August 15, 2013

ALL AGREEMENT STATES

DISTRIBUTION OF NRC INFORMATIONAL DOCUMENT, RIS-2013-10 (FSME-13-079)

Purpose: To inform the Agreement States of the U.S. Nuclear Regulatory Commission's (NRC) issuance of a Regulatory Issue Summary (RIS), "Permanent Implant Brachytherapy Medical Event Reporting Under 10 CFR Part 35." As discussed during a recent teleconference with the States, the NRC is providing this RIS to Agreement States for their information and for distribution to their medical licensees, as appropriate. You may find the RIS online at: http://pbadupws.nrc.gov/docs/ML1222/ML12228A606.pdf

Background: This RIS represents one of two measures involving medical event (ME) reporting that, in Staff Requirements Memorandum SRM-SECY-12-0053, the Commission directed staff to pursue. The purpose of the RIS is to clarify ME reporting requirements under the existing NRC rules.

Discussion: This RIS informs licensees about NRC's expectations for compliance with the current NRC requirements related to permanent implant brachytherapy. The RIS also announces that an Interim Enforcement Policy (IEP) has been developed and published. The IEP explains the enforcement discretion that NRC may provide to licensees until a new rule for ME reporting requirements is finalized. You may find the IEP online at: http://www.regulations.gov/#!documentDetail;D=NRC-2013-0114-0001

If you have any questions regarding this correspondence, please contact me at (301) 415-3340 or the individual named below.

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