Summary of Proposed Order With Respondent

The Proposed Order contains injunctive relief designed to prevent Respondents from engaging in the same or similar acts or practices in the future.

Part I prohibits Respondents from misrepresenting the extent to which (1) it collects, maintains, uses, discloses, deletes any covered information, and (2) the location data that Respondents collect, use, maintain, or disclose is deidentified.

Part II prohibits Respondents from selling, licensing, transferring, sharing, disclosing, or using sensitive location data in any products or services. Sensitive locations are defined as those locations in the United States associated with (1) medical facilities (*e.g.,* family planning centers, general medical and surgical hospitals, offices of physicians, offices of mental health physicians and practitioners, residential mental health and substance abuse facilities, outpatient mental health and substance abuse centers, outpatient care centers, psychiatric and substance abuse hospitals, and specialty hospitals); (2) religious organizations; (3) correctional facilities; (4) labor union offices; (5) locations of entities held out to the public as predominantly providing education or childcare services to minors; (6) associations held out to the public as predominantly providing services based on racial or ethnic origin; or (7) locations held out to the public as providing temporary shelter or social services to homeless people, survivors of domestic violence, refugees, or immigrants.

Part III requires that Respondents implement and maintain a sensitive location data Program to develop a comprehensive list of sensitive locations and to prevent the use, sale, license, transfer, or disclosure of sensitive location data.

Part IV requires that Respondents establish and implement policies, procedures, and technical measures designed to prevent recipients of Respondents' location data from associating consumers with locations providing services to LGBTQ+ individuals, locations of public gatherings of individuals during social demonstrations, marches, or protests, or using location data to determine the identity or location of an individual's home.

Part V requires Respondents to notify the Commission any time it determines that a third party shared Respondents' Location Data, in violation of a contractual requirement between Respondents and the third party.

Part VI requires that Respondents must not collect, use, maintain, and disclose location data (1) when consumers have opted-out, or otherwise declined targeted advertising, (2) without a record documenting the consumer's consent obtained prior to the collection of location data, and (3) in connection with Respondents' apps unless consumers receive a clear and conspicuous quarterly reminder about location data being collected.

Part VII requires that Respondents implement a supplier assessment program designed to ensure that consumers have provided consent for the collection and use of location data obtained by Respondents. Under this program, Respondents must conduct initial assessments of all their data suppliers within 30 days of entering into a data sharing agreement, or within 30 days of the initial date of data collection. The program also requires that Respondents confirm that consumers provide consent and create and maintain records of suppliers' assessment responses. Finally, Respondents must cease from using, selling, or disclosing location data for which consumers have not provided consent.

Part VIII requires that Respondents provide a clear and conspicuous means for consumers to request the identity of any entity, business, or individual to whom their location data has been sold, transferred, licensed, or otherwise disclosed or a method to delete the consumers' location data from the databases of Respondents' customers. Respondents must also provide written confirmation to consumers that the deletion requests have been sent to Respondents' customers.

Part IX requires that Respondents provide a simple, easily-located means for consumers to withdraw any consent provided and Part X requires that Respondents cease collecting location data within 15 days after Respondents receive notice that the consumer withdraws their consent.

Part XI also requires that Respondents provide a simple, easily-located means for consumers to request that Respondents delete location data that Respondents previously collected and to delete the location data within 30 days of receipt of such request unless a shorter period for deletion is required by law.

Part XII requires that Respondents (1) document and adhere to a retention schedule for the covered information it collects from consumers, including the purposes for which it collects such information, the specific business needs, and an established timeframe for its deletion, and (2) prior to collecting or using new type of information related to consumers that was not previously collected, and is not described in its retention schedule, Respondents must update its retention schedule.

Part XIII requires that Respondents delete or destroy all historic location data and all data products. Respondents have the option to retain historic location data if it has obtained affirmative express consent or it ensures that the historic location data is deidentified or rendered non-sensitive. Respondents must inform all customers that received location data from Respondents within 3 years prior to the issuance date of this Order, of the Commission's position that such data should be deleted, deidentified, or rendered non-sensitive.

Part XIV requires Respondents to establish and implement, and thereafter maintain, a comprehensive privacy program that protects the privacy of consumers' personal information.

Parts XV–XVIII are reporting and compliance provisions, which include recordkeeping requirements and provisions requiring Respondent to provide information or documents necessary for the Commission to monitor compliance.

Part XIX states that the Proposed Order will remain in effect for 20 years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the Proposed Order, and it is not intended to constitute an official interpretation of the complaint or Proposed Order, or to modify the Proposed Order's terms in any way.

By direction of the Commission.

April J. Tabor,

Secretary.

[FR Doc. 2024–00928 Filed 1–17–24; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

Pursuant to 5 U.S.C. 1009(d), 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, Office of Strategic Business Initiatives, 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)—RFA–IP–24–045, Network of Community Cohorts for Monitoring Changes in Respiratory Virus Epidemiology (Pandemic Preparedness Cohorts).

Dates: April 25–26, 2024.

Times: 10 a.m.–5 p.m., EDT.

Place: Videoconference.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Gregory Anderson, MS, M.P.H, Scientific Review Officer, National Center for HIV, Viral Hepatitis, STD, and TB Prevention, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H24–6, Atlanta, Georgia 30329–4027. Telephone: (404) 718–8833; Email: *GAnderson@cdc.gov.*

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

[FR Doc. 2024–00826 Filed 1–17–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10346]

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

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 20, 2024.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

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 without change of a currently approved collection; *Title of Information Collection:* Quality Bonus Payment Appeals; *Use:* Section 1853(o) of the Act requires CMS to make QBPs to MA organizations that achieve performance rating scores of at least 4 stars under a five-star rating system. While CMS has applied a Star Rating system to MA organizations for a number of years, prior to the QBP program these Star Ratings were used only to provide additional information for beneficiaries to consider in making their Part C and D plan elections. Beginning in 2012, the Star Ratings CMS assigns for purposes of QBPs directly affected the monthly payment amount MA organizations receive from CMS under their contracts. Additionally, section 1854(b)(1)(C)(v) of the Act, as added by the Affordable Care Act, also requires CMS to change the share of savings that MA organizations must provide to enrollees as the beneficiary rebate specified at § 422.266(a) based on the level of a sponsor's Star Rating for quality performance.

The information collected on the Request for Reconsideration form from MA organizations is considered by the reconsideration official and potentially the hearing officer to review CMS's determination of the organization's eligibility for a QBP. The form asks MA organizations to select the Star Ratings measure(s) they believe was miscalculated or used incorrect data and describe what they believe is the issue. Under § 422.260(c)(3)(ii) these are the only bases for appeals. In conducting the reconsideration, the reconsideration official will review the QBP determination, the evidence and findings upon which it was based, and any other written evidence submitted by the organization with their Request for Reconsideration or by CMS before the reconsideration determination is made. *Form Number:* CMS–10346 (OMB Control Number 0938–1129; *Frequency:* Yearly; *Affected Public:* Private Sector; *Number of Respondents:* 20; *Total Annual Responses:* 20; *Total Annual Hours:* 160. (For policy questions regarding this collection contact, Sarah Gaillot at 410–786–4637.)