it was stored in a precleaned polyethylene bottle.

This purified nitric acid was used to prepare 4N nitric acid by diluting 250 ml of purified nitric acid to 1000 ml by double distilled water. This 4N nitric acid was used in the analysis of heavy metals in sea water.

10 % nitric acid was used for cleaning all apparatus used in this study.

Ammonium Pyrrolidine Dithiocarbamate (APDC) Solution.

2% (w/v) aqueous solution of APDC was prepared by dissolving 2 g. of APDC in 100 ml of double distilled water. This solution was purified by shaking with methyl isobutyl ketone (MIBK) prior to use in order to make it free of metal impurities. Purification was made by extracting each solution 3 times with MIBK. At the first extraction, APDC solution was extracted with MIBK in a ratio 10:1 and 20:1 at the second and third extractions. Each extraction was performed by shaking for 5 minute in a separatory funnel and allowing for each phase to fully separate and then the lower layer was withdrawn. Discard the upper layer with was MIBK. The

APDC solution was freshly prepared each time prior use for solvent extraction.

Methyl Isobutyle Ketone (MIBK).

The MIBK was purified by redistillation at 115-116 °C in all glass apparatus.

Ammonium Hydroxide Solution (NH OH).

NH₄OH was purified by placing two 250 ml teflon beakers in which the first beaker contained concentrated ammonium hydroxide solution (AR grade) and another beaker contained double distilled water to about half of full in a vacuum desiccator. The pressure inside the desiccator was reduced by a vacuum pump and let stand for at least 24 hour. The purified ammonium hydroxide solution was obtained from the beaker originally contained double distilled water. This purified solution was a dilution of ammonium hydroxide solution.

Stock Standard Solution of Copper, Lead, Iron and Zinc.

The stock standard solutions (1000 ppm) of Cu, Pb, Fe and Zn were commercially available and were used to prepare a primary standard solution (100 ppm)

1. Primary Standard Solution (100 ppm).

Primary standard solution was prepared by diluting 500 μ l of stock standard solution to 5 ml with 4N HNO3, and then stored in the precleaned polyethylene vials and kept in a refrigerator.

2. Secondary Standard Solution (10 ppm).

Secondary standard solution was prepared by diluting 500 μ l of primary standard solution to 5 ml with 4N HNO3, and then stored in the precleaned polyethylene vials and kept in a refrigerator.

3. Third Standard Solution (1000 µg/1).

Third standard solution was prepared by diluting 500 $\,\mu$ l of secondary standard solution to 5 ml with 4N HNO3, and then stored in the precleaned polyethylene vials and kept in a refrigerator.

4. Working Standards Solution (Cu, Pb, Fe and Zn).

4.1 Copper.

The copper working standard solutions were 0, 1, 5, 10, 20 and 40 μ g/l which were prepared by diluting 0, 10, 50, 100, 200 and 400 μ l of third standard solution to 10 ml with 4N HNO₃.

4.2 <u>Lead</u>.

The lead working standard solutions were 0, 1, 3, 5, 10, 15 and 20 μ g/l which were prepared by diluting of 0, 10, 30, 50, 100, 150 and 200 μ l of third standard solution to 10 ml with 4N HNO₃.

4.3 Iron.

The iron working standard solutions were 0, 100, 500, 1000, 2000, 4000 and 8000 μ g/l which were prepared by diluting

of 0, 100, 500, 1000, 2000, 4000 and 8000 μl of secondary standard solution to 10 ml with 4N HNO3.

4.4 Zinc.

The zinc working standard solutions were 0, 50, 100, 200, 400, 1000 and 1500 μ g/l which were prepared by diluting of 0, 50, 100, 200, 400, 1000 and 1500 μ l of secondary standard solution to 10 ml with 4N HNO₃.

TOC Standard Solution Preparation.

0.85 g of potassium hydrogen phthalate was dissolved in distilled water and diluted to 1000 ml. The standard solution was equivalent to 0.4 mgC/ml = 400 ppmC.

