the availability of information for this action. You may obtain publiclyavailable information related to this action by any of the following methods:

- Federal Rulemaking Web site: Go to http://www.regulations.gov and search for Docket ID NRC-2017-0067.
- NRC's Agencywide Documents Access and Management System (ADAMS): You may obtain publiclyavailable documents online in the ADAMS Public Documents collection at http://www.nrc.gov/reading-rm/ adams.html. To begin the search, select "ADAMS Public Documents" and then select "Begin Web-based ADAMS Search." For problems with ADAMS, please contact the NRC's Public Document Room (PDR) reference staff at 1-800-397-4209, 301-415-4737, or by email to pdr.resource@nrc.gov. The supporting statement is available in ADAMS under Accession No. ML17191B158.
- NRC's PDR: You may examine and purchase copies of public documents at the NRC's PDR, Room O1–F21, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852.
- NRC's Clearance Officer: A copy of the collection of information and related instructions may be obtained without charge by contacting the NRC's Clearance Officer, David Cullison, Office of the Chief Information Officer, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001; telephone: 301–415–2084; email: INFOCOLLECTS.Resource@NRC.GOV.

## B. Submitting Comments

The NRC cautions you not to include identifying or contact information that you do not want to be publicly disclosed in your comment submission. The NRC posts all comment submissions at <a href="http://www.regulations.gov">http://www.regulations.gov</a> as well as entering the comment submissions into ADAMS. The NRC does not routinely edit comment submissions to remove identifying or contact information.

If you are requesting or aggregating comments from other persons for submission to the OMB, then you should inform those persons not to include identifying or contact information that they do not want to be publicly disclosed in their comment submission. Your request should state that the NRC does not routinely edit comment submissions to remove such information before making the comment submissions available to the public or entering the comment submissions into ADAMS.

#### II. Background

Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the NRC recently submitted a request for renewal of an existing collection of information to OMB for review entitled, "Licensing Requirements for Land Disposal of Radioactive Waste." The NRC hereby informs potential respondents that an agency may not conduct or sponsor, and that a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

The NRC published a **Federal Register** notice with a 60-day comment period on this information collection on May 10, 2017 (82 FR 21834).

- 1. The title of the information collection: 10 CFR part 61—Licensing Requirements for Land Disposal of Radioactive Waste.
  - 2. OMB approval number: 3150-0135.
  - 3. Type of submission: Extension.
- 4. The form number if applicable: Not applicable.
- 5. How often the collection is required or requested: Applications for licenses are submitted as needed. Other reports are submitted annually and as other events require.
- 6. Who will be required or asked to respond: Applicants for and holders of an NRC license (to include Agreement State licensees) for land disposal of low-level radioactive waste.
- 7. The estimated number of annual responses: 16 (12 reporting responses + 4 recordkeepers).
- 8. The estimated number of annual respondents: 4.
- 9. An estimate of the total number of hours needed annually to comply with the information collection requirement or request: 5,372 hours (56 hours reporting + 5,316 hours recordkeeping).
- 10. Abstract: Part 61 of title 10 of the Code of Federal Regulations (10 CFR), establishes the procedures, criteria, and license terms and conditions for the land disposal of low-level radioactive waste. The reporting and recordkeeping requirements are mandatory and, in the case of application submittals, are required to obtain a benefit. The information collected in the applications, reports, and records is evaluated by the NRC to ensure that the licensee's or applicant's disposal facility, equipment, organization, training, experience, procedures, and plans provide an adequate level of protection of public health and safety, common defense and security, and the environment.

Dated at Rockville, Maryland, this 22nd day of August, 2017.

For the Nuclear Regulatory Commission. **David Cullison**,

NRC Clearance Officer, Office of the Chief Information Officer.

[FR Doc. 2017–18141 Filed 8–25–17; 8:45 am] BILLING CODE 7590–01–P

# NUCLEAR REGULATORY COMMISSION

[NRC-2017-0166]

Information Collection: NRC Form 483, Registration Certificate—In Vitro Testing with Byproduct Material Under General License

**AGENCY:** Nuclear Regulatory Commission.

**ACTION:** Renewal of existing information collection; request for comment.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) invites public comment on the renewal of Office of Management and Budget (OMB) approval for an existing collection of information. The information collection is entitled, "NRC Form 483, Registration Certificate—In Vitro Testing with Byproduct Material Under General License." NRC Form 483 will be revised to update instructions and regulatory language.

**DATES:** Submit comments by October 27, 2017. Comments received after this date will be considered if it is practical to do so, but the Commission is able to ensure consideration only for comments received on or before this date.

