

US EPA ARCHIVE DOCUMENT



Module 5: Remedial Process



Module Overview

- ◆ Module Objective: Discuss the definition of a remedial action and explain the major steps and activities performed in conducting remedial actions under CERCLA
- ◆ Topics:
 - » Phases of the remedial process
 - » General requirements for remedial actions under CERCLA
 - » Role and responsibilities of the RPM
 - » General activities and requirements for each phase of the remedial process
 - » Management of documents and other records

Definition of Remedial Action

- ◆ Long-term action to achieve a permanent remedy
- ◆ Typical remedial actions



Major Phases in Remedial Process

- ◆ NCP defines six major phases in the remedial process
 - » RI
 - » FS
 - » Selection of remedy
 - » RD
 - » RA
 - » O&M (including site closeout)



Remedial Program Goal and Management Principles

- ◆ National goal of remedy selection is to select remedies that are:
 - » Protective of human health and the environment
 - » Maintain protection over time
 - » Minimize untreated waste
- ◆ Management principles:
 - » Sites should generally be remediated in OUs
 - » OUs should not be inconsistent with final remedy
 - » Data collection and alternative analysis reflects scope and complexity of site problems

Expectations for Developing Appropriate Remedial Alternatives

- ◆ Use treatment to address principal threats wherever practicable
- ◆ Use engineering controls to address long-term, lower level threats
- ◆ Use a combination of treatment, engineering controls, and institutional controls
- ◆ Use institutional controls to supplement engineering controls
- ◆ Consider using innovative technologies
- ◆ Return usable groundwater to its beneficial uses wherever practicable

Risk Management Principles: Remedial Action Objectives

- ◆ RAOs specify contaminants and media of concern, potential exposure pathways, and remediation goals
- ◆ PRGs are used initially and adjusted based on site-specific information
- ◆ Final remediation goals are determined when the remedy is selected

Risk Management Principles: Remediation Goals

- ◆ Establish acceptable exposure levels
- ◆ Must consider ARARs and facility siting laws and the following factors:
 - » For systemic toxicants, levels that do not cause adverse effects
 - » For carcinogens, levels corresponding to excess lifetime cancer risk between 10^{-4} and 10^{-6}
 - » For carcinogens, use 10^{-6} level as point of departure for PRGs when ARARs are not available or sufficiently protective because of multiple contaminants and pathways
 - » Factors related to technical limitations and uncertainty
 - » Other pertinent information

Risk Management Principles: Additional Considerations for RAOs

- ◆ Attain MCLG set above zero where relevant and appropriate
- ◆ Attain MCL when MCLG is set at zero
- ◆ Attain water quality criteria where relevant and appropriate
- ◆ Establish alternative concentration level in accordance with CERCLA
- ◆ Evaluate environmental threats

Responsibilities of the RPM

- ◆ The RPM has the authority and the responsibility to:
 - » Participate in all decision-making processes
 - » Coordinate with all parties involved



Overview of the RI/FS Process

- ◆ RI/FS supports selection of the remedy
- ◆ RI is performed to characterize the site
- ◆ FS is used to develop and analyze remedial action alternatives
- ◆ Various lead agencies may conduct the RI/FS



Interdependency of the RI and FS



Main Activities of the RI

- ◆ Site characterization
- ◆ Baseline risk assessment
- ◆ Treatability studies



Main Activities of the FS

- ◆ Development and screening of alternatives
- ◆ Detailed analysis of alternatives



Development and Screening of Alternatives

- ◆ Development and screening involve seven general steps:
 - » Refine the RAOs
 - » Develop general response actions
 - » Identify volumes or areas of media
 - » Identify remedial technologies
 - » Identify and screen process options
 - » Assemble alternatives
 - » Screen alternatives



Scoping the RI/FS

- ◆ Goal of RI/FS is to support the ROD
- ◆ RI/FS process begins with scoping
- ◆ Key Activities
 - » Form a site team
 - » Conduct a kickoff meeting with the site team
 - » Develop a CSM
 - » Identify initial DQOs
 - » Identify RAOs, general response actions, and ARARs
 - » Collect and evaluate existing data
 - » Conduct a site visit

RI/FS Project Plans

- ◆ The RPM is responsible for overseeing the preparation of several work plans
 - » RI/FS work plan
 - » SAP
 - QAPP
 - FSP
 - » HASP
 - » CIP



RI/FS Report

- ◆ The RI/FS report
 - » Forms foundation of support in remedy selection process
 - » Documents the development and screening of remedial alternatives

Detailed Analysis of Alternatives

- ◆ Nine evaluation criteria are the basis for remedy selection
- ◆ The nine criteria are separated into three levels
 - » Threshold criteria
 - » Balancing criteria
 - » Modifying criteria



