

SUBJECT:

UNITED STATES NUCLEAR REGULATORY COMMISSION WASHINGTON, D.C. 20555-0001

February 24, 2020

IN RESPONSE, PLEASE REFER TO: M200128

MEMORANDUM TO:	Margaret M. Doane Executive Director for Operations
FROM:	Annette L. Vietti-Cook, Secretary

STAFF REQUIREMENTS – DISCUSSION OF MEDICAL USES OF RADIOACTIVE MATERIALS, 9:00 A.M., TUESDAY, JANUARY 28, 2020, COMMISSIONERS' CONFERENCE ROOM, ONE WHITE FLINT NORTH, ROCKVILLE, MARYLAND (OPEN TO PUBLIC ATTENDANCE)

The Commission was briefed by the NRC staff and external stakeholders on the NRC's program for medical uses of radioactive materials. The NRC staff presentations included: an overview of the NRC's program for medical uses of radioactive materials, a status of recent NRC staff activities, innovation opportunities and initiatives, efforts to prepare for the review of emerging medical technologies, and a Regional perspective on licensing and oversight of medical licensees. A representative from the U.S. Food and Drug Administration, the American Society for Radiation Oncology, and NorCal CarciNET Community provided federal, medical community, and patient suggestions, respectively, regarding transformation and innovation opportunities for the NRC to explore. The Chair of the Organization of Agreement States discussed emerging issues regarding the national program for the regulation of medical uses of radioactive materials. Additionally, the President of the Society of Nuclear Medicine and Molecular Imaging provided perspectives on recent NRC staff initiatives related to medical uses of radioactive materials.

There were no requirements identified for staff action.

cc: Chairman Svinicki Commissioner Baran Commissioner Caputo Commissioner Wright OGC CFO OCA OIG OPA ODs, RAs, ACRS, ASLBP (via E-Mail) PDR