Organizational Planning Meeting for the Chemical Data Reporting (CDR) Inorganic Byproducts Negotiated Rulemaking

May 9-10, 2017

South American Rooms, Capitol Hilton, 1001 16th Street NW, Washington, DC

Agenda

Tuesday, May 9, 2017, 9:00 AM – 5:00 PM

Welcome, Introductions and Agenda Review (9:00-9:45 AM)

Meeting Guidelines Review, Revisions and Approval (9:45-10:00 AM)

Overview of the Mandate for the CDR Inorganic Byproducts Negotiated Rulemaking Committee (10:00-10:15 AM)
EPA will provide brief background on why the Committee was formed and the Agency's mandate.

FACA Expectations (10:15-10:45 AM)
EPA will provide an overview of FACA's legal requirements and EPA's policy positions.

Break (10:45-11:00 AM)

Introduction to Chemical Data Reporting (CDR), Toxics Release Inventory (TRI), and RCRA (11:00-12:15 PM)

- Background on TSCA Section 8(a), including CDR
- Introduction to TRI
- Introduction to RCRA
- Byproduct-related requirements under CDR, TRI, and RCRA

Lunch (12:15-1:30 PM)

Industry Reporting Practices (1:30-2:15 PM)
A sampling of industry members will provide an overview of industry practices associated with reporting inorganic byproducts.

Examples of Past Reporting under CDR, TRI, and RCRA (2:15-3:30 PM)
EPA will describe examples displaying the reporting requirements across these three EPA programs.

Break (3:30-3:45 PM)

Summary of Potential Topics for Committee Discussion (3:45-4:30 PM)
The group will review initial issues for discussion outlined in the Situation Assessment Report and will discuss whether any additional issues need to be considered, based on additional questions that arose during the education and information exchange, or other significant concerns.

Public Comment Period (4:30-5:00 PM)

Wednesday, May 10, 2019, 9:00 AM – 3:00 PM

Check-In and Agenda Review (9:00-9:15 AM)

Discussion of Draft Operating Protocol (9:15-10:15 AM)
The group will discuss any suggested modifications to the Draft Operating Protocol, which includes expectations of Committee Members, consensus decision-making approaches, use of a sub-group to develop agendas, communication protocols, expectations for timeframe and final products, etc. This is also an opportunity to discuss potential approaches for engaging stakeholders that are not Committee Members.

Recap of Day 1 and Refine Issues for Future Discussion (10:15-11:45 PM)
The group will build upon the previous day's discussion to clarify topics for future meetings. This will involve identifying specific questions, information needs and sequencing in order to develop future agendas.

Lunch (11:45-1:00 PM)

Public Comment Period or Continued Discussion (1:00-1:30 PM)

Identify Volunteers for Reg-Neg Process Workgroup to Develop Meeting Agendas (1:30-1:45 PM)

Refine Issues for Future Discussion (Continued) (1:45-2:30 PM)

Meeting Wrap-Up (2:30-3:00)
The group will review next steps and any logistical reminders and share feedback on the process.