implement the protections of the Commission's final Rule.

In 2001, the Commission approved the Children's Advertising Review Unit (''CARU'') as a COPPA safe harbor program. In July 2009, Respondent joined CARU's COPPA safe harbor program. Thereafter, Respondent began disseminating statements regarding its participation in CARU's COPPA safe harbor program. Respondent remained a member of CARU's COPPA Safe Harbor Program until July 6, 2015, when CARU terminated Respondent's participation in the program. After CARU terminated Respondent from its safe harbor program, Respondent continued to make claims that it participated in the program.

The Commission's proposed one-count complaint alleges that Respondent violated Section 5(a) of the Federal Trade Commission Act. Specifically, the proposed complaint alleges that Respondent engaged in a deceptive act or practice by falsely representing that it was a current participant in the CARU COPPA safe harbor program when it was not.

Part I of the proposed order prohibits Respondent from making misrepresentations about its membership in any privacy or security program sponsored by the government or any other self-regulatory or standard-setting organization, including, but not limited to, the CARU COPPA safe harbor.

Parts II through V of the proposed order are reporting and compliance provisions. Part II requires acknowledgement of the order and dissemination of the order now and in the future to persons with responsibilities relating to the subject matter of the order. Part III ensures notification to the FTC of changes in corporate status and mandates that the company submit an initial compliance report to the FTC. Part IV requires the company to create certain documents relating to its compliance with the order for ten (10) years and to retain those documents for a five-year period. Part V mandates that the company make available to the FTC information or subsequent compliance reports, as requested.

Part VI is a provision ''sun-setting'' the order after twenty (20) years, with certain exceptions.

The purpose of this analysis is to aid public comment on the proposed order. It is not intended to constitute an official interpretation of the complaint or proposed order, or to modify in any way the proposed order's terms.

By direction of the Commission.

April J. Tabor,

Acting Secretary.

[FR Doc. 2020–11098 Filed 5–21–20; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Availability of Program Application Instructions for Medicare Improvements for Patients and Providers Act (MIPPA) Program Funds

AGENCY: Administration for Community Living, HHS.

ACTION: Notice

SUMMARY: The purpose of MIPPA funding is to enhance statewide and local coalition building focused on outreach, education, and one-to-one assistance activities to Medicare beneficiaries likely to be eligible for the Low Income Subsidy program (LIS) or the Medicare Savings Programs (MSP), above and beyond those regular activities planned in response to other funding. ACL will provide funding to State Health Insurance Assistance Programs (SHIP), Area Agencies on Aging (AAA), and Aging and Disability Resource Center (ADRC).

ACL seeks plans from applicants that will describe how the MIPPA funds will be used for outreach, education, and one-on-one application assistance over the next year. ACL requests that applicants submit a one (1) year state plan with specific project strategies to:

1. Enhance their one-on-one assistance, education, and outreach efforts to eligible Medicare beneficiaries regarding their preventive, wellness, and limited income benefits;

2. Describe how the SHIP, AAA, and ADRC efforts will be coordinated to provide outreach to beneficiaries with limited incomes statewide including rural areas and tribal entities;

3. Review and update previous MIPPA plans to reflect successes achieved to date and direct their efforts to enhance and expand their MIPPA outreach activities; and

4. Set performance goals, taking into account the MIPPA Performance Measures (PMs) implemented in Grant Year 2019 [Performance Measures include: Overall MIPPA Contacts; Overall Persons Reached through Outreach; MIPPA Target Populations (Under 65, Rural, Native American, English as a Secondary Language); and Contacts with Applications Submitted].

Additionally, programs should ensure MIPPA counselors familiarity with integrated care programs that support beneficiaries' independence at home and in the community, including Programs of All-Inclusive Care for the Elderly, Medicare Advantage Special Needs Plans and Supplemental benefits, and other integrated care programs for Medicare and Medicaid dual eligible beneficiaries. Applicant plans should go above and beyond those regular activities planned in response to other funding. State agencies may prepare either one statewide plan or separate plans for each eligible agency.

Announcement Type: Initial.

Funding Opportunity Number: CIP–MI–20–001.