Nine Evaluation Criteria

- ◆ Threshold criteria
 - » Overall protection of human health and the environment
 - » Compliance with ARARs
- ◆ Balancing criteria
 - » Long-term effectiveness and permanence
 - » Reduction of toxicity, mobility, or volume through treatment
 - » Short-term effectiveness
 - » Implementability
 - » Cost
- ◆ Modifying criteria
 - » State (support agency) acceptance
 - » Community acceptance

Overview of the Remedy Selection Process

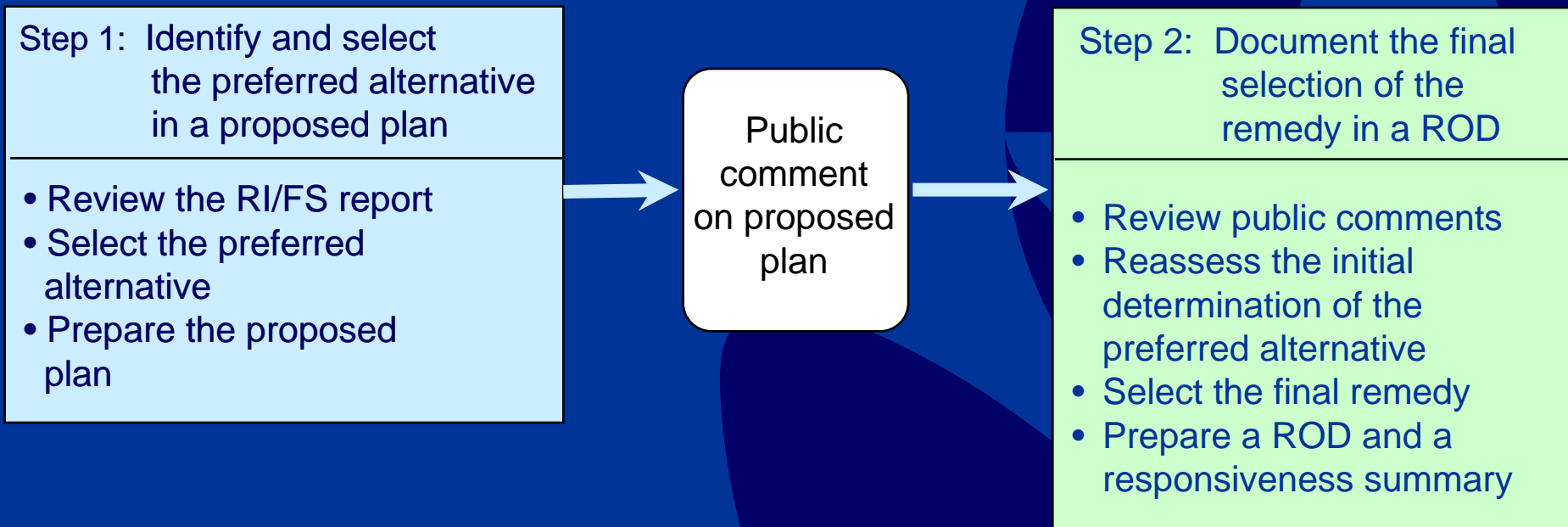
- ◆ CERCLA establishes specific requirements
- ◆ The NCP establishes procedures for proposing and documenting final remedy
- ◆ Changes that occur after selection of the remedy need to be addressed
- ◆ EPA reforms to improve the quality of selected remedies



CERCLA Requirements for Remedy Selection

- ◆ Protect human health and the environment
- ◆ Attain (or waive) ARARs
- ◆ Be cost-effective
- ◆ Use permanent solutions and alternative treatment technologies to the maximum extent practicable
- ◆ Satisfy the preference for treatment as a principal element of the remedy
- ◆ Involve states in a substantial and meaningful manner
- ◆ Consistency with the NCP

The Remedy Selection Process



Proposed Plan

- ◆ Highlights key aspects of the RI/FS
- ◆ Describes remedial alternatives
- ◆ Explains the rationale for selection of the preferred alternative
- ◆ Requests comments from the public
- ◆ Includes the views of support agencies



