

# Soil Extraction Procedure for Dioxin

### **Extraction Procedure**

- 1. Weigh 5 gm of soil into a 30 mL HDPE bottle.
- 2. Add 6 gm of anhydrous sodium sulfate and mix until sample is free flowing.
- 3. Add 1 steel mixing ball.
- 4. Add 20 mL of 20% acetone in hexane. Shake for 1 hour.
- 5. Allow to settle for 1 hour or centrifuge for 5 min. at 3000g.
- 6. Transfer extract to a 40 mL screw cap vial.

#### **Sulfuric Acid Treatment**

- 1. Add 8 mL of concentrated sulfuric acid to vial containing extract.
- 2. Shake by hand for 2 min. (Be sure to release pressure before shaking.)
- 3. Allow phases to separate and transfer the extract (top layer) to a clean 40 mLvial.
- 4. Transfer 5 mL of the extract to a 7 mL test tube and evaporate to 1 mL withnitrogen. This 1 mL will be applied to the multilayer silica column.

## Multilayer Silica Column Clean Up

Multilayer Silica Column Preparation

- 1. Activate Silica Gel Wash 60-100 mesh silica gel with methylene chloride. Allow the methylene chloride to evaporate under a fume hood with frequent stirring of the silica gel. When all of the methylene chloride has evaporated, bake the silica gel at 180°C for at least 1 hour. Cool in a desiccator. Store ina pre-cleaned glass jar at room temperature in a desiccator.
- 2. Prepare 44% Sulfuric Acid Silica Gel In a clean glass jar; add 4.28 mL of sulfuric acid to 10 gm of activated silica gel. Stir and shake until a uniform mixture without any lumps is obtained. Cap and store in a desiccator at roomtemperature for at least 24 hours prior to use.
- 3. Prepare Basic (2%NaOH) Silica Gel In a clean glass jar, add 4.85 mL of 1N sodium hydroxide to 10 gm of activated silica gel. Stir and shake until a uniform mixture without any lumps is obtained. Cap and store in a desiccatorat room temperature for at least 24 hours prior to use.
- 4. Prepare Multilayer Silica Column Obtain a 20 mL filtration column. Placea frit in the bottom. Add the following layers: 0.5 gm activated silica, 1 gmbasic silica, 0.5 gm activated silica, 2 gm sulfuric acid silica gel, 0.5 gm activated silica, 1 gm sodium sulfate. Place a frit over the top layer (sodium sulfate). Store in desiccator at room temperature until use.

### Extract Clean Up Using Multilayer Silica Column

- 1. Pre-elute multilayer silica column with 30 mL of hexane.
- 2. Add the 1 mL of extract from the sulfuric acid treatment step to the column. Rinse the test tube with 2-1 mL portions of hexane and apply to the column. Discard the solvent.
- 3. Elute the column with 30 mL of hexane and collect in a 30 mL or larger testtube.
- 4. Evaporate with nitrogen to 4 mL or less.
- 5. Transfer to a 7 mL test tube. Rinse the 30 mL test tube with 3-1 mL portions of hexane and add to the 7 mL tube.
- 6. Evaporate with nitrogen to 0.5 mL or less (do not completely evaporate todryness). Add 100 µL of DMSO with 0.02% Triton X-100 (DMSOT).
- 7. Evaporate <u>all</u> of the hexane so only DMSOT is left. It is very important to that no hexane is left. It will interfere with the ELISA.

- 8. Add 400  $\mu$ L of DMSOT. Add 500  $\mu$ L of deionized water to bring to 1 mL. Transfer to a 4 mL vial with a Teflon lined lid.
- 9. Dilute 1:10 with 50%DMSO in deionized water with 0.01% Triton X-100.Analyze in ELISA.

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