Federal Regulations (10 CFR) the NRC is providing notice of the receipt of the application; providing the opportunity to submit written comments concerning the application; and providing the opportunity to request a hearing or petition for leave to intervene, for a period of 30 days after publication of this notice in the **Federal Register**.

A hearing request or petition for leave to intervene must include the information specified in 10 CFR 110.82(b). Any request for hearing or petition for leave to intervene shall be served by the requestor or petitioner in accordance with 10 CFR 110.89(a), either by delivery, by mail, or filed with the NRC electronically in accordance with the NRC's E-Filing rule (72 FR 49139; August 28, 2007, as amended at 77 FR 46562; August 3, 2012). Detailed guidance on making electronic submissions may be found in the Guidance for Electronic Submissions to the NRC and on the NRC website at *https://www.nrc.gov/site-help/e-submittals.html*.

To comply with the procedural requirements of E-Filing, at least 10 days prior to the filing deadline, the participant should contact the Office of the Secretary by email at *Hearing.Docket@nrc.gov*, or by telephone at 301–415–1677, to (1) request a digital identification (ID) certificate, which allows the participant

## the E-Filing system for any proceeding in which it is participating; and (2) advise the Secretary that the participant will be submitting a petition or other adjudicatory document (even in instances in which the participant, or its counsel or representative, already holds an NRC-issued digital ID certificate). Based upon this information, the Secretary will establish an electronic docket for the hearing in this proceeding if the Secretary has not already established an electronic docket. The information concerning this

(or its counsel or representative) to

digitally sign submissions and access

application for an export license follows:

## NRC EXPORT LICENSE APPLICATION

Application Information	
Name of Ap- plicant.	Perma-Fix Northwest Richland, Inc. (PFNW).
Date of Appli- cation.	October 12, 2022.
Date Received	October 17, 2022.
Application No	XW027/01.
Docket No	11006380.
ADAMS Ac-	ML22292A007.
cession No.	
Description of Material	
Material Type	The incoming material from Germany consists of liquid, shredded and combustible material produced from research, medical, and other industries (excluding nuclear power plants). After treatment by PFNW, radioactive waste returned to Germany will consist of residual ash and residual metal or non-combustible material that cannot be recycled. Radionuclides in the waste include carbon-14, cesium-137, cobalt-60, nickel-63, radium-226, and strontium-90.
Total Quantity	Authorization to export a total maximum quantity of waste will not exceed 800,000 kilograms. The maximum activity returned to the originating Eckert and Ziegler Nuclitech GmbH facility will not exceed 8.5 TBq.
End Use	Disposal in Germany.
Country of	Germany.
Destination.	

Dated: November 17, 2022.

For the Nuclear Regulatory Commission. Peter J. Habighorst,

Acting Deputy Director, Office of International Programs. [FR Doc. 2022–25376 Filed 11–21–22; 8:45 am]

BILLING CODE 7590-01-P

# NUCLEAR REGULATORY COMMISSION

# Advisory Committee on the Medical Uses of Isotopes: Call for Nominations

**AGENCY:** U.S. Nuclear Regulatory Commission.

**ACTION:** Call for Nominations.

**SUMMARY:** The U.S. Nuclear Regulatory Commission (NRC) is soliciting nominations for the position of Radiation Oncologist Physician (Brachytherapy) on the Advisory Committee on the Medical Uses of Isotopes (ACMUI). Nominees should be currently practicing radiation oncologists.

**DATES:** Nominations are due on or before January 23, 2023.

Nomination Process: Submit an electronic copy of resume or curriculum vitae, along with a cover letter, to Dr. Celimar Valentin-Rodriguez, celimar.valentin-rodriguez@nrc.gov. The cover letter should describe the nominee's current duties and responsibilities and express the nominee's interest in the position. Please ensure that the resume or curriculum vitae includes the following information, if applicable: education; certification(s); professional association and committee membership activities; duties and responsibilities in current and previous clinical, research, and/or academic position(s).

# FOR FURTHER INFORMATION CONTACT: $\ensuremath{\mathrm{Dr}}.$

Celimar Valentin-Rodriguez, U.S. Nuclear Regulatory Commission, Office of Nuclear Material Safety and Safeguards; (301) 415–7124; *celimar.valentin-rodriguez@nrc.gov.*  SUPPLEMENTARY INFORMATION: ACMUI members possess the medical and technical skills needed to address evolving issues. The current membership is comprised of the following professionals: (a) nuclear medicine physician; (b) nuclear cardiologist; (c) two radiation oncologists; (d) diagnostic radiologist; (e) therapy medical physicist; (f) nuclear medicine physicist; (g) nuclear pharmacist; (h) health care administrator; (i) radiation safety officer; (j) patients' rights advocate; (k) Food and Drug Administration representative; and (l) Agreement State representative. For additional information about membership on the ACMUI, visit the ACMUI Membership web page, http:// www.nrc.gov/about-nrc/regulatory/ advisorv/acmui/membership.html.