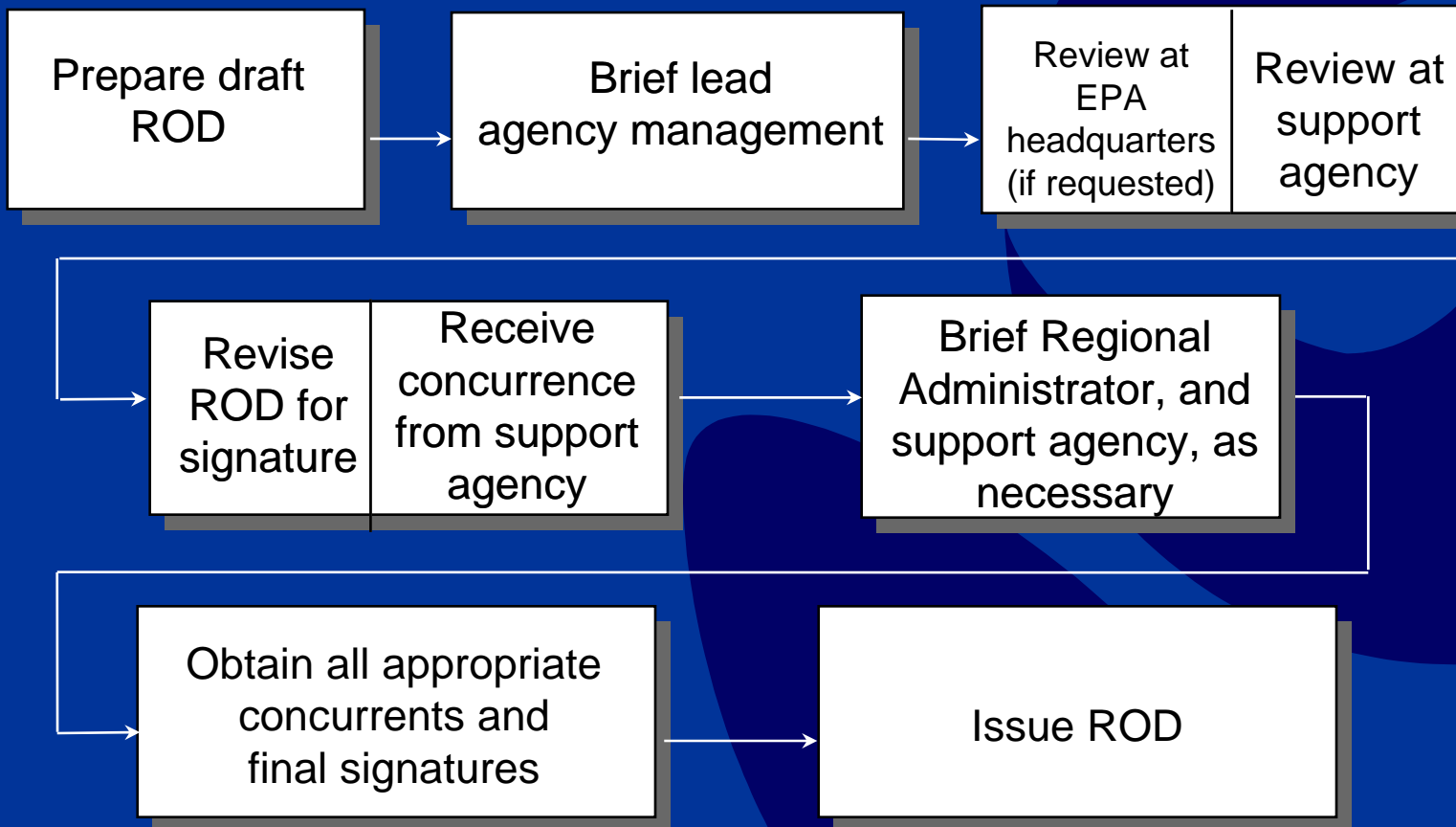
Document the Final Selection of the Remedy in a ROD

- ◆ Review public comments
 - » Address significant changes in the selected remedy
 - » Solicit additional comments
- ◆ Reassess initial determination of the preferred alternative
- ◆ Select the final remedy
 - » The lead agency has responsibility
 - » The PRP should not influence the decision
 - » The best balance among the criteria should be evaluated

ROD and Responsiveness Summary

- ◆ A ROD is:
 - » The official decision document on remedy selection
 - » A technical, legal, and public document
- ◆ A responsiveness summary is:
 - » A written summary of responses to each significant comment submitted during the public comment period

Preparing the ROD



(continued)

Preparing the ROD

- ◆ The lead agency prepares the ROD
- ◆ EPA retains final authority over remedy selection
- ◆ The ROD is supported by documents in the AR
- ◆ Follow procedures for review, concurrence, and signature

Content of the ROD

- ◆ Declaration
 - » Formal statement signed by RA
- ◆ Decision summary
 - » Overview of problems and risks
 - » Rationale for remedy selection
- ◆ Responsiveness summary
 - » Addresses comments received



Post-ROD Changes

- ◆ Changes are prompted by new information, enforcement agreements, or developments during RD/RA
- ◆ There are three types of ROD changes
 - » Minor
 - » Significant
 - » Fundamental
- ◆ Different requirements and procedures apply to each type of post-ROD change



Superfund Reforms — Remedy Selection

- ◆ Superfund reforms improve consistency in remedy selection

