January 13, 2010

MEMORANDUM TO: Michael T. Lesar, Chief
Rulemaking, Directives and Editing Branch
Division of Administrative Services
Office of Administration

FROM: Edward M. Lohr
Rulemaking Branch B **/RA/**
Division of Intergovernmental Liaison
and Rulemaking, FSME

SUBJECT: REGULATORY HISTORY INDEX – DIRECT FINAL RULE: MEDICAL USE OF BYPRODUCT MATERIAL – "AUTHORIZED USER CLARIFICATION" (74 FR 33901) RIN 3150-AI59

Enclosed is the regulatory history index of those documents of central relevance to the subject proposed rule. The ADAMS number is provided for each document or package. The direct final rule amended several sections of 10 CFR Part 35 to clarify that individuals who do not need to comply with the training requirements under § 35.57 may serve as preceptors and work experience supervisors for individuals seeking recognition on an NRC license for the same uses. Additionally, several minor administrative changes were included in this rulemaking. The direct final rule titled "Authorized User Clarification" was published in the *Federal Register* on July 14, 2009 (74 FR 33901). A confirmation of effective date for the direct final rule was published in the *Federal Register* on August 27, 2009 (74 FR 43619).

Enclosure: Regulatory History Index

CONTACT: Edward M. Lohr, FSME/DILR
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(301) 415-0253

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OFFICE	RB:DILR	RB:DILR	
NAME	ELohr	MDelligatti	
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Accession Number	Title/Description	Document Date	Author	Public Availability
ML090090335	Technical Basis for Part 35 "Medical Use of Byproduct Material – Authorized User Clarification" Rulemaking	1/15/2009		Non-Publicly Available
ML091170198	Office Concurrence Package for Part 35 "Medical Use of Byproduct Material – Authorized User Clarification" Rulemaking			Non-Publicly Available
ML091260782	Agreement State Package for Part 35 "Medical Use of Byproduct Material – Authorized User Clarification" Rulemaking			Non-Publicly Available
ML091170198	Office Concurrences for Part 35 "Medical Use of Byproduct Material – Authorized User Clarification" Rulemaking			Publicly Available

Accession Number	Title/Description	Document Date	Author	Public Availability
ML091620631	EDO Package for Part 35 “Medical Use of Byproduct Material – Authorized User Clarification” Rulemaking			Non-Publicly Available
ML091870428	ADM Package for Part 35 “Medical Use of Byproduct Material – Authorized User Clarification” Rulemaking			Publicly Available
ML091801075	DFR Signed by EDO - Part 35 “Medical Use of Byproduct Material – Authorized User Clarification” Rulemaking	6/29/2009		Non-Publicly Available
ML092300018	DFR – Confirmation of Effective Date - Part 35 “Medical Use of Byproduct Material – Authorized User Clarification” Rulemaking	2/26/2008	Thevenot L I	Publicly Available
ML100130683	Regulatory History Index – Direct Final Rule: Medical Use of Byproduct Material – “Authorized User Clarification” (74 FR 33901) RIN 3150-AI59	1/13/2010	Lohr, E. M.	Publicly Available