

Submission of Federal Rules Under the Congressional Review Act

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A. With respect to this rule, did your agency prepare an analysis of costs and benefits? 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)? 2. prepare a final Regulatory Flexibility Analysis under 5 U.S.C. § 604(a)? C. With respect to this rule, did your agency prepare a written statement under § 202 of the Unfunded Mandates Reform Act of 1995? 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)? E. Does this rule contain a collection of information requiring OMB approval under the Paperwork Reduction Act of 1995? F. Did you discuss any of the following in the preamble to the rule? • E.O. 12612, Federalism • E.O. 126630, Government Actions and Interference with Constitutionally Protected Property Rights • E.O. 12866, Regulatory Planning and Review • E.O. 12875, Enhancing the Intergovernmental Partnership • E.O. 12988, Civil Justice Reform • E.O. 13045, Protection of Children from Environmental Health Risks	No	N/A
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and Safety Risks	\bigcirc	•
 Other statutes or executive orders discussed in the preamble concerning the rulemaking process (please specify) 		



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 bl	lack pen or #2 pencil.
Name of Department or Agency	2. Subdivision or Office
U.S. Nuclear Regulatory Commission	FSME
3. Rule Title Medical Use of Byproduct Material - A	Authorized User Clarification
4. Regulation Identifier Number (RIN) or Other I	Unique Identifier (if applicable)
5. Major Rule Non-major Rule	
6. Final Rule Other	
7. With respect to this rule, did your agency soli	icit public comments? Yes No N/A
8. Priority of Regulation (fill in one) Economically Significan Significant; or Substantive, Non Significant	Informational/Administrative/Other
9. Effective Date (if applicable)	
10. Concise Summary of Rule (fill in one or both	h) attached stated in rule
Submitted by: Name: Rebecca Schmidt	(signature)
Title: Director, Office of Congress	sional Affairs
For Congressional Use Only:	
Date Received:	
Committee of Jurisdiction:	

3/23/99



		Yes	No	N/A
Δ.	With respect to this rule, did your agency prepare an analysis of costs and benefits?	\circ	\bigcirc	•
В.	With respect to this rule, by the final rulemaking stage, did your agency			
	 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)? 	\bigcirc	\bigcirc	•
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	E.O. 12866, Regulatory Planning and Review	\bigcirc	\bigcirc	\odot
	E.O. 12875, Enhancing the Intergovernmental Partnership	\bigcirc	\bigcirc	\odot
	E.O. 12988, Civil Justice Reform	\bigcirc	\bigcirc	\odot
	 E.O. 13045, Protection of Children from Environmental Health Risks and Safety Risks 	\circ	\bigcirc	•
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Name of Department or Agency	2. Subdivision or Office
U.S. Nuclear Regulatory Commission	FSME
3. Rule Title Medical Use of Byproduct Material - A	uthorized User Clarification
Regulation Identifier Number (RIN) or Other U RIN 3150-AI59	Inique Identifier (if applicable)
5. Major Rule Non-major Rule	•
6. Final Rule Other	
7. With respect to this rule, did your agency solid	cit public comments? Yes No N/A
8. Priority of Regulation (fill in one) Economically Significant Significant; or Substantive, Non Significant	Informational/Administrative/Other
9. Effective Date (if applicable)	
10. Concise Summary of Rule (fill in one or both	attached stated in rule •
Submitted by: Name: Rebecca Schmidt	(signature)
Title: Director, Office of Congressi	ional Affairs
For Congressional Use Only:	
Date Received:	
Committee of Jurisdiction:	



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