The ACMUI brachytherapy radiation oncologist provides advice on issues associated with radiation oncology and the clinical use of brachytherapy, including the use of permanently implanted microspheres. This advice includes providing input on NRC proposed rules and guidance, providing recommendations on the training and experience requirements for physicians specializing in this use, identifying medical events associated with this use, evaluating non-routine uses of byproduct material and emerging medical technologies, bringing key issues in the radiation oncology community to the attention of NRC staff, and other radiation oncology issues as they relate to radiation safety and NRC medical use policy.

The ACMUI advises the NRC on policy and technical issues that arise in the regulation of the medical use of byproduct material. Responsibilities of an ACMUI member include providing comments on changes to the NRC regulations and guidance; evaluating certain non-routine uses of byproduct material; providing technical assistance in licensing, inspection, and enforcement cases; and bringing key issues to the attention of the NRC staff, for appropriate action. Committee members currently serve a four-year term and may be considered for reappointment to an additional term.

Nominees must be U.S. citizens and be able to devote approximately 160 hours per year to ACMUI business. Members are expected to attend semiannual meetings at NRC headquarters in Rockville, Maryland and to participate in teleconferences or virtual meetings, as needed. Members who are not Federal employees at the time of their appointment are compensated for their service. In addition, members are reimbursed for travel (including per diem in lieu of subsistence) and are reimbursed secretarial and correspondence expenses. Full-time Federal employees are reimbursed for travel expenses only.

Security Background CHECK: The selected nominee will undergo a thorough security background check. Security paperwork may take the nominee several weeks to complete. Nominees will also be required to complete a financial disclosure statement to avoid conflicts of interest.

Dated at Rockville, Maryland, this 17th day of November, 2022.

For the U.S. Nuclear Regulatory Commission.

## Brooke P. Clark,

Secretary of the Commission. [FR Doc. 2022–25403 Filed 11–21–22; 8:45 am] BILLING CODE 7590–01–P

# POSTAL REGULATORY COMMISSION

[Docket Nos. MC2023-47 and CP2023-45]

### **New Postal Products**

**AGENCY:** Postal Regulatory Commission. **ACTION:** Notice.

**SUMMARY:** The Commission is noticing a recent Postal Service filing for the Commission's consideration concerning a negotiated service agreement. This notice informs the public of the filing, invites public comment, and takes other administrative steps.

**DATES:** *Comments are due:* November 23, 2022.

ADDRESSES: Submit comments electronically via the Commission's Filing Online system at *http:// www.prc.gov.* Those who cannot submit comments electronically should contact the person identified in the FOR FURTHER INFORMATION CONTACT section by telephone for advice on filing alternatives.

# FOR FURTHER INFORMATION CONTACT:

David A. Trissell, General Counsel, at 202–789–6820.

# SUPPLEMENTARY INFORMATION:

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## I. Introduction

II. Docketed Proceeding(s)

### I. Introduction

The Commission gives notice that the Postal Service filed request(s) for the Commission to consider matters related to negotiated service agreement(s). The request(s) may propose the addition or removal of a negotiated service agreement from the Market Dominant or the Competitive product list, or the modification of an existing product currently appearing on the Market Dominant or the Competitive product list.

Section II identifies the docket number(s) associated with each Postal Service request, the title of each Postal Service request, the request's acceptance date, and the authority cited by the Postal Service for each request. For each request, the Commission appoints an officer of the Commission to represent the interests of the general public in the proceeding, pursuant to 39 U.S.C. 505 (Public Representative). Section II also establishes comment deadline(s) pertaining to each request.

The public portions of the Postal Service's request(s) can be accessed via the Commission's website (*http:// www.prc.gov*). Non-public portions of the Postal Service's request(s), if any, can be accessed through compliance with the requirements of 39 CFR 3011.301.<sup>1</sup>

The Commission invites comments on whether the Postal Service's request(s) in the captioned docket(s) are consistent with the policies of title 39. For request(s) that the Postal Service states concern Market Dominant product(s), applicable statutory and regulatory requirements include 39 U.S.C. 3622, 39 U.S.C. 3642, 39 CFR part 3030, and 39 CFR part 3040, subpart B. For request(s) that the Postal Service states concern Competitive product(s), applicable statutory and regulatory requirements include 39 U.S.C. 3632, 39 U.S.C. 3633, 39 U.S.C. 3642, 39 CFR part 3035, and 39 CFR part 3040, subpart B. Comment deadline(s) for each request appear in section II.

## **II. Docketed Proceeding(s)**

1. Docket No(s).: MC2023–47 and CP2023–45; Filing Title: USPS Request to Add Priority Mail, First-Class Package Service & Parcel Select Contract 4 to Competitive Product List and Notice of Filing Materials Under Seal; Filing Acceptance Date: November 15, 2022; Filing Authority: 39 U.S.C. 3642, 39 CFR 3040.130 through 3040.135, and 39 CFR 3035.105; Public Representative: Christopher C. Mohr; Comments Due: November 23, 2022.

This Notice will be published in the **Federal Register**.

# Erica A. Barker,

Secretary.

[FR Doc. 2022–25335 Filed 11–21–22; 8:45 am] BILLING CODE 7710–FW–P

<sup>&</sup>lt;sup>1</sup> See Docket No. RM2018–3, Order Adopting Final Rules Relating to Non-Public Information, June 27, 2018, Attachment A at 19–22 (Order No. 4679).