

Submission of Federal Rules Under the Congressional Review Act

☒ President of the Senate☐ Speaker of the House of Representatives☐ GAO

Please fill the circles electronically or with black pen or #2 pencil.

1. Name of Department or Agency

U.S. Nuclear Regulatory Commission

2. Subdivision or Office

Office of Nuclear Material Safety and Safeguards

3. Rule Title

Medical Use of Byproduct Material: Clarifying and Minor Amendments (10 CFR Part 35)

4. Regulation Identifier Number (RIN) or Other Unique Identifier (if applicable)

RIN 3150-AH08

5. Major Rule ☐ Non-major Rule ☒

6. Final Rule ☒ Other ☐

7. With respect to this rule, did your agency solicit public comments?

Yes ☒ No ☐ NA ☐

8. Priority of Regulation (fill in one)

☐ Economically Significant; or
Significant; or
Substantive, Non Significant

☒ Routine and Frequent or
Informational/Administrative/Other
(Do not complete the other side of this form
if filled in above.)

9. Effective Date (if applicable)

75 days after publicatuin in FR unless significant adverse comments are recieved

10. Concise Summary of Rule (fill in one or both)

attached ☒ stated in rule ☐

Submitted by: _____ (signature)

Name: _____

Title: _____

For Congressional Use Only:

Date Received: _____

Committee of Jurisdiction: _____

	Yes	No	N/A
A. With respect to this rule, did your agency prepare an analysis of costs and benefits?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
B. With respect to this rule, by the final rulemaking stage, did your agency			
1. certify that the rule would not have a significant economic impact on a substantial number of small entities under 5 U.S.C. § 605(b)?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
2. prepare a final Regulatory Flexibility Analysis under 5 U.S.C. § 604(a)?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
C. With respect to this rule, did your agency prepare a written statement under § 202 of the Unfunded Mandates Reform Act of 1995?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
D. With respect to this rule, did your agency prepare an Environmental Assessment or an Environmental Impact Statement under the National Environmental Policy Actg (NEPA)?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
E. Does this rule contain a collection of information requiring OMB approval under the Paperwork Reduction Act of 1995?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
F. Did you discuss any of the following in the preamble to the rule?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
• E.O. 12612, Federalism	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
• E.O. 126630, Government Actions and Interference with Constitutionally Protected Property Rights	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
• E.O. 12866, Regulatory Planning and Review	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
• E.O. 12875, Enhancing the Intergovernmental Partnership	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
• E.O. 12988, Civil Justice Reform	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
• E.O. 13045, Protection of Children from Environmental Health Risks and Safety Risks	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
• Other statutes or executive orders discussed in the preamble concerning the rulemaking process (please specify)			

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