
A Review of National Dose and Risk Criteria

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R. Ferch

The LTSC Initiative of the RWMC-RF:

- RWMC's Regulators' Forum formed in 1999
 - Comparative Study of Regulatory Structures
 - Included an initial comparison of long-term radiological protection criteria
 - Long-Term Safety Criteria Group formed to investigate in more detail
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Information sources:

- ICRP-81(1999), IAEA WS-R-4(2006)
- National criteria based on information from the NEA RWMC Regulators' Forum



International recommendations - ICRP

- ICRP-81: 0.3 mSv/a dose constraint (risk on the order of $10^{-5}/a$)
 - Beyond a few hundred years, calculated doses are considered as performance indicators, not measures of health detriment
 - Quantitative calculations for $10^3 - 10^4$ years, stylized or qualitative calculations further into the future
 - Note that with current dose/risk conversion factors, 0.3 mSv/a $\sim 2 \times 10^{-5}/a$ risk
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International recommendations - IAEA

- WS-R-4: 0.3 mSv/a dose constraint, on the order of 10^{-5} /a risk constraint
 - On long time scales the criteria may no longer serve as a reasonable basis for decision making
 - Criteria = targets rather than limits in the long term
 - Appendix suggests comparison with natural background at very long time scales (10^6 yr)
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National Criteria(1)

- Belgium: 0.1-0.3 mSv/a (high probability), $10^{-5}/a$ risk (low p) (interim working values)
 - Canada: 0.3 mSv/a (interim working value), timescales guidance under development
 - Czech Republic: 0.25 mSv/a for 10^6 yr, $10^{-6}/a$ probability cutoff
 - Finland: 0.1 mSv/a for several thousand yr, equivalent risk for unlikely events
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National Criteria(2)

- France: 0.25 mSv/a for 10^4 yr, **target** later
 - Germany: 0.1 mSv/a for 10^6 yr; dose targets for different classes of likelihood (under revision)
 - Hungary: 0.1 mSv/a, 10^{-5} /a risk for disruptive events, 10^{-7} /a probability cutoff
 - Rep. of Korea: 0.1 mSv/a, 1 mSv for intrusion, 10^{-6} /a risk for disruptive events (under development)
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National Criteria(3)

- Netherlands: 0.1 mSv/a limit (**target** of 0.04 mSv/a), normal evolution
 - Slovakia: 0.1 mSv/a, 1 mSv/a for intrusion (under development)
 - Spain: 0.1 mSv/a, 10^{-6} /a risk for low probability (under revision)
 - Sweden: 10^{-6} /a risk (0.014 mSv/a) for 1000 y, general consideration of scenarios after that
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National Criteria(4)

- Switzerland: 0.1 mSv/a, 10^{-6} /a risk for low probability, applied at all times
 - UK: 10^{-6} /a risk **target** at all times, dose constraint of 0.3 mSv/a before withdrawal of control
 - USA: 0.15 mSv/a for 10^4 yr, (proposed) 3.5 mSv/a up to 10^6 yr (both apply to human intrusion as well as normal evolution)
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Bases for Criteria (Implied or Explicit)

- ICRP 0.3 mSv/a dose constraint based on radiation protection arguments, often reduced (e.g. to 0.1 mSv/a)
- Absolute risk arguments (e.g. from $10^{-4}/a$ barely tolerable to $10^{-6}/a$ broadly acceptable level)
- Arguments based on natural background (~ 3 mSv/a) or on variability of background

Supplementary Criteria

- Dose for normal scenarios, risk for low likelihood (e.g. Finland, Spain, Switzerland)
 - Radionuclide fluxes (Finland at long time scales)
 - Quantitative assessments at long time scales or for low-probability events (e.g. Sweden)
 - Optimization (ALARP) approach (e.g. UK, France at long time scales)
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Design-based criteria and BAT

- Covered only incidentally in the questionnaire, may include:
 - engineering design requirements
 - complete containment for several 100's of years
 - restrictions on predicted release rates
 - restrictions on the geology
 - BAT or BAT(neec) (Ref EC IPPC Directive)
 - Optimization applied not to calculated doses/risks as in ALARP, but to parameters more directly related to design
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Treatment of Uncertainties

- Probabilistic, e.g. low-likelihood scenarios
 - e.g. risk criterion replaces dose for low-likelihood events
 - dose limits varying with class of likelihood (normal vs. disruptive, or more detailed classification)
 - Parameter uncertainties
 - e.g. sensitivity analysis, probabilistic analysis
 - Uncertainties inherent in long time scales
 - e.g. unknown characteristics of the receiving population, geological uncertainties, performance modelling uncertainties
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Other Sources of Differences

- Modelling strategies
 - choice of parameters (design centre vs. conservative, etc.)
 - choice of critical group (or most exposed individual)
 - choice of analysis scenarios
 - Differences of interpretation
 - use of hard limits vs. targets
 - allowances for judgment, application of other lines of reasoning
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What does it mean?

- Are the readily observed differences in criteria correlated with differences in the actual level of protection?
 - No way of knowing – other uncertainties and differences dominate
 - There is a large national cultural component
 - Comparison with experience in environmental protection
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Some possibilities for further investigation:

■ Ethical questions:

- ❑ Level of protection for distant future – constant or varying? constant duty but varying ability to guarantee?
- ❑ Transfer of burdens, responsibilities and resources to near-future generations

■ Socio-political questions:

- ❑ Decision-making in the broad sense vs. regulation
 - ❑ Importance of national culture vs. international standards
 - ❑ Investigation of fundamental protection goals
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Some possibilities for further investigation:

- Technical questions:
 - Harmonization of limits/constraints?
 - Compliance criteria for distant future?
 - “Complementary” criteria (not dose- or risk-based)
 - BAT?
 - Technical performance criteria (concentrations, fluxes)?
 - Design standards?
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