

IN RESPONSE, PLEASE
REFER TO: M930624D

August 5, 1993

MEMORANDUM TO: James M. Taylor
Executive Director for Operations

FROM: Samuel J. Chilk, Secretary /s/

SUBJECT: STAFF REQUIREMENTS - BRIEFING ON INTERNAL
MANAGEMENT REVIEW OF NRC PROGRAM FOR MEDICAL
USE OF BYPRODUCT MATERIAL, 2:00 P.M.,
THURSDAY, JUNE 24, 1993, COMMISSIONERS'
CONFERENCE ROOM, ONE WHITE FLINT NORTH,
ROCKVILLE, MARYLAND (OPEN TO PUBLIC
ATTENDANCE)

The Commission* was briefed by the NRC staff on the internal management review of the NRC program for medical use of byproduct material.

The Commission advised the staff to ensure that regulatory burdens placed on licensees are not just additive, but that consideration is also given to replacing existing requirements, and to work for consistency in areas where the NRC may differ from the States and where revisions related to the medical use area may impact other materials areas. The staff should proceed with recommended actions to ensure the Quality Management program is implemented by licensees, with development of a Management Directive for followup on therapeutic misadministration reports and with other recommended actions. The staff should be realistic in estimating associated resource requirements.

There were no specific requirements identified for action.

cc: The Chairman
Commissioner Rogers
Commissioner Remick
Commissioner de Planque
OGC
OIG
OCA
Office Directors, Regions, ACRS, ACNW, ASLBP (via E-Mail)
PDR - Advance
DCS - P1-24

* Commissioner Rogers was on leave and did not attend this briefing.