**ADDRESSES:** You may submit comments by any of the following methods:

- Federal Rulemaking Web site: Go to http://www.regulations.gov and search for Docket ID NRC-2017-0166. Address questions about NRC dockets to Carol Gallagher; telephone: 301-415-3463; email: Carol.Gallagher@nrc.gov. For technical questions, contact the individual listed in the FOR FURTHER INFORMATION CONTACT section of this document.
- Mail comments to: David Cullison, Office of the Chief Information Officer, Mail Stop: T–5 F53, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001.

For additional direction on obtaining information and submitting comments, see "Obtaining Information and Submitting Comments" in the SUPPLEMENTARY INFORMATION section of this document.

#### FOR FURTHER INFORMATION CONTACT:

David Cullison, Office of the Chief Information Officer, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001; telephone: 301–415– 2084; email: INFOCOLLECTS.Resource@NRC.GOV.

#### SUPPLEMENTARY INFORMATION:

# I. Obtaining Information and Submitting Comments

#### A. Obtaining Information

Please refer to Docket ID NRC–2017– 0166 when contacting the NRC about the availability of information for this action. You may obtain publiclyavailable information related to this action by any of the following methods:

- Federal Rulemaking Web site: Go to http://www.regulations.gov and search for Docket ID NRC-2017-0166. A copy of the collection of information and related instructions may be obtained without charge by accessing Docket ID NRC-2017-0166 on this Web site.
- NRC's Agencywide Documents Access and Management System (ADAMS): You may obtain publiclyavailable documents online in the ADAMS Public Documents collection at http://www.nrc.gov/reading-rm/ adams.html. To begin the search, select "ADAMS Public Documents" and then select "Begin Web-based ADAMS Search." For problems with ADAMS, please contact the NRC's Public Document Room (PDR) reference staff at 1-800-397-4209, 301-415-4737, or by email to pdr.resource@nrc.gov. A copy of the collection of information and related instructions may be obtained without charge by accessing ADAMS under Accession No. ML17128A454. The supporting statement is available in ADAMS under Accession No. ML17128A131.
- NRC's PDR: You may examine and purchase copies of public documents at the NRC's PDR, Room O1–F21, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852.
- NRC's Clearance Officer: A copy of the collection of information and related instructions may be obtained without charge by contacting NRC's Clearance Officer, David Cullison, Office of the Chief Information Officer, U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001; telephone: 301-415-2084; email: INFOCOLLECTS.Resource@ NRC.GOV.

#### B. Submitting Comments

Please include Docket ID NRC–2017–0166 in the subject line of your comment submission, in order to ensure that the NRC is able to make your comment submission available to the public in this docket.

The NRC cautions you not to include identifying or contact information that you do not want to be publicly disclosed in your comment submission.

The NRC posts all comment submissions at http://www.regulations.gov as well as entering the comment submissions into ADAMS. The NRC does not routinely edit

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## II. Background

In accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the NRC is requesting public comment on its intention to request the OMB's approval for the information collection summarized below:

- 1. The title of the information collection: NRC Form 483, Registration Certificate—In Vitro Testing With Byproduct Material Under General License.
  - 2. OMB approval number: 3150-0038.
- 3. *Type of submission:* Revision.
- 4. The form number, if applicable: NRC Form 483.
- 5. How often the collection is required or requested: 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 the change.
- 6. Who will be required or asked to respond: 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.
- 7. The estimated number of annual responses: 6.
- 8. The estimated number of annual respondents: 6.
- 9. The estimated number of hours needed annually to comply with the information collection requirement or request: 1.10 hours.
- 10. Abstract: Section 31.11 of title 10 of the Code of Federal Regulations (10 CFR), established 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. The licensee can use the validated copy of NRC Form 483 to obtain byproduct material from a specifically licensed supplier. The NRC incorporates this information into a database which is used to verify that a general licensee is authorized to receive the byproduct material.

#### **III. Specific Requests for Comments**

The NRC is seeking 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 estimate of the burden of the information collection 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 on respondents be minimized, including the use of automated collection techniques or other forms of information technology?

Dated at Rockville, Maryland, this 23rd day of August, 2017.

For the Nuclear Regulatory Commission. **David Cullison**,

NRC Clearance Officer, Office of the Chief Information Officer.

[FR Doc. 2017–18147 Filed 8–25–17; 8:45 am]

# PENSION BENEFIT GUARANTY CORPORATION

Agency Information Collection Activities: Submission of Information Collection for OMB Review; Comment Request; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

**AGENCY:** Pension Benefit Guaranty Corporation.

**ACTION:** Notice of request for extension of OMB approval.