June 6, 2005

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: None

Frequency of Response: One time

Reasons for Changes in Burden/Responses: Although, the overall burden estimate for NRC Form 483 has remained the same; the cost estimates have changed since the last clearance, as a result of an increase in the fee per hour from $144 to $197/hour.

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:
Not Applicable.

Search of the Information Requirements Control Automated System (IRCAS):
IRCAS was searched and no duplication was found

Abstract: Section 31.11 of 10 CFR establishes a general license authorizing any physician, clinician, clinical laboratory, veterinarian in the practice of veterinary medicine, or hospital to possess certain small quantities of byproduct material for in vitro clinical or laboratory test not involving the internal or external administration of the byproduct material or the radiation there from to human beings or animals. Possession of byproduct material under 10 CFR 31.11 is not authorized until the physician, clinical laboratory, veterinarian in the practice of veterinary medicine, or hospital has filed NRC Form 483 and received from the Commission a validated copy on NRC Form 483 with a registration number.

cc: Chris Colburn
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