(f) Notwithstanding any provision to the contrary in the rules of the Authority, the Authority may make public disclosure of any relevant information at any time, including prior to delivery of notice of a violation, if the Authority determines that such disclosure:

(1) Concerns a violation or circumstance that poses a serious and imminent risk of harm to Covered Persons, Covered Horses, or the public; or

(2) Is otherwise in the best interest of horseracing conducted at Covered Horseraces.

(g) The Authority shall publicly disclose the resolution of an alleged violation no later than 20 calendar days after the earlier of:

(1) The imposition of a final civil sanction;

(2) A resolution between the Authority and the Covered Person; or

(3) The dismissal of the allegation or a finding of no violation by the Authority.

(h) Public disclosure under paragraph (g)(1) & (2) shall include the following:

(1) The name of the Covered Person who committed the violation and any Covered Horse affected by the violation;

(2) The Rule violated;

(3) The sanction imposed;

(4) The order or other ruling issued in the matter; and

(5) The results of any appellate decisions concerning the violation.

(i) Public Disclosure shall not be required under this Rule if the Covered Person alleged to have committed a violation is a minor. Public disclosure concerning a case involving a minor shall be at the discretion of the Authority and in proportion to the facts and circumstances of the case.

(j) Publication shall be accomplished at a minimum by placing the required information on the Authority's website or publishing it through other means.

(k) Pursuant to 15 U.S.C. 3054, this Rule shall preempt any provision of State law or regulation, including those pertaining to data practice and privacy laws.

8400. Investigatory Powers

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

(1) Shall have free access to:

(i) With regard to Covered Persons, books, records, offices, racetrack facilities, and other places of business of Covered Persons that relate to the care, treatment, training, and racing of Covered Horses; and

(ii) With regard to any person who owns a Covered Horse or performs services on a Covered Horse, books, records, offices, facilities, and other places of business that relate to the care, treatment, training, and racing of Covered Horses.

(2) May seize any medication, drug, substance, or paraphernalia in violation or suspected violation of any provision of 15 U.S.C. Chapter 57A or the regulations of the Authority, and any object or device reasonably believed to have been used in furtherance of the violation or suspected violation.

(b) Upon final resolution of a violation, the Commission, the Authority, or their designees shall return seized property, including but not limited to phones, computers, and other repositories of electronic data, the possession of which is not specifically prohibited by the Act or the rules of the Authority.

(c) 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.

(d) 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. Chapter 57A or the regulations of the Authority.

(e) 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.

(f) 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.

(g) 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.

April J. Tabor,

Secretary.

[FR Doc. 2022–15972 Filed 7–25–22; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; ACF Performance Progress Report, ACF–OGM–SF–PPR–B (OMB #0970–0406)

AGENCY: Office of Grants Management, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Grants Management (OGM), in the Administration for Children and Families (ACF) is requesting a 3-year extension of the form ACF–OGM–SF–PPR–B (OMB #0970–406, expiration 11/30/2022). There are minor changes requested to the form.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

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. You can also obtain copies of the proposed collection of information by emailing *infocollection@acf.hhs.gov.* Identify all emailed requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: ACF's OGM is proposing the continued collection of program performance data for ACF's discretionary grantees using the existing ACF–OGM–SF–PPR–B (OMB #0970–0406, expiration 11/30/2022) form with minor changes to improve the function of the form. Revisions include collection of the Unique Entity Identifier instead of the Data Universal Numbering System, a rewording of the submission instructions to be more inclusive of all possible report submission methods utilized across ACF, and the addition of a program indicator to collect information on activities recipients conducted during the reporting period to address or advance equity. The form, developed by OGM, was created from the basic template of the OMB-approved reporting format of the Program

Performance Report. OGM uses this data to ensure grantees are proceeding in a satisfactory manner in meeting the approved goals and objectives of the project and if funding should be continued for another budget period.

OMB grants policy requires grantees to report on performance. Specific citations are contained in 45 CFR part 75 Uniform Administrative Requirements, Cost Principles, and Audit Requirements for HHS Awards.

Respondents: All ACF discretionary grantees. State governments, Native American Tribal governments, Native American Tribal Organizations, local governments, universities, and nonprofits with or without 501(c)(3) status with the IRS.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
ACF–OGM–SF–PPR–B	6,000	2	1	12,000

Estimated Total Annual Burden Hours: 12,000.

Authority: 45 CFR part 75.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–15940 Filed 7–25–22; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–0863]

Agency Information Collection Activities; Proposed Collection; Comment Request; Monthly Monitoring Study

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish a notice in the **Federal Register** concerning each proposed collection of information and to allow 60 days for public comment in response to the notice. This notice solicits comments on a proposed information collection entitled ''Monthly Monitoring Study.''

DATES: Either electronic or written comments on the collection of information must be submitted by September 26, 2022.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of September 26, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2022–N–0863 for the ''Monthly Monitoring Study'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*