

July 18, 2018

10 CFR PART 32

Please Note: The bracket A [] A around a compatibility category designation means that the Section may have been adopted elsewhere in a State rules and it is not necessary to adopt it again.

NRC Regulation Section	Section Title	State Section	Compatibility Category	Difference Yes/No	Significant Yes/No	If Difference, Why or Why Not was a Comment Generated
§32.1 (a), (b), & (c)(2)	Purpose and Scope		D	N/A		
§32.1 (c)(1)	Purpose and Scope		NRC			
§32.2	Definitions					
	Committed dose		D	N/A		
	Dose commitment		[A]			
	Lot Tolerance Percent Defective		B			
	Nationally tracked Source		B			
	Sealed source and device registry		D	N/A		

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§32.3	Maintenance of records		D	N/A		
§32.8	Information collection requirements: OMB approval		D	N/A		
§32.11	Introduction of byproduct material in exempt concentrations into products or materials and transfer of ownership or possession: Requirements for license		NRC			
§32.12	Same: Records and material transfer reports		NRC			
§32.13	Same: Prohibition of introduction		C			

NRC Regulation Section	Section Title	State Section	Compatibility Category	Difference Yes/No	Significant Yes/No	If Difference, Why or Why Not was a Comment Generated
§32.14	Certain items containing byproduct material; requirements for license to apply or initially transfer		NRC			
§32.15	Same: Quality assurance, prohibition of transfer and labeling		NRC			
§32.16	Certain items containing byproduct material: Records and reports of transfer		NRC			
§32.18	Manufacture, distribution and transfer of exempt quantities: Requirements for license		NRC			

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§32.19	Same: Conditions of licenses		NRC			
§32.20	Same: Records and material transfer reports		NRC			
§32.21	Radioactive drug: Manufacture, preparation or transfer for commercial distribution of capsules containing carbon-14 urea each for Δ in vivo Δ diagnostic use for humans to persons exempt from licensing; requirements for a license		NRC			
§32.21a	Same: Conditions of license		NRC			
§32.22			NRC			

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	Self luminous products containing tritium, krypton-85 or promethium-147: Requirements for license to manufacture, process, produce, or initially transfer:					
§32.23	Same: Safety criteria		NRC			
§32.24	Same: Table of organ doses		B			
§32.25	Conditions of licenses issued under §32.22: Quality Control, labeling and reports of transfer		NRC			
§32.26	Gas and aerosol detectors containing byproduct		NRC			

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	material: Requirements for license to manufacture, process, produce or initially transfer					
§32.27	Same: Safety criteria		NRC			
§32.28	Same: Table of organ doses		NRC			
§32.29	Conditions of licenses issued under §32.26: Quality control, labeling and reports of transfer		NRC			
§32.30	Certain industrial devices containing byproduct material: Requirements for license to manufacture, process, produce,		NRC			

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	or initially transfer					
§32.31	Certain industrial devices containing byproduct material: Safety criteria		NRC			
§32.32	Conditions of licenses issued under § 32.30: Quality control, labeling, and reports of transfer		NRC			
§32.51	Byproduct material contained in devices for use under §31.5: Requirements for license to manufacture or initially transfer		B			
§32.51a	Same: Conditions of licenses		B			

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§32.52	Same: Material transfer reports and records		B			
§32.53	Luminous safety devices for use in aircraft: Requirements for license to manufacture, assemble, repair or initially transfer		B			
§32.54	Same: Labeling of devices		B			
32.55	Same: Quality assurance; prohibition of transfer		B			
§32.56	Same: Material transfer reports		B			
§32.57	Calibration or reference sources Am-241 or		B			

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	Ra-226: Requirements for license to manufacture or initially transfer					
§32.58	Same: Labeling of devices		B			
§32.59	Same: Leak testing of each source		B			
§32.60	[Reserved]					
§32.61	Ice detection devices containing strontium-90; Requirements for license to manufacture or initially transfer		B			
§32.62	Same: Quality Assurance; prohibition of transfer		B			

NRC Regulation Section	Section Title	State Section	Compatibility Category	Difference Yes/No	Significant Yes/No	If Difference, Why or Why Not was a Comment Generated
§32.71	Manufacture and distribution of byproduct material for certain in vitro clinical or laboratory testing under general license		B			
§32.72	Manufacture, preparation or transfer for commercial distribution of radioactive drugs containing byproduct material for medical use under Part 35		B			
§32.74	Manufacture and distribution of sources or devices containing byproduct material for medical use		B			

NRC Regulation Section	Section Title	State Section	Compatibility Category	Difference Yes/No	Significant Yes/No	If Difference, Why or Why Not was a Comment Generated
§32.101	Removed (see RATS ID 2012-4)					
§32.102	Removed (see RATS ID 2012-4)					
§32.103	Removed (see RATS ID 2012-4)					
§32.110	Removed (see RATS ID 2012-4)					
§32.201	Serialization of nationally tracked sources		B (only required if a State has a licensee who manufacturers sources with greater than Cat 2 amounts of radioactive material)			
§32.210(a), (b), (c), (d), (e), (f) & (g)	Registration of product information		B- States with authority for sealed source and device			

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			(SS&D) evaluations D- for States that do not perform SS&D evaluations			
§32.210(h)	Registration of product information		C- States with authority for sealed source and device (SS&D) evaluations D- for States that do not perform SS&D evaluations			
§32.211	Inactivation of certificates of registration of sealed sources and devices		B - States with authority for sealed source and device (SS&D) evaluations			

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NRC Regulation Section	Section Title	State Section	Compatibility Category	Difference Yes/No	Significant Yes/No	If Difference, Why or Why Not was a Comment Generated
			D - States without SS&D authority			
§32.301	Violations		D	N/A		
§32.303	Criminal penalties		D	N/A		