The public scoping meeting date is: Tuesday, September 12, 2017, from 6:30 p.m. to 8:30 p.m., Eastern Daylight Time (EDT).

ADDRESSES: CHI Center, 10501 New Hampshire Avenue, Silver Spring, Maryland 20903.

FOR FURTHER INFORMATION CONTACT: Paul Gyamfi, GSA, National Capital Region, Public Buildings Service, Office of Planning and Design Quality, at 202–440–3405. Please contact this number if special assistance is needed to attend and participate in the scoping meeting.

SUPPLEMENTARY INFORMATION: GSA intends to prepare an Environmental Impact Statement (EIS) to analyze the potential impacts resulting from the proposed Master Plan to support the FDA Headquarters consolidation at the Federal Research Center (FRC) at White Oak, located in Silver Spring, Maryland.

Background

In 1997, GSA completed an EIS that analyzed the impacts from the consolidation of 5,975 FDA employees at the FRC. In 2005, GSA completed a Supplemental Environmental Impact Statement (SEIS) that analyzed the impacts of increasing the number of employees from 5,947 to 7,720 and the impacts of adding a new eastern access entrance point into the FRC. In 2009, GSA completed its second SEIS that analyzed the impacts of increasing the number of employees (from 7,720 to 8,889) needed to conduct the complex and comprehensive reviews mandated by new legislation. To accommodate future growth and further consolidate FDA operations, GSA is preparing an EIS to assess the impacts of an employee population increase, of up to an approximately 18,000 employees, over a period of 15 years.

The purpose of the proposed action is to provide a Master Plan for the FDA Campus at FRC to accommodate the projected growth. The need for the proposed action is to continue to support the FDA Headquarters consolidation at FRC, and provide the necessary office and laboratory space, in order to conduct the complex and comprehensive reviews mandated by Congress.

Alternatives Under Consideration

GSA will analyze a range of alternatives (including the no action alternative) for the proposed Master Plan of the FDA Headquarters, to increase the campus population by up to an approximately 18,000 employees over 15 years. As part of the EIS, GSA will study the impacts of each alternative on the human environment.

Scoping Process

In accordance with NEPA, a scoping process will be conducted to aid in determining the alternatives to be considered and the scope of issues to be addressed, as well as for identifying the significant issues related to the proposed Master Plan. Scoping will be accomplished through a public scoping meeting, direct mail correspondence to potentially interested persons, agencies, and organizations, and meetings with agencies having an interest in the Master Plan. It is important that Federal, regional, State, and local agencies, and interested individuals take this opportunity to identify environmental concerns that should be addressed during the preparation of the Draft EIS.

Public Scoping Meeting

A public scoping meeting will be held on Tuesday, September 12, 2017, from 6:30 p.m. to 8:30 p.m., EDT. The meeting will be an informal open house along with a brief presentation, where visitors may come, receive information, and give comments. GSA is publishing notices in the Washington Post, Montgomery County Sentinel, and Prince George's Sentinel announcing the meeting.

Written Comments

Agencies and the public are encouraged to provide written comments on the scoping issues in addition to, or in lieu of, giving their comments at the public scoping meeting. Written comments regarding the environmental analysis for the proposed Master Plan must be postmarked between Monday, August 21, 2017, and Monday, September 25, 2017, and sent to the following address: General Services Administration, Public Buildings Service, Office of Planning and Design Quality, Attention: Paul Gyamfi, 301 7th Street SW., Room 4004, Washington, DC 20407. Email: *paul.gyamfi@gsa.gov* using the subject line: FDA White Oak Master Plan Comment.

Dated: August 4, 2017.

Mina Wright,

Director, Office of Planning and Design Quality, Public Buildings Service, National Capital Region, General Services Administration.

[FR Doc. 2017–16945 Filed 8–10–17; 8:45 am]

BILLING CODE 6820–Y1–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10454]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. 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 or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected; and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by September 11, 2017.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 *OR,* Email: *OIRA_submission@omb.eop.gov.*

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at Web site address at *https://www.cms.gov/Regulations-and-*

Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. No comments were received in response to the 60-day comment period. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Disclosure of State Rating Requirements; *Use:* The final rule ''Patient Protection and Affordable Care Act; Health Insurance Market Rules; Rate Review'' implements sections 2701, 2702, and 2703 of the Public Health Service Act (PHS Act), as added and amended by the Affordable Care Act, and sections 1302(e) and 1312(c) of the Affordable Care Act. The rule directs that states submit to CMS certain information about state rating and risk pooling requirements for their individual, small group, and large group markets, as applicable. Specifically, states will inform CMS of age rating ratios that are narrower than 3:1 for adults; tobacco use rating ratios that are narrower than 1.5:1; a state-established uniform age curve; geographic rating areas; whether premiums in the small and large group market are required to be based on average enrollee amounts (also known as composite premiums); and, in states that do not permit any rating variation based on age or tobacco use, uniform family tier structures and corresponding multipliers. In addition, states that elect to merge their individual and small group market risk pools into a combined pool will notify CMS of such election. This information will allow CMS to determine whether state-specific rules apply or Federal default rules apply. It will also support the accuracy of the Federal risk adjustment methodology. *Form Number:* CMS–10454 (OMB control number: 0938–1258); *Frequency:* On Occasion; *Affected Public:* State, Local, or Tribal Governments, Private Sector; *Number of Respondents:* 47; *Total Annual Responses:* 47; *Total Annual Hours:* 2,239. (For policy questions regarding this collection contact Russell Tipps at 301–492–4371.)

Dated: August 8, 2017.

William N. Parham, III,

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

[FR Doc. 2017–17020 Filed 8–10–17; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–7046–N]

Health Insurance Marketplace℠, Medicare, Medicaid, and Children's Health Insurance Programs; Meeting of the Advisory Panel on Outreach and Education (APOE), September 13, 2017

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Notice.

SUMMARY: This notice announces the next meeting of the Advisory Panel on Outreach and Education (APOE) (the Panel) in accordance with the Federal Advisory Committee Act. The Panel advises and makes recommendations to the Secretary of the U.S. Department of Health and Human Services (HHS) and the Administrator of the Centers for Medicare & Medicaid Services (CMS) on opportunities to enhance the effectiveness of consumer education strategies concerning the Health Insurance Marketplace℠, Medicare, Medicaid, and the Children's Health Insurance Program (CHIP). This meeting is open to the public.

DATES: *Meeting Date:* Wednesday, September 13, 2017, 8:30 a.m. to 4:00 p.m. eastern daylight time (e.d.t).

Deadline for Meeting Registration, Presentations, Special Accommodations and Comments: Wednesday, August 30, 2017, 5:00 p.m. (e.d.t.).

ADDRESSES: *Meeting Location:* U.S. Department of Health & Human Services, Hubert H. Humphrey Building, 200 Independence Avenue SW., Room 505A, Conference Room, Washington, DC 20201.

Presentations and Written Comments: Presentations and written comments should be submitted to: Thomas Dudley, Designated Federal Official (DFO), Office of Communications, Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Mailstop S1–05–06, Baltimore, MD 21244–1850 or via email at *Thomas.Dudley@cms.hhs.gov.*

Registration: The meeting is open to the public, but attendance is limited to the space available. Persons wishing to attend this meeting must register at the Web site *https://www.regonline.com/apoesept2017meeting* or by contacting the DFO as listed in the **FOR FURTHER INFORMATION CONTACT** section of this notice, by the date listed in the **DATES** section of this notice. Individuals requiring sign language interpretation or other special accommodations should contact the DFO at the address listed in the **ADDRESSES** section of this notice by the date listed in the **DATES** section of this notice.

FOR FURTHER INFORMATION CONTACT: Thomas Dudley, Designated Federal Official, Office of Communications, CMS, 7500 Security Boulevard, Mail Stop S1–05–06, Baltimore, MD 21244–1850, 410–786–1442, email *Thomas.Dudley@cms.hhs.gov.* Additional information about the APOE is available on the Internet at: *http://www.cms.gov/Regulations-and-Guidance/Guidance/FACA/APOE.html.* Press inquiries are handled through the CMS Press Office at (202) 690–6145.

SUPPLEMENTARY INFORMATION:

I. Background

The Advisory Panel for Outreach and Education (APOE) (the Panel) is governed by the provisions of Federal Advisory Committee Act (FACA) (Pub. L. 92–463), as amended (5 U.S.C. App. 2), which sets forth standards for the formation and use of federal advisory committees. The Panel is authorized by section 1114(f) of the Social Security Act (42 U.S.C. 1314(f)) and section 222 of the Public Health Service Act (42 U.S.C. 217a).

The Secretary of the U.S. Department of Health and Human Services (HHS) (the Secretary) signed the charter establishing the Citizen's Advisory Panel on Medicare Education [1] (the

[1] We note that the Citizen's Advisory Panel on Medicare Education is also referred to as the Advisory Panel on Medicare Education (65 FR

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