Statutory Authority: The statutory authority for grants under this program announcement is contained in the 2006 Reauthorization of the Older Americans Act—Section 202 and the Medicare Improvements for Patients and Providers Act of 2008—Section 119, Public Law (Pub. L.) 110–275 as amended by the Patient Protection and Affordable Care Act of 2010 (Affordable Care Act), reauthorized by the American Taxpayer Relief Act of 2012 (ATRA), the Protecting Access to Medicare Act of 2014, and the Medicare Access and CHIP Reauthorization Act of 2015 (MACRA), the Bipartisan Budget Act of 2018, and the Coronavirus Aid, Relief, and Economic Security (CARES) Act of 2020.

Catalog of Federal Domestic Assistance (CFDA) Number: 93.071.

DATES: The deadline date for the submission of MIPPA Program State Plans is 11:59 p.m. EST July 20, 2020.

I. Award Information

1. Funding Instrument Type

These awards will be made in the form of grants to agencies for each MIPPA Priority Area:

Priority Area 1—Grants to state agencies (State Units on Aging or State Departments of Insurance) that administer the SHIP to provide enhanced outreach to eligible Medicare beneficiaries regarding their preventive, wellness, and limited income benefits; application assistance to individuals who may be eligible for LIS or MSPs; and outreach activities aimed at preventing disease and promoting wellness.

Priority Area 2—Grants to state agencies for AAA and Native American programs to provide enhanced outreach to eligible Medicare beneficiaries regarding their preventive, wellness, and limited income benefits; application assistance to individuals who may be eligible for LIS or MSPs; and outreach

activities aimed at preventing disease and promoting wellness.

Priority Area 3—Grants to agencies that are established ADRCs who have received an ADRC/No Wrong Door System (NWD) grant to provide outreach regarding Medicare Part D benefits related to LIS and MSPs, and conduct outreach activities aimed at preventing disease and promoting wellness.

2. Anticipated Total Priority Area Funding per Budget Period

ACL intends to make available, under this program announcement, grant awards for the three MIPPA priority areas. Funding will be distributed through a formula as identified in statute. The amounts allocated are based upon factors defined in statute and will be distributed to each priority area based on the formula. ACL will fund total project periods of up to one (1) year contingent upon availability of federal funds.

Priority Area 1—SHIP: $12.4 million in FY 2020 for state agencies that administer the SHIP Program.

Priority Area 2—AAA: $7.1 million in FY 2020 for State Units on Aging for Area Agencies on Aging and for Native American programs. Funding for Native American Programs ($285,000) is deducted from Priority 2 and is being allocated through a separate process.

Priority Area 3—ADRC: $4.7 million in FY 2020 for agencies that are established ADRCs who have received an ADRC/NWD grant.

II. Eligibility Criteria and Other Requirements

1. Eligible Applicants for MIPPA Priority Areas 1, 2 and 3

Priority Area 1: Only existing SHIP grant recipients are eligible to apply.

Priority Area 2: Only State Units on Aging are eligible to apply.

Priority Area 3: Only agencies that are established ADRCs who have received an Aging and Disability Resource Center (ADRC)/No Wrong Door System (NWD) grant.

Eligibility may change if future funding is available.

2. Cost Sharing or Matching is not Required

3. DUNS Number

All grant applicants must obtain and keep current a D–U–N–S number from Dun and Bradstreet. It is a nine-digit identification number, which provides unique identifiers of single business entities. The D–U–N–S number can be obtained from: *https://iupdate.dnb.com/iUpdate/viewiUpdateHome.htm.*

4. Intergovernmental Review

Executive Order 12372, Intergovernmental Review of Federal Programs, is not applicable to these grant applications.

III. Submission Information

1. Application Kits

Application Kits/Program Instructions are available at *www.grantsolutions.gov.* Instructions for completing the application kit will be available on the site.

2. Submission Dates and Times

To receive consideration, applications must be submitted by 11:59 p.m. Eastern Time on July 20, 2020, through *www.GrantSolutions.gov.*

VI. Agency Contacts

Direct inquiries regarding programmatic issues to:

Margaret Flowers, Phone: 202.795.7315, Email: *Margaret.Flowers@acl.hhs.gov.*

Dated: May 14, 2020.

Mary Lazare,

Principal Deputy Administrator.

[FR Doc. 2020–11050 Filed 5–21–20; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–1064]

Agency Information Collection Activities; Proposed Collection; Comment Request; State Petitions for Exemption From Preemption

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) 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 notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection provisions of our reporting requirements contained in existing FDA regulations governing state petitions for exemption from preemption.

DATES: Submit either electronic or written comments on the collection of information by July 21, 2020.

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 July 21, 2020. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of July 21, 2020. 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–1064 for ''Agency Information