pursuant to the Authority's racetrack safety rules; or

(3) Prohibit a Racetrack from conducting any Covered Horserace.

8370. Final Civil Sanction

Any decision rendered by the Board of the Authority under Rule 8350, or the Authority under Rule 8360, shall constitute a final civil sanction subject to appeal and review in accordance with the provisions of 15 U.S.C. 3058.

8400. Investigatory Powers

(a) The Commission, the Authority, or their designees:

(1) Shall have free access to the books, records, offices, racetrack facilities, and other places of business of Covered Persons that are used in the care, treatment, training, and racing of Covered Horses, and to the books, records, offices, facilities, and other places of business of any person who owns a Covered Horse or performs services on a Covered Horse; and

(2) May seize any medication, drug, substance, paraphernalia, object, or device in violation or suspected violation of any provision of 15 U.S.C. 57A or the regulations of the Authority.

(b) A Covered Person shall:

(1) Cooperate with the Commission, the Authority or their designees during any investigation; and

(2) Respond truthfully to the best of the Covered Person's knowledge if questioned by the Commission, the Authority, or their designees about a racing matter.

(c) A Covered Person or any officer, employee or agent of a Covered Person shall not hinder a person who is conducting an investigation under or attempting to enforce or administer any provision of 15 U.S.C. 57A or the regulations of the Authority.

(d) The Commission or the Authority may issue subpoenas for the attendance of witnesses in proceedings within their jurisdiction and for the production of documents, records, papers, books, supplies, devices, equipment, and all other instrumentalities related to matters within the jurisdiction of the Commission or the Authority.

(e) Failure to comply with a subpoena or with the other provisions of this Rule may be penalized by the imposition of one or more penalties set forth in Rule 8200.

(f) The Commission or the Authority may administer oaths to witnesses and require witnesses to testify under oath in matters within the jurisdiction of the Commission or the Authority.

By direction of the Commission.

Joel Christie,

Acting Secretary.

[FR Doc. 2022–01663 Filed 1–25–22; 8:45 am]

BILLING CODE 6750–01–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–0246; Docket No. 2022–0001; Sequence No. 1]

Information Collection; General Services Administration Regulation; Packing List Clause

AGENCY: Office of Acquisition Policy, General Services Administration (GSA).

ACTION: Notice and request for comments.

SUMMARY: Under the provisions of the Paperwork Reduction Act of 1995, GSA invites the public to comment on a request to review and approve an extension of a previously approved information collection requirement regarding the packing list clause.

DATES: *Submit comments on or before:* March 28, 2022.

ADDRESSES: Submit comments identified by Information Collection 3090–0246 via *http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link ''Submit a Comment'' that corresponds with ''Information Collection 3090–0246, Packing List Clause''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 3090–0246, Packing List Clause'' on your attached document.

Instructions: Please submit comments only and cite Information Collection 3090–0246, Packing List Clause, in all correspondence related to this collection. Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two-to-three days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Mr. Clarence Harrison Jr, Procurement Analyst, at telephone 202–227–7051, or via email at *gsarpolicy@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

GSAR clause 552.211–77, Packing List, requires a contractor to include a packing list or other suitable document that verifies placement of an order and identifies the items shipped. In addition to information contractors would normally include on packing lists, the identification of cardholder name, telephone number and the term ''Credit Card'' is required.

B. Annual Reporting Burdens

Respondents: 14,923.
Responses per Respondent: 19.
Total Annual Responses: 283,233.
Hours per Response: .05.
Total Burden Hours: 14,161.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division, by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.*

Jeffrey A. Koses,

Senior Procurement Executive, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2022–01490 Filed 1–25–22; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10157 and CMS–R–262]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed

extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by February 25, 2022.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* The HIPAA Eligibility Transaction System (HETS); *Use:* CMS created the HETS application to provide Health Insurance Portability and Accountability Act of 1996 (HIPAA) compliant 270/271 health care eligibility inquiries (270) and responses (271) on a real-time basis. In creating the HETS application, federal law requires that CMS take precautions to minimize the security risk to federal information systems. Accordingly, CMS is requiring that trading partners who wish to connect to the HETS 270/271 application via the CMS Extranet and/or internet agree to specific trading partner terms as a condition of receiving access to Medicare eligibility information. Applicants will complete the entire Trading Partner Agreement form to indicate agreement with CMS trading partner terms and provide sufficient information to establish connectivity to the service and assure that those entities that access the Medicare eligibility information are aware of applicable provisions and penalties for the misuse of information.

CMS uses the Trading Partner Agreement Form to capture certain information whereby a person certifies that they are fully aware of any and all penalties related to the use of PHI and their access to this data from the HETS application. The information is an attestation by the authorized representative of an entity that wishes to access the Medicare eligibility information to conduct real-time eligibility transactions. The authorized representative is a person responsible for business decisions on behalf of the Organization who is submitting the access request. The data captured includes the authorized representative's name, title contact number and the name of the submitting entity. Other data captured is the submitter's National Provider Identifier (NPI), business name, billing address, physical address, and telephone number. *Form Number:* CMS–10157 (OMB control number: 0938–0960); *Frequency:* Annually; *Affected Public:* Private Sector, Businesses or other for-profits; *Number of Respondents:* 1,000; *Total Annual Responses:* 1,000; *Total Annual Hours:* 250. (For policy questions regarding this collection contact Rupinder Singh at 410–786–7484.)

2. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Contract Year 2023 Plan Benefit Package (PBP) Software and Formulary Submission; *Use:* Under the Medicare Modernization Act (MMA), Medicare Advantage (MA) and Prescription Drug Plan (PDP) organizations are required to submit plan benefit packages for all Medicare beneficiaries residing in their service area. The plan benefit package submission consists of the Plan Benefit Package (PBP) software, formulary file, and supporting documentation, as necessary. MA and PDP organizations use the PBP software to describe their organization's plan benefit packages, including information on premiums, cost sharing, authorization rules, and supplemental benefits. They also generate a formulary to describe their list of drugs, including information on prior authorization, step therapy, tiering, and quantity limits.

CMS requires that MA and PDP organizations submit a completed PBP and formulary as part of the annual bidding process. During this process, organizations prepare their proposed plan benefit packages for the upcoming contract year and submit them to CMS for review and approval. CMS uses this data to review and approve the benefit packages that the plans will offer to Medicare beneficiaries. This allows CMS to review the benefit packages in a consistent way across all submitted bids during with incredibly tight timeframes. This data is also used to populate data on Medicare Plan Finder, which allows beneficiaries to access and compare Medicare Advantage and Prescription Drug plans. *Form Number:* CMS–R–262 (OMB control number: 0938–0763); *Frequency:* Yearly; *Affected Public:* Private sector (Business or other for-profits and Not-for-profit institutions), State, Local, or Tribal Governments; *Number of Respondents:* 785; *Total Annual Responses:* 8,405; *Total Annual Hours:* 76,378. (For policy questions regarding this collection contact Kristy L Holtje at 410–786–2209.)

Dated: January 21, 2022.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2022–01495 Filed 1–25–22; 8:45 am]

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