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. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration for Children and Families.

Mary Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2017–24901 Filed 11–16–17; 8:45 am]

BILLING CODE 4184–23–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity: Comment Request

Proposed Projects:

Community Services Block Grant (CSBG) State Plan Application for States

Community Services Block Grant (CSBG) Eligible Entity Master List

Community Services Block Grant (CSBG) ACSI Survey of Eligible Entities

Title: Community Services Block Grant (CSBG) State Plan Application

OMB Number: 0970–0382

Description: Section 676 of the Community Services Block Grant (CSBG) Act requires states, including the District of Columbia and the Commonwealth of Puerto Rico, and U.S. territories applying for CSBG funds to submit an application and plan (CSBG State Plan). The CSBG State Plan must meet statutory requirements prior to states and territories being funded with CSBG funds. Applicants have the option to submit a detailed plan annually or biannually. Entities that submit a biannual plan must provide an abbreviated plan the following year if substantial changes to the initial plan will occur.

This request is to revise the automated CSBG State Plan format for states and territories by revising questions for clarity and system compatibility. It is not anticipated that these revisions will cause any additional burden to states as they have been completing the automated plan for three years. It is anticipated that the burden will continue to diminish in subsequent years due to improved automation.

In addition to the CSBG State Plan, states will be requested to complete a CSBG Eligible Entity Master List in year one, and then make updates as necessary in subsequent years. As the states have the information about their eligible entities (or sub-grantees), the burden will be minimal to the states to complete this the first year.

Lastly, the request includes a survey for the CSBG eligible entities (or sub-grantees). The survey focuses on the customer service that the eligible entities receive from the CSBG states. The survey is optional, and this will be the third time that the eligible entities that chose to submit will complete it.

Respondents: State Governments, including the District of Columbia and the Commonwealth of Puerto Rico, and U.S. territories, and local level sub-grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
CSBG State Plan Application for States	56	1	31	1,736
CSBG State Plan Eligible Entity List	56	1	1	56
CSBG ACSI Survey of Eligible Entities	1,019	1	.15	152.85

Estimated Total Annual Burden Hours: 1,792 hours for states and territories; 152.85 for eligible entities.

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chap. 35), the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attn: ACF Reports Clearance Officer. Email address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–24905 Filed 11–16–17; 8:45 am]

BILLING CODE 4184–27–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–4179]

Cardiac Troponin Assays; Public Workshop; Request for Comments; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA or Agency) is extending the comment period provided in the notice entitled ''Cardiac Troponin Assays; Public Workshop; Request for Comments'' that appeared in the **Federal Register** on July 31, 2017. That notice announced the public workshop and requested comments by November 28, 2017; FDA is extending the public

workshop's comment period by 30 days to December 28, 2017, in response to requests for an extension to allow interested persons additional time to submit comments.

DATES: FDA is extending the comment period for the public workshop ''Cardiac Troponin Assays'' published on July 31, 2017 (82 FR 35532). Submit either electronic or written comments by December 28, 2017.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before December 28, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of December 28, 2017. 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–2017–N–4179 for ''Cardiac Troponin Assays; Public Workshop; Request for Comments.'' 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.

- 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.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Paula Caposino, Food and Drug Administration, Center for Devices and Radiological Health, 10903 New Hampshire Ave., Bldg. 66, Rm. 4644, Silver Spring, MD 20993, 301–796–6160, *Paula.Caposino@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 31, 2017 (82 FR 35532), FDA published a notice announcing the public workshop entitled ''Cardiac Troponin Assays; Public Workshop; Request for Comments'' with a 120-day comment period to request comments.

The Agency has received requests for a 30-day extension of the comment period for the public workshop. The request conveyed concern that the current 120-day comment period does not allow sufficient time to develop a meaningful or thoughtful response to the public workshop.

FDA has considered the request and is extending the comment period for the public workshop for 30 days, until December 28, 2017. The Agency believes that a 30-day extension allows adequate time for interested persons to submit comments.

Dated: November 13, 2017.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2017–24922 Filed 11–16–17; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–D–6154]

Evaluation of Devices Used With Regenerative Medicine Advanced Therapies; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft document entitled ''Evaluation of Devices Used with Regenerative Medicine Advanced Therapies; Draft Guidance for Industry.'' The draft guidance document, when finalized, will provide device manufacturers, applicants, and sponsors engaged in the development of regenerative medicine therapies, with our current thinking regarding evaluation of devices used in the recovery, isolation or delivery of regenerative advanced therapies, which FDA generally refers to as ''regenerative medicine advanced therapies'' or ''RMATs.'' Specifically, as required by the 21st Century Cures Act (Cures Act),