Date: February 14, 2002

DRAFT SUMMARY OF INFORMATION COLLECTION REQUEST

Title: NRC Form 483, "Registration Certificate - In Vitro Testing with Byproduct Material under General License"

Current Burden/Responses: 42 hours/364 responses

Proposed Burden/Responses: 42 hours/364 responses

Burden Attributable to Third-Party Collections: 30 hours

Frequency of Response: One time

Reasons for Changes in Burden/Responses: The overall burden estimate for NRC Form 241 has remained the same.

Level of Concurrence: Chief, Material Safety Branch
Division of Industrial and Medical Nuclear Safety
Office of Nuclear Material Safety and Safeguards

Recordkeeping Requirements in Accordance with the Retention Periods for Records Rule: N/A

Search of the Information Requirements Control Automated System (IRCAS):
IRCAS was searched on February 12, 2002. No duplication was found.

Abstract: Section 31.11 of 10 CFR establishes a general license authorizing any physician, veterinarian, clinical laboratory or hospital to possess certain small quantities of byproduct material for in vitro clinical or laboratory tests not involving humans or animals. Possession of byproduct under 10 CFR 31.11 is not authorized until an NRC Form 483 is filed and validated with a registration number from the NRC.

cc: B. St. Mary