DEPARTMENT OF LABOR

Occupational Safety and Health Administration

[Docket No. OSHA–2007–0083]

Applied Research Laboratories of South Florida, LLC; Voluntary Termination of Recognition as a Nationally Recognized Testing Laboratory

AGENCY: Occupational Safety and Health Administration (OSHA), Labor.

ACTION: Notice.

SUMMARY: In this notice, OSHA announces the voluntary termination of recognition granted to Applied Research Laboratories of South Florida, LLC as a Nationally Recognized Testing Laboratory (NRTL).

DATES: The voluntary termination of recognition specified by this notice becomes effective on August 1, 2023.

FOR FURTHER INFORMATION CONTACT: Information regarding this notice is available from the following sources:

Press inquiries: Contact Mr. Frank Meilinger, Director, OSHA Office of Communications, U.S. Department of Labor; telephone: (202) 693–1999; email: *meilinger.francis2@dol.gov.*

General and technical information: Contact Mr. Kevin Robinson, Director, Office of Technical Programs and Coordination Activities, Directorate of Technical Support and Emergency Management, Occupational Safety and Health Administration, U.S. Department of Labor; telephone: (202) 693–2110; email: *robinson.kevin@dol.gov.*

SUPPLEMENTARY INFORMATION:

I. Notice of Voluntary Termination of Recognition as a Nationally Recognized Testing Laboratory

On April 25, 2018, OSHA granted Applied Research Laboratories of South Florida, LLC (ARL) recognition as a Nationally Recognized Testing Laboratory (NRTL). The NRTL Program regulation, 29 CFR 1910.7 App. A, provides that, ''[a]t any time, a recognized NRTL may voluntarily terminate its recognition, either in its entirety or with respect to any area covered in its recognition, by giving written notice to OSHA,'' that ''[t]he written notice shall state the date as of which the termination is to take effect, and that [t]he Assistant Secretary shall inform the public of any voluntary termination by **Federal Register** notice.''

ARL notified OSHA by a letter dated April 19, 2023 (OSHA–2007–0083–0057), that it would be closing its business effective July 31, 2023. In a subsequent email, dated May 25, 2023 (OSHA–2007–0038–0058), ARL confirmed that this letter constituted a voluntary and complete termination of ARL's recognition as a NRTL, effective August 1, 2023, as well as a voluntary withdrawal of any open NRTL scope expansions and NRTL renewal applications previously submitted to OSHA, also effective August 1, 2023.

Therefore, pursuant to the NRTL Program regulation, OSHA is informing the public that ARL has voluntary terminated its recognition.

II. Authority and Signature

James S. Frederick, Deputy Assistant Secretary of Labor for Occupational Safety and Health, 200 Constitution Avenue NW, Washington, DC 20210, authorized the preparation of this notice. Accordingly, the agency is issuing this notice pursuant to Section 29 U.S.C. 657(g)(2), Secretary of Labor's Order No. 8–2020 (85 FR 58393, Sept. 18, 2020), and 29 CFR 1910.7.

Signed at Washington, DC, on July 24, 2023.

James S. Frederick,

Deputy Assistant Secretary of Labor for Occupational Safety and Health.

[FR Doc. 2023–16131 Filed 7–31–23; 8:45 am]

BILLING CODE 4510–26–P

NUCLEAR REGULATORY COMMISSION

[NRC–2023–0065]

Information Collection: Medical Use of Byproduct Material

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, ''Medical Use of Byproduct Material.''

DATES: Submit comments by October 2, 2023. 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; however, the NRC encourages electronic comment submission through the Federal rulemaking website:

- *Federal Rulemaking Website:* Go to *https://www.regulations.gov* and search for Docket ID NRC–2023–0065. Address questions about Docket IDs in *Regulations.gov* to Stacy Schumann; telephone: 301–415–0624; email: *Stacy.Schumann@nrc.gov.* For technical questions, contact the individual listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.
- *Mail comments to:* David C. Cullison, Office of the Chief Information Officer, Mail Stop: T–6 A10M, 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 C. 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–2023–0065 when contacting the NRC about the availability of information for this action. You may obtain publicly available information related to this action by any of the following methods:

- *Federal Rulemaking Website:* Go to *https://www.regulations.gov* and search for Docket ID NRC–2023–0065.
- *NRC's Agencywide Documents Access and Management System (ADAMS):* You may obtain publicly available documents online in the ADAMS Public Documents collection at *https://www.nrc.gov/reading-rm/adams.html.* To begin the search, 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, at 301–415–4737, or by email to *PDR.Resource@nrc.gov.* The supporting statement and burden spreadsheet are available in ADAMS under Accession Nos. ML23137A012 and ML23136B177.
- *NRC's PDR:* The PDR, where you may examine and order copies of publicly available documents, is open by appointment. To make an appointment to visit the PDR, please send an email to *PDR.Resource@nrc.gov* or call 1–800–397–4209 or 301–415–4737, between 8 a.m. and 4 p.m. eastern time (ET), Monday through Friday, except Federal holidays.
- *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 C. 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 encourages electronic comment submission through the Federal rulemaking website (*https://www.regulations.gov*). Please include Docket ID NRC–2023–0065, in your comment submission.

