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 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 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

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, ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.'' 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 9, 2022, 87 FR 27664.

1. *The title of the information collection:* ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery''.
2. *OMB approval number:* 3150–0217.
3. *Type of submission:* Extension.
4. *The form number, if applicable:* Not applicable.
5. *How often the collection is required or requested:* On occasion and annually.
6. *Who will be required or asked to respond:* Individuals and households; businesses and organizations; State, local, or Tribal governments.
7. *The estimated number of annual responses:* 4,200.
8. *The estimated number of annual respondents:* 4,200.
9. *The estimated number of hours needed annually to comply with the information collection requirement or request:* 1,087.5.
10. *Abstract:* The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner, for the purpose of improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management. Feedback collected under this generic clearance will provide useful information, but it will not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: The target population to which generalizations will be made, the sampling frame, the sample design (including stratification and clustering), the precision requirements or power calculations that justify the proposed sample size, the expected response rate, methods for assessing potential nonresponse bias, the protocols for data collection, and any testing procedures that were or will be undertaken prior to fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

Dated: December 15, 2022.

For the Nuclear Regulatory Commission.

David Cullison,

NRC Clearance Officer, Office of the Chief Information Officer.

[FR Doc. 2022–27593 Filed 12–19–22; 8:45 am]

BILLING CODE 7590–01–P

POSTAL REGULATORY COMMISSION

[Docket Nos. MC2023–90 and CP2023–91; MC2023–91 and CP2023–92; MC2023–92 and CP2023–93; MC2023–93 and CP2023–94]

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:* December 22, 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:

Table of Contents

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.[1]

The Commission invites comments on whether the Postal Service's request(s) in the captioned docket(s) are consistent

[1] *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).

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).:* MC2023–90 and CP2023–91; *Filing Title:* USPS Request to Add Priority Mail Contract 773 to Competitive Product List and Notice of Filing Materials Filed Under Seal; *Filing Acceptance Date:* December 14, 2022; *Filing Authority:* 39 U.S.C. 3642, 39 CFR 3040.130 through 3040.135, and 39 CFR 3035.105; *Public Representative:* Kenneth R. Moeller; *Comments Due:* December 22, 2022.

2. *Docket No(s).:* MC2023–91 and CP2023–92; *Filing Title:* USPS Request to Add Priority Mail Express, Priority Mail, First-Class Package Service & Parcel Select Contract 104 to Competitive Product List and Notice of Filing Materials Filed Under Seal; *Filing Acceptance Date:* December 14, 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:* December 22, 2022.

3. *Docket No(s).:* MC2023–92 and CP2023–93; *Filing Title:* USPS Request to Add Parcel Select Contract 55 to Competitive Product List and Notice of Filing Materials Filed Under Seal; *Filing Acceptance Date:* December 14, 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:* December 22, 2022.

4. *Docket No(s).:* MC2023–93 and CP2023–94; *Filing Title:* USPS Request to Add Parcel Select Contract 56 to Competitive Product List and Notice of Filing Materials Filed Under Seal; *Filing Acceptance Date:* December 14, 2022; *Filing Authority:* 39 U.S.C. 3642, 39 CFR 3040.130 through 3040.135, and 39 CFR 3035.105; *Public Representative:* Gregory S. Stanton; *Comments Due:* December 22, 2022.

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

Erica A. Barker,

Secretary.

[FR Doc. 2022–27619 Filed 12–19–22; 8:45 am]

BILLING CODE 7710–FW–P

OFFICE OF SCIENCE AND TECHNOLOGY POLICY

Request for Information; Identifying Ambiguities, Gaps, Inefficiencies, and Uncertainties in the Coordinated Framework for the Regulation of Biotechnology

AGENCY: Office of Science and Technology Policy (OSTP).

ACTION: Notice of request for information (RFI).

SUMMARY: The National Biotech and Biomanufacturing Initiative (NBBI) identified biotechnology regulation clarity and efficiency as a priority of the Administration. Thus, the White House Office of Science and Technology Policy (OSTP)—on behalf of the primary agencies that regulate the products of biotechnology, the U.S. Environmental Protection Agency (EPA), the Food and Drug Administration (FDA), and the U.S. Department of Agriculture (USDA)—requests relevant data and information, including case studies, that may assist in identifying any regulatory ambiguities, gaps, inefficiencies, or uncertainties in the Coordinated Framework for the Regulation of Biotechnology, particularly with regard to new and emerging biotechnology products. The information provided will inform regulatory agency efforts to improve the clarity and efficiency of the regulatory processes for biotechnology products.

DATES: Interested persons and organizations are invited to submit comments on or before 5 p.m. ET February 3, 2023.

ADDRESSES: USDA is managing this docket and is listed as the primary addressee below. All three agencies and OSTP will be considering all submitted comments as part of their efforts to identify regulatory ambiguities, gaps, or uncertainties in the Coordinated Framework.

You may submit information by any of the following methods (Due to time constraints, the eRulemaking Portal is strongly preferred):

- *Federal eRulemaking Portal:* Go to *http://www.regulations.gov.* Enter ''APHIS-2022-0076'' in the Search field. Select the Documents tab, then select the Comment button in the list of documents and follow the instructions to submit your comment.
- *Postal Mail:* Send your comment to the following address. Please include Docket No. APHIS-2022-0076 in the subject line.

Animal and Plant Health Inspection Service, US Department of Agriculture, 4700 River Road, Riverdale, MD 20737, Attn: Alan Pearson

- *Listening Sessions:* The regulatory agencies and OSTP will host a virtual public listening session on January 12, 2023. If you are interested in registering for the virtual listening session, go to *https://www.zoomgov.com/webinar/register/WN_IhbckX4VTiacK0AsyiikKQ.* If you are interested in additional listening sessions, please contact Dominique Carter at *biotech-regulation@ostp.eop.gov.* Summaries of the comments offered during the public listening session and any small listening sessions will be posted to the docket on *regulations.gov.*

Response to this request for information (RFI) is voluntary. Each individual or institution is requested to submit only one response. Responses should include the name of the person(s) or organization(s) filing the response. Please identify your answers by referring to a specific question number within the response.

Comments submitted in response to this notice are subject to the Freedom of Information Act (FOIA). Responses to this RFI may be posted without change online. No proprietary information, copyrighted information, or personally identifiable information should be submitted in response to this RFI.

This RFI is issued solely for information and planning purposes and does not constitute a solicitation. There will be no response to individual submissions. Please note that the United States Government will not pay for response preparation, or for the use of any information contained in a response. If submitting a response by mail, please allow sufficient time for mail processing and include the docket number and title.

FOR FURTHER INFORMATION CONTACT:

OSTP: Dominique Carter, *biotech-regulation@ostp.eop.gov,* tel: 202–456–4444.

EPA: Mike Mendelsohn, *Mendelsohn.Mike@epa.gov.*

FDA: Eric Flamm, *Eric.Flamm@fda.hhs.gov.*

USDA: Alan Pearson, *alan.pearson@usda.gov.*

SUPPLEMENTARY INFORMATION: