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Texas Commission on Environmental Quality

CHECKLIST WORKSHEET

IHW RCRA INTERIM STATUS FACILITY CME CHECKLIST 8

Reg Ent Name:

Date:

Add ID:

Investigator Name:

| Item No | Description | Answer | Citations | Notes |
|---------|--|--------|---|-------|
| | ANALYTICAL PROCEDURES FOR INTERIM STATUS FACILITIES | | | |
| | SECTION A: REVIEW OF ANALYTICAL PROCEDURES | | | |
| 1 | From the Sampling and Analysis Plan, include a tabulation of analytical methods used for Groundwater samples. Indicate directly on the Attachment which analyses are performed at: off-site contract laboratory (*); on-site operator laboratory (**); field measurement (***) (Attachment). | | | |
| 2 | Laboratory analysis procedures: | | 265.90(a) 335.116(a) 335.112(a)(5) 265.92(a) | |
| 2A | Are all samples analyzed using an EPA-recommended method (SW-846 or other EPA recommended procedures)? | | 335.112(a)(5) 265.92(a) | |
| 2B | Are appropriate QA/QC measures used in laboratory analysis (e.g., blanks, spikes, standards)? | | 265.92(a) 335.112(a)(5) | |
| 2C | Are detection limits and percent recovery (if applicable) provided for each parameter? | | | |
| 2D | If a different analytical method or laboratory is used, are split samples run for comparison purposes? | | | |
| 2E | Describe any data inconsistencies and how the operator has tried to resolve them: | | | |
| 2F | Are samples analyzed within specified holding times? | | 265.92(a) 335.112(a)(5) | |
| 2G | What is the sample analysis turn-around time (i.e., the time required to receive analytical results from the laboratory)? | | | |
| 2H | Example of analytical results and/or QA/QC results as reported by the laboratory to the operator (Attachment). | | | |
| 3 | Laboratory logbook | | 265.90(a) 265.92(a) 335.116(a) 335.112(a)(5) | |
| 3A | Is a laboratory logbook maintained? | | 335.112(a)(5) 265.92(a) | |
| 3B | Are experimental conditions (e.g., temperature, humidity, etc.) noted? | | 265.92(a) 335.112(a)(5) | |
| 3C | If a sample for volatile analysis is received with headspace, is this noted? | | | |
| 3D | Are the results for all QC samples identified? | | | |
| 3E | Is the time, date, and name of person noted for each processing step? | | 335.112(a)(5) 265.92(a) | |
| 3F | Is the date and time of each instrument's calibration noted? | | 265.92(a) 335.112(a)(5) | |

CHECKLIST WORKSHEET

IHW RCRA INTERIM STATUS FACILITY CME CHECKLIST 8 (Cont)

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| 3G | Are notations made on which standards are used and when they were mixed? | · | 335.112(a)(5) 265.92(a) | |
| | SECTION B: REVIEW OF QUALITY ASSURANCE/QUALITY CONTROL | | | |
| 1 | Does the QA/QC program include: | | | |
| 1A | Documentation of any deviations from approved procedures? | · | 265.92(a) 335.112(a)(5) | |
| 1B | Collection and analysis of trip blanks, field blanks and equipment blanks? | · | 335.112(a)(5) 265.92(a) | |
| 1C | Documentation of analytical results for: i. Laboratory blanks? ii. Standards? iii. Duplicates? iv. Other (specify) | · | 335.112(a)(5) 265.92(a) | |
| 2 | Are field QC samples compared with field sample results? (NOTE: If concentrations in blanks are greater by an order of magnitude than the field samples, then resampling is recommended.) | · | 335.112(a)(5) 265.92(a) | |
| 3 | Does the operator critically examine the results to ensure that they have been properly calculated and reported? | · | 335.112(a)(5) 265.92(a) | |
| 4 | Is the validity and reliability of the laboratory and field generated data ensured by a QA/QC program? | · | 335.112(a)(5) 265.92(a) | |