Comment No.	Commenter	Location	Comment	Resolution
1	OAS	Page 3	There is an extra comma in the paragraph which should be removed and Figure 2.1 is missing the Region I Header.	Change accepted. The extra comma has been removed. The "Region I" title was added to Figure 2.1
2	OAS	Section 4.4, Page 16	"Applicable Regulations," includes subsections for select types of devices. The section does not include verbiage for portable gauges or XRF devices. The Board recommends verbiage for these devices be included.	Change not accepted. The sections in Chapter 4 of Volume 3 refer to the different types of devices listed in the regulations in 10 CFR 31 and 32. There is no separate part in 10 CFR 31 or 32 that specifically discusses portable gauges or XRF devices.
3	OAS	Page 35, Section 6.1	States that all document should be printed on standard 8-1/2 inch x 11 inch paper and then Section 6.2 states that drawings should be no larger than 4 inches by 6 inches. The Board recommends deleting the statement from Section 6.2	Change accepted. The reference to limiting the drawing size has been deleted.
4	OAS	Page 43	The third paragraph in Section 9.1 states "an acceptance review may be performed." The Board recommends changing "may" to "will" as it our belief that an acceptance review must be performed when the application is received.	Change was not accepted. Some reviews will not require an acceptance review. Instances where a review would not be required include name change and address changes.
5	OAS	Page 54	The Board recommends including regulation 32.74 into Table 10.2 for medical use design requirements.	The regulation has been added.
6	OAS	Page 57	The Board recommends including regulation 32.74 into Table 10.3 for the labeling requirements of medical sources.	Reference to 10 CFR 32.74 was already included in the last row of the table. No change is required.
7	OAS	Page 60	The Board recommends including regulation 32.74 into Table 10.4 for the prototype testing requirements of medical sources.	The regulation has been added.

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8	OAS	Page 62	The Board recommends including regulations 32.74 and 36.39 for the radiation levels of medical and irradiator sources/devices.	10 CFR 32.74 was added to the table. However, 10 CFR 36.39 was not included as the radiation levels pertain to the irradiator facility and not the sources.
9	OAS	Page 64	The Board recommends including regulations 34.20 and 36.41 for the QA/QC requirements of radiography and irradiator sources/devices.	Change not accepted. 10 CFR 34.20 does not include QA/QC information regarding the manufacturing of sources or devices used in radiography and 10 CFR 36.41 is used for construction monitoring and acceptance testing of an irradiator facility.