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ADD: Tom Pham, Mary Neely Comment (6) Publication Date: 11//08/2021 Citation: 86 FR 61795

Office of Administration Mail Stop: TWFN-7-A60M U.S. Nuclear Regulatory Commission Washington, DC 20555-0001 ATTN: Program Management, Announcements and Editing Staff

Reference: 1) Federal Register, September 23, 2021 (86 FR 52926) 2) SNM, 1257, Docket 70-1257

Subject: Framatome's Comments on Draft NUREG-2159, Docket ID NRC-2021-0170.

Dear Madam or Sir:

Framatome Inc. appreciates the opportunity to comment on NRC's draft NUREG-2159 "Acceptable Standard Format and Content for the Material Control and Accounting Plan Required for Special Nuclear Material of Moderate Strategic Significance: (Reference 1).

Draft NUREG-2159 is intended to provide guidance to facilitate compliance with applicable provisions in Subpart D of 10 CFR Part 74. Framatome is providing comments to clarify, and risk inform sections in the draft NUREG.

Attached are Framatome's specific comments regarding daft NUREG-2159.

If you have any information concerning this information, please contact me at 509 375 8237.

Sincerely,

Calvin D Manning

C D. Manning, Manager Licensing and Compliance

CDM/rd

Cc: R. Jervey, USNRC NMSS

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Attachment

Number	NUREG Section	Comment	Proposed Resolution
1	Section 2, Item 4; 6 Pg 6, lines 8- 18	The proposed guidance far exceeds the requirements found in Subpart D 74.41 (c) as well as the similar requirements found in Subpart E for SSNM and is not a credible concern.	Remove the noted text entirely. Replace with a paragraph stating: "Where
		74.43(b) (1) A management structure shall be established, documented, and maintained that assures:	the organizational structure is such that a position having responsibility within the 1 CFR Part 74 MC&A program also has
		(i) Clear overall responsibility for material control and accounting (MC&A) functions.	responsibility within the 10 CFR Part 73, "Physical Protection of Plants and Materials," system, the plan will address
		(ii) Independence from production and manufacturing responsibilities; and	roles and responsibilities to ensure the MC&A program is not subject to
		(iii) Separation of key responsibilities.	compromise by the actions of a single insider filling the position. A safeguards
		This provides sufficient independence of activities.	manager could be an example of such a position."
2	Section 2.2	For most licensees, such a position would require access to SGI and as such the individual would be granted access to SGI in accordance with 73.22(b) which includes a background check to determine the individual is trustworthy and reliable, and for some licensees such a position would require either an L or Q clearance.	
2	Section 3.3	Section 3.3 "MC&A Organization" states: "An organizational chart and position-by-position description of the entire MC&A organization should be provided. A licensee should designate an	Replace this wording with similar guidance used in NUREG-1520, Section 2.4.2(8)(3) to read:
me Inc.		individual as the overall manager of the MC&A program, and the MC&A plan must demonstrate the assurance of independence of action and objectivity of decision for the MC&A manager. Two options for meeting the organizational independence are: (1) report directly to the plant or site	"In the organizational hierarchy, the MC& organization(s) is independent of the operations organization(s), allowing it to provide objective MC&A audit, review, or