IC Standard Solution Preparation.

1.40 g of sodium hydrogen carbonate and 1.77 g of sodium carbonate (dried at 500-600 °C for 30 min and then allowed to cooled in a desiccator) was dissolved in distilled water and diluted to 1000 ml. The standard solution was equivalent to 0.4 mgC/ml = 400 ppmC.

Calibration Curves Preparation of TOC and IC.

The working standard solution of 0, 10, 20, 30 and 40 ppm standard solution were dilutions of 0, 2.5, 5.0, 7.5 and 10.0 ml of TOC and IC standard solution stock (400 ppmC) to 100 ml with distilled water.

APPENDIX B

STATISTICS ANALYSIS

Table B-1 Analysis of Variance, Completely Randomized (CRD)

Source of Variation	Sum of Squares	Degree of Freedom	Mean Square	Variance Ratio
Among group	SS _{among} = $\sum_{j=1}^{k} n_j (X_{.j} - X_{})^2$ $= \sum_{j=1}^{k} \frac{T^2_{.j}}{n_j} - \frac{T^2_{}}{N}$	k - 1	MS _{among} = SS _{among} (k-1)	V.R. = MS _{among} MS _{within}
Within group	SS _{within} = $\sum_{j=1}^{k} \sum_{i=1}^{\Sigma} (X_{i,j} - X_{.,j})^2$ = $\sum_{j=1}^{k} \sum_{i=1}^{n_j} x^2_{i,j} - \sum_{i=1}^{k} \frac{(T_{.,j})^2}{n_j}$	- N - k	MS _{within} = SS _{within} (N - k)	
Total	$ss_{total} = \sum_{\substack{j=1 \ j=1 \ i=1}}^{k} \sum_{j=1}^{n_j} (x_{ij} - x_{})^2$ $= \sum_{\substack{j=1 \ j=1 \ i=1}}^{k} \sum_{j=1}^{n_j} x_{ij}^2 - \frac{x_{}}{N}$	N - 1	ากร ทยา	ลัย

When
$$x_{ij} = \text{Observed data}$$
 $i = 1,2,3,...,n$
 $j = 1,2,3,...,k$
 $T_{.j} = \sum_{i=1}^{n_j} x_{ij} = \text{sum of column } j$
 $X_{.j} = \frac{T_{.j}}{n_j} = \text{mean of column } j$

$$T.. = \begin{cases} k \\ j=1 \end{cases} T.j$$

$$= \begin{cases} k \\ E \end{cases} \qquad \begin{cases} n_j \\ j=1 \end{cases} \qquad x_{ij} = sum \text{ of total}$$

$$x = \frac{T..}{N}, \qquad N = \begin{cases} k \\ E \end{cases} \qquad n_j$$

where,

SS_{among} = the among groups sum of squares

SS_{within} = the within groups sum of squares

.SStotal = the total sum of squares

MS_{among} = among group mean square

MSwithin = within group mean square

V.R. = the variance ratio

Decision

The variance ratio (V.R.) value was compared with the critical value F listed in a percentile of F-distributions table at degrees of freedom (k-1) and (N-k). If the V.R. was greater than F-value, the null hypothesis H_O was then rejected and could be concluded that the mean of samples were significantly different. If the V.R. was smaller than F-value, the null hypothesis H_O was accepted and could be concluded that the mean of samples were not significantly different.

APPENDIX C



Oil in the fuel tank and engine room is transferred to storage tank for sale.



Remaining oil which cannaot be removed is eliminated by burning.



Pieces of steel directly falls into the sea and weathering.



Some parts of the vessel presented on the sand beach.



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Sompongchaiyakul, P., M. Hungspreugs and S. Lim, "Baseline Values of Petroleum Hydrocarbon in the Upper Gulf of Thailand and in the Eastern Seaboard," Paper presented at the Third Seminar on Marine Science arranged by the National Research Council of Thailand.