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SSINS No.: 6835
IN 86-84

UNITED STATES
NUCLEAR REGULATORY COMMISSION
OFFICE OF INSPECTION AND ENFORCEMENT
WASHINGTON, D.C. 20555

September 30, 1986

~~IS~~ INFORMATION NOTICE NO. 86-84: RUPTURE OF A NOMINAL 40-MILLCURIE IODINE-125 BRACHYTHERAPY SEED CAUSING SIGNIFICANT SPREAD OF RADIOACTIVE CONTAMINATION

Addressees:

All NRC medical institution licensees.

Purpose:

This notice is to alert licensees of a spread of iodine-125 contamination resulting from the inadvertent cutting of the seed encapsulation during removal of the seed from Heyer-Schulte coaxial catheters. The seed was one of eight seeds used for brachytherapy treatment of a brain tumor. It is expected that licensees will review this information for applicability to their facilities and consider actions, if appropriate, to preclude similar problems from occurring at their facilities. However, suggestions contained in this information notice do not constitute NRC requirements; therefore, no specific action or written response is required.

Description of Circumstances:

The seeds, which are manufactured by 3M Company, are intended to be reusable because of the initial high activity of the seeds (40 millicuries per seed). Users of the seeds are motivated to reuse them for several patients because of the relatively high cost of the seeds. The seeds are removed from the old catheters and loaded into new catheters for implant into other patients. It was during removal of the seeds from the catheter that one of the seeds ruptured. The rupture was believed to have been caused by cutting the catheter to free the seeds with a sharp object, such as a razor blade or scissors. The licensee did not know that a seed had been ruptured. The seeds, including the ruptured seed, were reloaded into new catheters and implanted into a patient. As a result, the patient sustained a thyroid burden of about 557 microcuries and a radiation dose to the thyroid of about 2087 rads.

Licensee personnel were not immediately aware that one or more seeds had ruptured until they found iodine-125 contamination in a source/transport bucket stored in the brachytherapy source storage room (BSSR) on the day after the implant into a patient following reloading of the seeds into the new catheters.

The seeds were removed from the old catheters and loaded into new catheters in the BSSR an area not ventilated by a fume hood. Consequently, approximately 60 hospital personnel, including those involved in cleanup operations, received thyroid uptakes of iodine-125 from 0.04 to 209 nanocuries. A proper radiation survey meter was used, but the high background radiation in the BSSR masked the positive indication of contamination.

Discussion:

The isolated incident described is the only incident of its kind known by the NRC involving high-activity iodine-125 seeds. However, there have been several other similar incidents involving the use of low-activity seeds containing 0.1 to 1 millicuries used as permanent implants. Both types of seeds are manufactured by 3M Company.

The risk of iodine-125 seed rupture is relatively high when the seeds are used for several patients. The incident was caused because:

- . the seeds are susceptible to damage from sharp tools such as razor blades or scissors
- . the discolored or stained condition of the catheters after use in therapy makes viewing of the seeds difficult

The consequences of the seed rupture (involving patient exposures, other personnel iodine-125 uptakes, and facility contamination) can be mitigated by performing (1) adequate radiation surveys of the work area, using a radiation detection instrument capable of detecting the low-energy photons (average of 28 keV), (2) the proper handling of tools used to remove the seeds from the catheters, and (3) leak tests of the seeds. Also, contamination of the facilities probably can be prevented if the seed removal operation is performed in a fume hood. Personnel uptakes of the radioactive materials and facility contamination also might be mitigated by using radiation safety procedures designed to detect seed leakage promptly.

No specific action or written response is required by this notice. If you have any questions regarding this information notice, please contact the Regional Administrator of the appropriate NRC regional office or this office.



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