conducts independent reviews of the cost of travel and the operation of privately owned airplanes and motorcycles on an annual basis to determine their corresponding mileage reimbursement rates. These reviews evaluate various factors, such as the cost of fuel, depreciation of the original vehicle cost, maintenance and insurance, state and Federal taxes, and consumer price index data. FTR Bulletin 20–03 establishes and announces the new CY 2020 POV mileage reimbursement rates for official temporary duty and relocation travel. This notice is the only notification to agencies of revisions to the POV mileage rates for official travel, and relocation, other than the changes posted on GSA's website at *https://gsa.gov/mileage*.

Jessica Salmoiraghi,

Associate Administrator, Office of Government-wide Policy.

[FR Doc. 2020–00390 Filed 1–13–20; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—Funding Opportunity Announcement (FOA), PAR 16–098, Cooperative Research Agreements to the World Trade Center Health Program (U01).

Dates: March 10, 2020; and March 11, 2020.

Times: Day One: 8:00 a.m.–5:00 p.m., EDT; and Day Two: 8:00 a.m.–12:00 p.m., EDT.

Place: Courtyard Marriott Decatur Downtown/Emory, 130 Clairmont Avenue, Decatur, Georgia 30030, Telephone: (404) 371–0204.

Agenda: To review and evaluate grant applications.

FOR FURTHER INFORMATION CONTACT: Nina Turner, Ph.D., Scientific Review Officer, CDC/NIOSH, 1095 Willowdale Road, Mailstop G905, Morgantown, West Virginia 26505, Telephone: (304) 285–5975.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2020–00372 Filed 1–13–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–588, CMS–855B and CMS–R–262]

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 February 13, 2020.

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' website address at website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

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

2. 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. 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:* Extension of a currently

approved collection; *Title of Information Collection:* Electronic Funds Transfer Authorization Agreement; *Use:* Section 1815(a) of the Social Security Act provides the authority for the Secretary of Health and Human Services to pay providers/suppliers of Medicare services at such time or times as the Secretary determines appropriate (but no less frequently than monthly). Under Medicare, CMS, acting for the Secretary, contracts with Fiscal Intermediaries and Carriers to pay claims submitted by providers/suppliers who furnish services to Medicare beneficiaries. Under CMS' payment policy, Medicare providers/suppliers have the option of receiving payments electronically. Form number CMS–588 authorizes the use of electronic fund transfers (EFTs). *Form Number:* CMS–588 (OMB control number: 0938–0626); *Frequency:* On occasion; *Affected Public:* Business or other for-profit and Not-for-profit institutions; *Number of Respondents:* 100,000; *Total Annual Responses:* 100,000; *Total Annual Hours:* 100,000. (For questions regarding this collection contact Kim McPhillips at 410–786–5374.)

2. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* Medicare Enrollment Application for Clinics/Group Practices and Other Suppliers Revision; *Use:* The primary function of the CMS–855B Medicare enrollment application for suppliers, also known as Health Diagnosing and Treating Practitioners, is to gather information from the supplier that tells CMS who the supplier is, whether the supplier meets certain qualifications to be a Medicare health care provider or supplier, where the supplier practices or renders services, and other information necessary to establish correct claims payments.

The CMS–855B form includes an attachment for Opioid Treatment Programs (OTPs). This attachment is only used to capture the OTP personnel and consists of limited data fields (name, Social Security Number, National Provider Identifier, and license number) in response to the ''SUPPORT for Patients and Communities Act'' that was signed into law on October 24, 2018. This legislation was designed to alleviate the nationwide opioid crisis by: (1) Reducing the abuse and supply of opioids; (2) helping individuals recover from opioid addiction and supporting the families of these persons; and (3) establishing innovative and long-term solutions to the crisis. Section 2005 of the SUPPORT Act establishes a new Medicare Part B benefit for opioid use disorder (OUD) treatment services furnished by opioid treatment programs (OTPs) beginning on or after January 1, 2020. *Form Number:* CMS–855B (OMB control number: 0938–New); *Frequency:* Annually; *Affected Public:* Individuals and households; *Number of Respondents:* 327,696; *Total Annual Responses:* 327,696; *Total Annual Hours:* 522,041. (For questions regarding this collection contact Kim McPhillips at 410–786–5374.)

3. *Type of Information Collection Request:* Revision with change of a currently approved collection; *Title of Information Collection:* Contract Year 2021 Plan Benefit Package (PBP) Software and Formulary Submission; *Use:* Under the Medicare Modernization Act (MMA), Medicare Advantage (MA) and Prescription Drug Plan (PDP) organizations are required to submit plan benefit packages for all Medicare beneficiaries residing in their service area. The plan benefit package submission consists of the Plan Benefit Package (PBP) software, formulary file, and supporting documentation, as necessary. MA and PDP organizations use the PBP software to describe their organization's plan benefit packages, including information on premiums, cost sharing, authorization rules, and supplemental benefits. They also generate a formulary to describe their list of drugs, including information on prior authorization, step therapy, tiering, and quantity limits.

CMS requires that MA and PDP organizations submit a completed PBP and formulary as part of the annual bidding process. During this process, organizations prepare their proposed plan benefit packages for the upcoming contract year and submit them to CMS for review and approval. CMS uses this data to review and approve the benefit packages that the plans will offer to Medicare beneficiaries. This allows CMS to review the benefit packages in a consistent way across all submitted bids during with incredibly tight timeframes. This data is also used to populate data on Medicare Plan Finder, which allows beneficiaries to access and compare Medicare Advantage and Prescription Drug plans. *Form Number:* CMS–R–262 (OMB control number: 0938–0763); *Frequency:* Yearly; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 672; *Total Annual Responses:* 7,264; *Total Annual Hours:* 67,368. (For policy questions regarding this collection contact Kristy L. Holtje at 410–786–2209.)

Dated: January 9, 2020.

William N. Parham, III,

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

[FR Doc. 2020–00426 Filed 1–13–20; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–P–0015A]

Agency Information Collection Activities: Proposed Collection; 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 (the 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 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates 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 must be received by March 16, 2020.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following