

# NIOSH 10-Year Program Review Implementation

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**December 2011**

**Tampa, FL**

# Five Review Reports

- Reports focus on the following topic areas:
  - Dose Reconstruction
  - Quality of Service
  - Timeliness of Program Products
  - SEC Petitions
  - Quality of Science
- Summary of the reports and recommendations are available on the NIOSH Web site at:

<http://www.cdc.gov/niosh/docket/review/docket194/default.html>)

# Quality of Service

- **Issues related to customer-supplied information**
  - Develop system(s) for collecting all customer comments along with disposition action
- **Issues related to understandability of information**
  - Improve understandability and quality of communication vehicles: Web site, information sheets, letters to claimants or petitioners, etc.
- **Issues related to access to information**
  - Ensure Board and Work Group work products are posted on the Web site as soon as practical

# Timeliness

- Give higher priority to claims returned by DOL for new dose reconstruction
- Continue to adopt aggressive timeliness objectives for dose reconstruction
- Adopt aggressive time limits for the completion of the review of SEC petitions

# SEC Petitions

- Separate “policy” issues from “science” issues
  - Include more detail on the evolution of a science decision in SEC Evaluation Reports (e.g., thorium exposures)
  - Develop a policy memo with each Evaluation Report that identifies policy decision and the thought process behind the decisions
- Work to define “sufficient accuracy”
- Utilize staff “other than health physicists” when appropriate to guard against “professional orientation” toward accepting adequacy of techniques

# Quality of Science

- **Consider subjecting program documents to formal peer review**
  - NIOSH is developing an implementation guide to identify levels of review of program documents
- **Assess validity of indirect exposure methods**
  - NIOSH is conducting trial-run validation using Savannah River Site data

# Quality of Science—cont.

- Characterize degree of claimant-favorability in current methods
  - Compare “claimant-favorable” approaches to assessment methods commonly used in other applications
- Evaluate utility of EPA surrogate data protocol
- NIOSH review of EPA protocol indicates its recommendations are similar to information in “The Use of Data from Other Facilities in the Completion of Dose Reconstructions Under EEOICPA” (DCAS-IG-004)
  - Non-NIOSH reviewers will review EPA protocol against methods used by NIOSH

# Dose Reconstruction

- **Work with Subcommittee on Dose Reconstruction Reviews on QA/QC evaluation**
  - Develop duplicate analysis program in order to identify potential issues with dose reconstruction process
  - Review recent dose reconstruction review findings to determine cause of findings
- **Prioritize elimination of overestimating dose reconstructions relative to implementation of other program review actions**
  - Analyze cost of eliminating overestimating dose reconstructions
  - Consider impact on long-range budget planning