

Next Steps towards Revising Radiation Protection Regulations (10 CFR Part 20)

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Background

- **ICRP revised recommendations announced in December, 2007**
- **NRC staff analysis indicated areas warranting consideration for revisions – SECY-08-0197, December, 2008**
- **Commission approved staff recommendation to engage stakeholders and initiate development of technical basis materials on April 2, 2009**
- **Staff recommendations – SECY-12-0064, April 25, 2012**

SRM-SECY-12-0064

Recommendations for Policy and Technical Direction to Revise Part 20

- The Commission issued the Staff Requirements Memorandum (SRM) to the staff on December 17, 2012.
- The Commission approved in part, and disapproved in part, the staff's recommendation from SECY-12-0064.
- The staff is moving forward to implement the Commission's direction.

Revise Methodology and Terminology

- **Commission Direction:**
 - Develop a regulatory basis for a revision to 10 CFR Part 20 to align with the most recent methodology and terminology for dose assessment.
 - Develop a regulatory basis for parallel alignment of 10 CFR Part 50, Appendix I.
 - Make corresponding changes in other portions of the regulations.
- **Proposal:**
 - TEDE becomes TED
 - New W_T and W_R values incorporated into definitions
 - Appendix B revised with new ALI and DAC values

Revise Methodology and Terminology

- **Key Questions:**
 - **What would be an appropriate time frame and approach to transition of terminology?**
 - **Consistent methodology for calculations, particularly for members of the public.**

Limit for Occupational TEDE

- **Commission Direction:**
 - **Disapproved staff's recommendation to develop the regulatory basis to reduce the occupational total effective dose equivalent (TEDE).**
 - **Continue discussions with stakeholders on alternative approaches to deal with individual protection at or near the current dose limit.**
- **Objective:**
 - **Regulatory requirements and guidance that will ensure that cumulative exposures are examined, and that progressive restrictions can be taken as cumulative exposures increase.**

Individual Protection - ALARA

- **Options:**
 - Performance based requirement added to ALARA and Radiation Protection Programs, with guidance
 - Prescriptive requirements
 - Require licensees to establish an administrative control level (ACL) as part of their radiation protection program and to establish specific procedures for individual protection
 - Require licensees to have a record of all occupational doses (lifetime) if exposures are permitted to exceed 20 mSv per year
 - Require that licensees not allow occupational exposures to exceed 20 mSv in a year if the cumulative occupational exposure exceeds xxx mSv
 - Require licensees be provided with record of all other sources of occupational exposure
 - Other options.

Individual Protection - ALARA

- **Key Questions:**
 - How does each approach work for different classes of licensed use?
 - Should licensees be allowed to establish different ACL's for different groups of individuals?
 - Is there another mechanism to look at cumulative exposures?
 - Should States be allowed to use prescriptive requirement if NRC decides to use performance based approach?
 - What is impact on licensee activities? State regulatory programs?

Occupational Limit - Lens of the Eye

- **Commission Direction:**
 - **Continue discussions with stakeholders regarding possible revisions to the dose limit for the lens of the eye**
- **Proposal:**
 - **Develop regulatory basis for reducing limit to 50 mSv LDE**

Occupational Limit - Lens of the Eye

- **Key Questions:**
 - Are there alternatives to keep cumulative exposure below threshold?
 - Viewpoints on the relative importance of health endpoint?
 - What methods should be allowed for measurement or assessment?
 - What methods should be allowed for recording dose when eye is protected?
 - What is impact on licensee activities? State regulatory programs?

Occupational Limit - Embryo/Fetus

- **Commission Direction:**
 - **Continue discussions with stakeholders**
- **Proposal:**
 - **Develop regulatory basis for reducing limit to 1 mSv**

Occupational Limit - Embryo/Fetus

- **Key Questions:**

- **Apply to post declaration or entire gestation period?**
- **What should be done if 1 mSv has already been reached at declaration?**
- **What methods should be allowed for measurement or assessment?**
- **What is impact on licensee activities? State regulatory programs?**

Units of Exposure and Dose

- **Commission Direction:**
 - Disapproved the elimination of traditional units from NRC regulations. Both units should be maintained.
- **Proposal:**
 - Implement Commission Policy Statement – SI units first, traditional units in parenthesis

Units of Exposure and Dose

- **Key Questions:**
 - How do we avoid confusion?
 - Should Appendix B be given in SI units, or traditional, or both?
 - Should licensees be allowed to report in SI?
 - What is impact on licensee activities? State regulatory programs?

Reporting of Occupational Dose

- **Commission Direction:**
 - Improve reporting of occupational exposure by NRC and Agreement State licensees, some of which do not currently submit reports.
- **Proposal:**
 - Add categories of licensed use: e.g., Part 35, medical
 - Modify requirements for compatibility
 - Explore mechanisms for central repository of data for all to use

Reporting of Occupational Dose

- **Key Questions:**
 - **What categories should be included?**
 - **What is the rationale for reporting?**
 - **What are health and safety, and/or trans-boundary considerations?**
 - **How to deal with occupational exposure of machine produced radiations?**
 - **What is impact on licensee activities? State regulatory programs?**

Next Steps

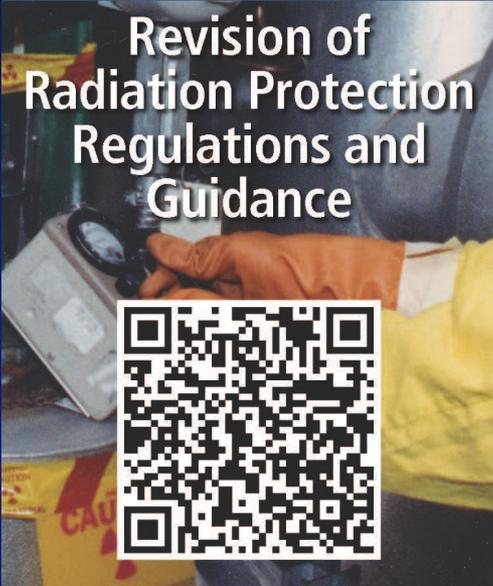
- Engage Federal Agencies, States, licensees, and with public stakeholders on each of the topics.
- The staff will develop regulatory basis using Commission direction for each technical issue.
- Develop *Federal Register* Notice with specific proposed options and questions – plan to publish for input this summer.
- Possibility of webinars.
- Further opportunities for comment in 2014 with more specific proposals.
- The tentative date for development of the regulatory basis is December, 2015.

Questions? Questions?

<http://www.nrc.gov/about-nrc/regulatory/rulemaking/potential-rulemaking/opt-revise.html>


United States Nuclear Regulatory Commission
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**Revision of
Radiation Protection
Regulations and
Guidance**



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