The NRC cautions you not to include identifying or contact information in comment submissions that you do not want to be publicly disclosed in your comment submission. All comment submissions are posted at *https://www.regulations.gov* and entered into ADAMS. Comment submissions are not routinely edited to remove identifying or contact information.

If you are requesting or aggregating comments from other persons for submission to the NRC, 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 comment submissions are not routinely edited to remove such information before making the comment submissions available to the public or entering the comment into ADAMS.

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:* 10 CFR part 35, Medical Use of Byproduct Material.

2. *OMB approval number:* 3150–0010.

3. *Type of submission:* Revision.

4. *The form number, if applicable:* Not applicable.

5. *How often the collection is required or requested:* Reports of medical events, doses to an embryo/fetus or nursing child, or leaking source are reportable on occurrence. A specialty board certifying entity desiring to be recognized by the NRC must submit a one-time request for recognition and infrequently revise the information.

6. *Who will be required or asked to respond:* Physicians and medical institutions holding an NRC license authorizing the administration of byproduct material or radiation from this material to humans for medical use. A specialty board certification entity desiring to have its certifying process and board certificate recognized by NRC.

7. *The estimated number of annual responses:* 313,982 (234,272 reporting responses + 7,327 recordkeepers + 72,383 third party disclosure responses).

8. *The estimated number of annual respondents:* 7,328 (862 NRC licensees + 6,465 Agreement State licensees + 1 specialty board certification entity).

9. *The estimated number of hours needed annually to comply with the information collection requirement or request:* 1,117,269 hours (61,506 reporting + 1,043,235 recordkeeping + 12,528 third party disclosure).

10. *Abstract:* Part 35 of title 10 of the *Code of Federal Regulations,* ''Medical Use of Byproduct Material,'' contains NRC's requirements and provisions for the medical use of byproduct material and for issuance of specific licenses authorizing the medical use of this material. These requirements and provisions provide for the radiation safety of workers, the general public, patients, and human research subjects. Part 35 contains mandatory requirements that apply to NRC licensees authorized to administer byproduct material or radiation to humans for medical use. These requirements also provide voluntary provisions for specialty boards to apply to have their certification processes recognized by the NRC so that their board-certified individuals can use the certifications as proof of training and experience.

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? Please explain your answer.

2. Is the estimate of the burden of the information collection accurate? Please explain your answer.

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: July 27, 2023.

For the Nuclear Regulatory Commission.

David C. Cullison,

NRC Clearance Officer, Office of the Chief Information Officer.

[FR Doc. 2023–16256 Filed 7–31–23; 8:45 am]

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NUCLEAR REGULATORY COMMISSION

[NRC–2023–0122]

Applications for Amendments to Facility Operating Licenses Involving Proposed No Significant Hazards Consideration(s) and Containing Sensitive Unclassified Non-Safeguards Information and Order Imposing Procedures for Access to Sensitive Unclassified Non-Safeguards Information

AGENCY: Nuclear Regulatory Commission.

ACTION: License amendment request; notice of opportunity to comment, request a hearing, and petition for leave to intervene; order imposing procedures.

SUMMARY: The U.S. Nuclear Regulatory Commission (NRC) received and is considering approval of two amendment requests. The amendment requests are for Millstone Power Station, Unit 3. For each amendment request, the NRC proposes to determine that it involves no significant hazards consideration (NSHC). Because each amendment request contains sensitive unclassified non-safeguards information (SUNSI), an order imposes procedures to obtain access to SUNSI for contention preparation by persons who file a hearing request or petition for leave to intervene.

DATES: Comments must be filed by August 31, 2023. A request for a hearing or petitions for leave to intervene must be filed by October 2, 2023. Any potential party as defined in section 2.4 of title 10 of the *Code of Federal Regulations* (10 CFR) who believes access to SUNSI is necessary to respond to this notice must request document access by August 11, 2023.

ADDRESSES: You may submit comments by any of the following methods; however, the NRC encourages electronic comment submission through the Federal rulemaking website:

- *Federal rulemaking website:* Go to *https://www.regulations.gov* and search for Docket ID NRC–2023–0122. Address questions about Docket IDs in *Regulations.gov* to Stacy Schumann; telephone: 301–415–0624; email: *Stacy.Schumann@nrc.gov.* For technical