

also be used to assign a performance score to each TIN/NPI in the virtual group. *Form Number:* CMS-10652 (OMB control number: 0938-NEW); *Frequency:* Annually; *Affected Public:* Private Sector: Business or other for-profits and Not-for-profit institutions and Individuals; *Number of Responses:* 16; *Total Annual Responses:* 16; *Total Annual Hours:* 160. (For policy questions regarding this collection contact Michelle Peterman at 410-786-2591.)

Dated: June 8, 2017.

William N. Parham, III,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Availability of Program Application Instructions for MIPPA Program Funds

Title: Medicare Improvements for Patients and Providers Act: State Plans for Medicare Savings Program, Low Income Subsidy & Prescription Drug Enrollment Outreach and Assistance.

Announcement Type: Initial.

Funding Opportunity Number: CIP-MI-17-001.

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 (PL) 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).

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

Dates: The deadline date for the submission of MIPPA Program State Plans is 11:59PM EST August 14, 2017.

I. Funding Opportunity Description

The purpose of MIPPA funding is to enhance state efforts to provide assistance to Medicare beneficiaries through statewide and local coalition building focused on intensified outreach activities to beneficiaries likely to be eligible for the Low Income Subsidy program (LIS) or the Medicare Savings

Program (MSP), and to assist those beneficiaries in applying for benefits. ACL will provide MIPPA program funding to State Health Insurance Assistance Programs (SHIPs), Area Agencies on Aging (AAAs), and Aging and Disability Resource Center programs (ADRCs) to inform Medicare beneficiaries about available Medicare program benefits. ACL seeks plans from states that will describe how the MIPPA program funds will be used for beneficiary outreach, education, and one-on-one application assistance over the next year.

ACL requests that states submit a one (1) year state plan with specific project strategies to expand, extend, or enhance their one-on-one assistance, education, and group outreach efforts to Medicare beneficiaries on Medicare and assistance programs for those with limited incomes. States should describe how the SHIP, AAA, and ADRC efforts will be coordinated to provide outreach to beneficiaries with limited incomes statewide. States that are eligible to apply are asked to review previous MIPPA plans and update these plans to reflect successes achieved to date and direct their efforts to enhance and expand their MIPPA outreach activities. State agencies may prepare either one statewide plan or separate plans for each eligible State agency.

II. Award Information

1. Funding Instrument Type

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

Priority Area 1—Grants to State Agencies (the State Unit on Aging or the State Department of Insurance) that administer the State Health Insurance Assistance Program (SHIP) to provide enhanced outreach to eligible Medicare beneficiaries regarding their benefits, enhanced outreach and application assistance to individuals who may be eligible for the Medicare Low Income Subsidy (LIS) or the Medicare Savings Program (MSP), and for the purposes of conducting outreach activities aimed at preventing disease and promoting wellness.

Priority Area 2—Grants to State Units on Aging for Area Agencies on Aging to provide enhanced outreach to eligible Medicare beneficiaries regarding their Medicare benefits, enhanced outreach and one-on-one application assistance to individuals who may be eligible for the LIS or the MSP, and for the purposes of conducting outreach activities aimed at preventing disease and promoting wellness.

Priority Area 3—Grants to State Units on Aging that administer the Aging and Disability Resource Centers to provide outreach to individuals regarding Medicare Part D benefits, benefits available under the LIS and MSP, and for the purposes of conducting 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: \$11.5 million in FY 2017 for state agencies that administer the SHIP Program.

Priority Area 2—AAA: \$7.9 million in FY 2017 for State Units on Aging for Area Agencies on Aging and for Native American programs. Funding for Native American Programs (\$270,000) is deducted from Priority 2 and is being allocated through a separate process.

Priority Area 3—ADRC: \$6 million in FY 2017 for State Agencies that received an ACL, Centers for Medicare and Medicaid Services (CMS), Veterans Health Administration (VHA) Aging and Disability Resource Center (ADRC)/No Wrong Door System (NWD) grant to support the development of their ADRC/NWD Systems.

III. Eligibility Criteria and Other Requirements

1. Eligible Applicants for MIPPA Priority Areas 1, 2 and 3: Awards made under this announcement, by statute, will be made only to agencies of State Governments.

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 State Agencies that received an ACL, CMS, VHA Aging and Disability Resource Center (ADRC)/No Wrong Door System (NWD) grant to support the development of their ADRC/NWD Systems are eligible for MIPPA funding in FY 2017.

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.

IV. 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 August 14, 2017, through www.GrantSolutions.gov.

V. Agency Contacts

Direct inquiries regarding programmatic issues to U.S. Department of Health and Human Services, Administration for Community Living, Office of Healthcare Information and Counseling, Washington, DC 20201, attention: Isaac C. Long or by calling 202-795-7315 or by email isaac.long@acl.hhs.gov.

Dated: June 9, 2017.

Daniel P. Berger,

Acting Administrator and Assistant Secretary for Aging.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0622]

Agency Information Collection Activities; Proposed Collection; Comment Request; Color Additive Certification Requests and Recordkeeping

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 FDA's regulations governing batch certification of color additives manufactured for use in foods, drugs, cosmetics, or medical devices in the United States.

DATES: Submit either electronic or written comments on the collection of information by August 14, 2017.

ADDRESSES: You may submit comments as follows. Late, untimely filed comments will not be considered. Electronic comments must be submitted on or before August 14, 2017. The <https://www.regulations.gov> electronic filing system will accept comments until midnight Eastern Time at the end of August 14, 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):** Division of

Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Division of Dockets Management, 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-2010-N-0622 for "Agency Information Collection Activities; Proposed Collection; Comment Request; Color Additive Certification Requests and Recordkeeping." 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 Division of Dockets Management 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 Division of Dockets Management. 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: <http://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