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		manager, or (2) report to an individual who reports to the plant or site manager through a management chain with no production responsibilities." This guidance is not performance based. Other options can easily meet the objectives of the regulation. It is unnecessarily to restrict organization structures and independent reporting options. This could be particularly problematic as advanced reactors look to limit the number of onsite personnel.	control activities. "Independent" means that neither organization reports to the other in an administrative sense. (However, both may report to a common manager.) Lines of responsibility and authority are clearly drawn."
3	Section 3.5	Section 3.5 states one of the roles to be an "overall MC&A program management (note that this individual should have no major responsibilities not related to MC&A)." This guidance is not performance based. Other options can easily meet the objectives of the regulation. Given the industry need to reduce overhead cost, there should be an allowance for MC&A management to have other duties not specifically related to MC&A.	Revise the first bullet to read: "overall MC&A program management (note that overall MC&A program management should be vested to a single individual at an organizational level sufficient to assure independence of action and objectiveness of decisions)."
4	Section 5.1.3; Pg 20, lines 36-39	This Section states that: "All contractor or offsite laboratory assessment findings and recommendations should be documented and submitted to both the measurement control program manager and the overall MC&A manager within 30 days of completion of the review." This time frame is not specified in the regulations and does not appear to be risk informed.	Reword to remove the 30 days and state "Findings and recommendations are to be addressed consistent with the licensees QA and Corrective Action Program."
5	Section 5.2; Pg 21, lines 6-7	This Section states that: "The FNMC plan should describe the replicate sampling program, which must include 6 the following, as appropriate" The language of this sentence mixes requirements, recommendations and provides a conditional out.	Reword sentence to state "FNMCP should address the following applicable attributes:"

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6	Section 6.2;	This Section states that: Reword to state "licensees should		
	Pg 37, lines 28-31	"Licensees should also commit to having at least two individuals independently verify the accuracy of the ID and SEID calculations for each total plant material balance."	commit to independently verify the accuracy of the ID and SEID calculations for each total plant	
7		This recommendation is not performance based. To invoke two additional verifications for such a highly specialized function is an extreme burden on the licensee with no clear regulatory basis or safety benefit.	material balance."	
7	Section 6.3; Pg 38, lines 29-30	This Section states that: "the NRC acceptance criteria do not normally call for applying bias corrections to either the accounting records or as an adjustment to ID unless the effect of a single significant bias or the net sum of all significant biases is unusually large."	Reword or remove "unusually large" or state the following: "The bias correction is only applied if it is significant with 95% confidence and exceeds the rounding error."	
		"unusually large" is an ambiguous term. NRC acceptance criteria does not normally call for applying bias corrections to either the accounting records or as an adjustment.		
8	Section 6.3; Pg 38, line 44	This Section states that: "the bias is greater than 0.01 percent relative"	Reword: "if such bias is statistically significant at the 95 percent confidence level, and exceeds the rounding error of the affected items. The bias correction is	
0	0 11 7 7	This appears to be a new criterion being introduced without a clear tie to regulation.	intended to correct the ID. It's impact on the SEID should also be propagated, resulting in an adjustment to the SEID."	
9	Section 7.5; Pg 44, lines 13-14	This Section states that: "physical inventories are performed so as to confirm that a loss or diversion of a significant quantity of SNM has not occurred: dynamic (i.e., in- process) materials, which is performed on a frequency not to exceed 3 calendar months"	Suggest removing the 3 months and state "at a frequency that takes onto account the potential for loss or diversion from the operation"	
		This requirement is not performance based and there is no direct regulatory basis for the 3 months stated.		

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10	Section 8.2; Pg 51, lines 29-39	This section provides examples of items that may be exempt from item control program coverage. This list should expand to include samples.	Also include the guidance from the previous draft regarding the exemption for laboratory samples and reference standards maintained in the laboratory material management system and containing uranium enriched to less than 20 percent in
11	Section 9.5	 Section 9.5, "Resolution of Significant Shipper-Receiver Differences - Commitments and Acceptance Criteria" states: "Each shipping container is inspected within 3 working days after receipt for loss or damage to the container or T/Ds to determine whether SNM could have been removed. If the integrity of a container is questionable, the presence of all items that were that were packaged in the shipping container will be confirmed within 24 hours of discovering the questionable integrity." 10 CFR 74.43(b)(7) contains neither a 3 working day nor a 24 hour requirement. The 24 hour requirement appears to be overly restrictive and depending on the type of package and the number of items in the package in some cases may not be achievable. 	the uranium-235 isotope. Recommend that this section be reworded to be more consistent with 74.15(a) which requires updated information to be generated within 10 days.