

U.S. NRC Discussion of Options to Revise Radiation Protection Recommendations

2011 ICRP International Symposium

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NRC - Who We Are

- **Atomic Energy Act of 1954 formed the Atomic Energy Commission**
- **The Energy Reorganization Act of 1974 established the independent U.S. NRC to regulate commercial uses of nuclear material.**
- **Energy Policy Act of 2005 added new types of Byproduct Material**



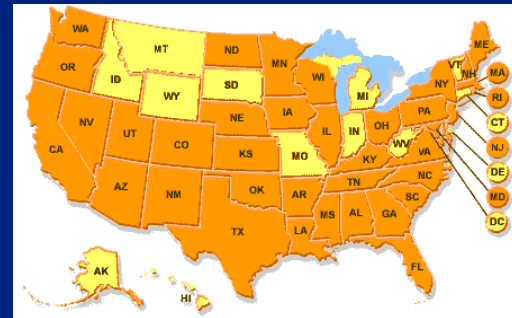
NRC - What We Regulate

- **Nuclear reactors** - commercial power reactors, research and test reactors, new reactor designs
- **Nuclear materials** - nuclear reactor fuel, radioactive materials for medical, industrial and academic use
- **Nuclear waste** – transportation, storage and disposal of nuclear material and waste, decommissioning of nuclear facilities



Who Does What?

- **NRC and States** regulate radioactive materials
 - Authority relinquished by Agreements to States (37)
 - Adequacy and Compatibility requirements
 - Some things essentially identical
 - Some things States can be more restrictive
 - Some things (reactors) reserved to NRC
- **States** regulate machine produced radiation
 - X-ray, CT, Mammography, Fluoroscopy, etc.
 - Most State radiation protection requirements match for materials and machine produced radiation



Other Responsible Organizations

- **Federal**
 - EPA – General and Environmental Standards
 - DOE – Military, Promotion, Education
 - HHS – Medical and Devices
 - DHS – Security and Emergency Response
- **Interagency Steering Committee on Radiation Standards**
 - Forum for Federal agencies to keep abreast of national and international radiation protection activities
 - Identifying interagency issues and coordinating their resolution



Background of Regulations

- NRC regulations last revised in 1991
- Requirements in Part 20, Licensing Parts
- NRC staff analysis indicated areas warranting consideration for revision
- Commission approved staff recommendation to engage stakeholders and initiate development of technical basis materials on April 2, 2009



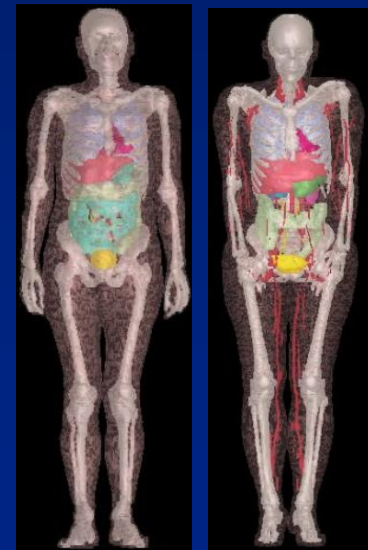
Outreach Activities

- **Phase I of outreach included:**
 - Presentations to numerous organizations and groups
 - FRN published inviting inputs (72 FR 32198)
- **Phase II Workshops**
 - FRN published with issues and questions (75 FR 59160)
 - Workshops in Washington, Los Angeles, and Houston
- **Phase III Comment – Lens of the Eye**
 - FRN published asking for feedback (76 FR 53847)
 - Comments due by October 31, 2011



TED and Numerical Values

- **Issue: Update terms and scientific information?**
- **Feedback:**
 - **General support for updating numerical values and scientific base**
 - **Mixed views on terminology**
 - **Many suggested delaying rulemaking until ICRP completes work on dose coefficients**
 - **Some discussion of moving from Regulation to Guidance**



Occupational Dose Limits

- **Issue: Change the Occupational Dose Limit?**
- **Feedback:**
 - Little support for change to regulation
 - Certain groups of licensees continue to have individuals above 20 mSv/yr (2 rem)
 - Legal Boundary for enforcement needs to remain as is
 - ALARA has resulted in achieving desired dose reductions
 - Many do not believe changes in risk justify change to limit



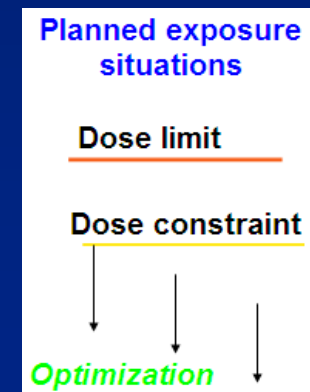
Lens of the Eye

- **Issue: New Recommendation from ICRP**
 - ICRP recommendation is now 20 mSv (2 rem) over 5 years, with a maximum of 50 mSv (5 rem) in any one year
 - Part 20 limit is 150 mSv (15 rem) per year
 - Fluoroscopy and other procedures contribute significantly
- **Feedback:**
 - Caution needed in making any changes
 - Numeric value for LDE could be the same as the numeric value of TED, to avoid compliance issues



ALARA Planning

- **Issue: Add to requirements for ALARA?**
- **Feedback:**
 - Most licensees do planning to reduce exposures and use a variety of criteria to trigger actions
 - Little support for using the term “constraint”
 - Many concerned that any numerical values in regulations will be a *de facto* limit
 - Some support for explicitly requiring planning, but with reservations of what inspectors would be expecting in licensing programs



Path Forward

- **Policy paper for Commission consideration in April 2012**
- **Development of Technical Basis to support Commission decisions**
- **It is still “too soon to tell” what the staff will ultimately recommend**
- **Comments and views welcomed**



Interagency

- NRC working with interagency through ISCORS to keep them up to date on stakeholder dialogue
- Federal Agencies funding for development of dose coefficients
- Discussions underway on need to update Federal Guidance documents (EPA lead)



Resources

- Web pages

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

- Email Address: regs4rp@nrc.gov

Rulemaking.Comments@nrc.gov

- Rulemaking Web Site:

<http://www.regulations.gov>

Docket ID: NRC-2009-0279

Questions?

