[7590-01-P]

NUCLEAR REGULATORY COMMISSION

[Docket No. NRC-2014-0155]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Nuclear Regulatory Commission.

ACTION: Notice of pending NRC action to submit an information collection request to the Office of Management and Budget (OMB) and solicitation of public comment.

SUMMARY: The NRC invites public comment about our intention to request the OMB's approval for renewal of an existing information collection that is summarized below. We are required to publish this notice in the *Federal Register* under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

Information pertaining to the requirement to be submitted:

1. *The title of the information collection:* NRC Form 483, "Registration Certificate –*In Vitro* Testing with Byproduct Material Under General License."

2. *Current OMB approval number:* 3150-0038.

3. *How often the collection is required:* There is a one-time submittal of information to receive a validated copy of NRC Form 483 with an assigned registration number. In addition, any changes in the information reported on NRC Form 483 must be reported in writing to the NRC within 30 days after the effective date of such change.

4. *Who is required or asked to report:* Any physician, veterinarian in the practice of veterinary medicine, clinical laboratory or hospital which desires a general license to receive, acquire, possess, transfer, or use specified units of byproduct material in certain *in vitro* clinical or laboratory tests.

5. *The number of annual respondents:* 8 respondents.

6. *The number of hours needed annually to complete the requirement or request:* 1.18 hours (1.07 hours reporting + 0.11 hour recordkeeping).

7. *Abstract:* Section 31.11 of Title 10 of the *Code of Federal Regulations* (10 CFR) establishes a general license authorizing any physician, 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 therefrom 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 of NRC Form 483 with a registration number.

Submit, by **(INSERT DATE 60 DAYS AFTER PUBLICATION IN THE *FEDERAL REGISTER***), comments that address the following questions:

1. Is the proposed collection of information necessary for the NRC to properly perform its functions? Does the information have practical utility?

2. Is the burden estimate accurate?

3. Is there a way to enhance the quality, utility, and clarity of the information to be collected?

4. How can the burden of the information collection be minimized, including the use of automated collection techniques or other forms of information technology?

The public may examine and have copied for a fee publicly-available documents, including the draft supporting statement, at the NRC's Public Document Room, Room O-1F21, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852. The OMB clearance requests are available at the NRC's Web site: http://www.nrc.gov/public-involve/doc-comment/omb/. The document will be available on the NRC's home page site for 60 days after the signature date of this notice.

Comments submitted in writing or in electronic form will be made available for public inspection. Because your comments will not be edited to remove any identifying or contact information, the NRC cautions you against including any information in your submission that you do not want to be publicly disclosed. Comments submitted should reference Docket No. NRC-2014-0155. You may submit your comments by any of the following methods: Electronic comments go to http://www.regulations.gov and search for Docket No. NRC-2014-0155. Mail comments to the Acting NRC Clearance Officer, Brenda Miles (T-5 F44), U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001.

Questions about the information collection requirements may be directed to the Acting NRC Clearance Officer, Brenda Miles (T-5 F44), U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001, telephone: 301-415-7884, or by e-mail to INFOCOLLECTS.Resource@NRC.GOV.

Dated at Rockville, Maryland, this 15th day of July, 2014.

For the Nuclear Regulatory Commission.

Brenda Miles,
Acting NRC Clearance Officer,
Office of Information Services.

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