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.

SUPPLEMENTARY INFORMATION:

Description: The COVID–19 pandemic has had profound effects on the U.S. economy, on the healthcare sector and on individuals and families across the country. The pandemic has also had broad implications for HPOG 2.0 programs—on how and how much healthcare training is delivered, on demand for healthcare workers, on interest in working in health care, and on the labor market more broadly. OPRE seeks to understand the particular experiences of those who apply for the HPOG Program during this period by surveying study participants enrolled after the onset of the pandemic. The COVID Cohort Survey would collect important information on participant experiences 15 months after randomization and would allow the impact study to compare impacts for pre-COVID participants with impacts for those enrolled after the onset the pandemic.

Respondents: HPOG impact study participants from the 27 non-tribal HPOG 2.0 grantees (treatment and control group members who enroll between May 2020 and September 2021).

Annual Burden Estimates: This request is specific to the COVID Cohort Survey. Currently approved materials and associated burden can be found at: *https://www.reginfo.gov/public/do/PRAICList?ref_nbr=201904-0970-006.*

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Instrument 12a: COVID–19 Cohort Survey	5,625	1	1	5,625	1,875

Estimated Total Annual Burden Hours: 1,875.

Authority: Section 2008 of the Social Security Act as enacted by Section 5507 of the Affordable Care Act and Section 413 of the Social Security Act, 42 U.S.C. 613.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–10732 Filed 5–20–21; 8:45 am]

BILLING CODE 4184–72–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Public Comment Request; State Performance Report [OMB #0985–New]

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living is announcing that the proposed collection of information listed above has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under section 506(c)(2)(A) of the Paperwork Reduction Act of 1995. This 30-Day notice collects comments on the information collection requirements related to the State Performance Report. This notice solicits comments on the new information collection requirements relating to the State Performance Report.

DATES: Comments on the collection of information must be submitted electronically by 11:59 p.m. (EST) or postmarked by June 21, 2021.

ADDRESSES: Submit written comments and recommendations for the proposed information collection within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find the information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. By mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

FOR FURTHER INFORMATION CONTACT: Susan Jenkins, Administration for Community Living, Washington, DC 20201, by email at *Susan.Jenkins@acl.hhs.gov* or by telephone at 202–795–7369.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, ACL has submitted the following proposed collection of information to OMB for review and clearance.

The Administration for Community Living (ACL) is requesting approval to collect data for the currently approved SPR under 0985–0008 expiring in FY 2022, which is the final reporting year for the currently approved OMB control number (0985–0008). In order to comply with requirements under the PRA it is necessary to place this ''new SPR'' IC under a new OMB control number while keeping the currently approved SPR under 0985–0008 active for remaining reporting in FY 2022.

The purpose of this data collection is to fulfill requirements of the Older Americans Act and the Government Performance and Results Modernization Act (GPRA Modernization Act) of 2010 and related program performance activities.

Section 202(a)(16) of the OAA requires the collection of statistical data regarding the programs and activities carried out with funds provided under the OAA and Section 207(a) directs the Assistant Secretary on Aging to prepare and submit a report to the President and Congress based on those data.

Section 202(f) directs the Assistant Secretary to develop a set of performance outcome measures for planning, managing, and evaluating activities performed and services provided under the OAA. Requirements pertaining to the measurement and evaluation of the impact of all programs authorized by the OAA described in section 206(a). The State Performance Report is one source of data used to develop and report performance outcome measures and measure program effectiveness in achieving the stated goals of the OAA.

The Administration on Aging (now within the Administration for

Community Living) first developed a State Program Report (SPR) in 1996 as part of its National Aging Program Information System (NAPIS). The SPR collects information about the national Aging Network, how State Agencies on Aging expend their OAA funds as well as funding from other sources for OAA authorized supportive services. The SPR also collects information on the demographic and functional status of the recipients, and is a key source for ACL performance measurement. This previously approved ''New SPR'' was a revision of the currently active version (effective 2019–2022) and approved in 2018, also assigned with the same OMB Control Number #0985–0001. This previously approved collection reduces the number of data elements reported by 70% compared to the 2019–2022 SPR.

ACL intends to seek a new OMB Control Number for the for the new SPR effective FY 2022–2025.

This request applies only to making an administrative change to the 2018 approved version of the State Performance Report for State Units on Aging (Older Americans Act Titles III and VII (Chapters 3 and 4) (''new SPR''). ACL intends to use this proposed data to collect information with the FY 2022 reporting year.

Comments in Response to the 60-Day Federal Register Notice: ACL published a 60-day notice in the **Federal Register** soliciting public comments on February 25, 2021, Volume 86, Number 36, pages 11541–11542. ACL received no comments.

Estimated Program Burden: ACL estimates an annual burden of 1,876 hours.

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Annual burden hours
SPR	56	1	33.5	1,876
Total	56	1	33.5	1,876

Dated: May 17, 2021.

Alison Barkoff,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2021–10708 Filed 5–20–21; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0441]

Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Cardiovascular and Renal Drugs Advisory Committee. The general function of the committee is to provide advice and recommendations to FDA on regulatory issues. The meeting will be open to the public. FDA is establishing a docket for public comment on this document.

DATES: The meeting will be held virtually on July 15, 2021, from 10 a.m. to 5 p.m. Eastern Time.

ADDRESSES: Please note that due to the impact of the COVID–19 pandemic, all meeting participants will be joining this advisory committee meeting via an online teleconferencing platform. Answers to commonly asked questions about FDA advisory committee meetings may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

FDA is establishing a docket for public comment on this meeting. The docket number is FDA–2021–N–0441. The docket will close on July 14, 2021. Submit either electronic or written comments on this public meeting by July 14, 2021. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before July 14, 2021. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of July 14, 2021. 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.

Comments received on or before June 30, 2021, will be provided to the committee. Comments received after that date will be taken into consideration by FDA. In the event that the meeting is cancelled, FDA will continue to evaluate any relevant applications or information, and consider any comments submitted to the docket, as appropriate.

You may submit comments as follows:

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–2021–N–0441 for ''Cardiovascular and Renal Drugs Advisory Committee; Notice of Meeting; Establishment of a