and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. *See* FTC Rule 4.9(c). Your comment will be kept confidential only if the General Counsel grants your request in accordance with the law and the public interest. Once your comment has been posted on the public FTC website—as legally required by FTC Rule 4.9(b)—we cannot redact or remove your comment from the FTC website, unless you submit a confidentiality request that meets the requirements for such treatment under FTC Rule 4.9(c), and the General Counsel grants that request.

Visit the FTC website at *http://www.ftc.gov* to read this Notice and the news release describing it. The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding, as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before January 17, 2020. For information on the Commission's privacy policy, including routine uses permitted by the Privacy Act, see *https://www.ftc.gov/site-information/privacy-policy*.

Analysis of Proposed Consent Order To Aid Public Comment

The Federal Trade Commission (''Commission'') has accepted, subject to final approval, an agreement containing a consent order from Incentive Services, Inc. (''Incentive Services'' or ''Respondent'').

The proposed consent order (''proposed order'') has been placed on the public record for 30 days for receipt of comments by interested persons. Comments received during this period will become part of the public record. After 30 days, the Commission will again review the agreement and the comments received, and will decide whether it should withdraw from the agreement and take appropriate action or make final the agreement's proposed order.

This matter concerns alleged false or misleading representations that Incentive Services made concerning its participation in the Privacy Shield framework agreed upon by the U.S. and the European Union (''EU''). The Privacy Shield framework allows for the lawful transfer of personal data from the EU to participating companies in the U.S. The framework consists of a set of principles and related requirements that have been deemed by the European Commission as providing ''adequate'' privacy protection. The principles include notice; choice; accountability for onward transfer; security; data integrity and purpose limitation; access; and recourse, enforcement, and liability. The related requirements include, for example, securing an independent recourse mechanism to handle any disputes about how the company handles information about EU citizens.

To participate in the framework, a company must comply with the Privacy Shield principles and self-certify that compliance to the U.S. Department of Commerce (''Commerce''). Commerce reviews companies' self-certification applications and maintains a public website, *https://www.privacyshield.gov/list,* where it posts the names of companies who have completed the requirements for certification. Companies are required to recertify every year in order to continue benefitting from Privacy Shield.

Incentive Services is a company that works with organizations to improve performance of individual employees through service award programs, performance incentives, and loyalty programs. According to the Commission's complaint, Incentive Services published on its website, *https://www.incentiveservices.com/,* a privacy policy containing statements related to its participation in Privacy Shield. However, it only initiated an application to Commerce for Privacy Shield certification, and did not complete the steps necessary to participate in the framework.

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 certified participant in the EU–U.S. and the Swiss-U.S. Privacy Shield frameworks.

Part I of the proposed order prohibits the company 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 EU–U.S. Privacy Shield framework, the Swiss-U.S. Privacy Shield framework, and the APEC Cross-Border Privacy Rules.

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 20 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 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. 2019–27237 Filed 12–17–19; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meetings

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Notice of five AHRQ subcommittee meetings.

SUMMARY: The subcommittees listed below are part of AHRQ's Health Services Research Initial Review Group Committee. Grant applications are to be reviewed and discussed at these meetings. Each subcommittee meeting will commence in open session before closing to the public for the duration of the meeting.

DATES: See below for dates of meetings:

1. *Healthcare Effectiveness and Outcomes Research (HEOR)*
   Date: February 12–13, 2020 (Open from 8:00 a.m. to 8:30 a.m. on February 12 and closed for remainder of the meeting)
2. *Healthcare Safety and Quality Improvement Research (HSQR)*
   Date: February 20–21, 2020 (Open from 7:30 a.m. to 8:00 a.m. on February 20 and closed for remainder of the meeting)
3. *Health System and Value Research (HSVR)*
   Date: February 27–28, 2020 (Open from 8:00 a.m. to 8:30 a.m. on February 27 and closed for remainder of the meeting)
4. *Health Care Research and Training (HCRT)*

Date: February 27–28, 2020 (Open from 8:00 a.m. to 8:30 a.m. on February 27 and closed for remainder of the meeting)

5. *Healthcare Information Technology Research (HITR)*

Date: February 27–28, 2020 (Open from 8:00 a.m. to 8:30 a.m. on February 27 and closed for remainder of the meeting)

ADDRESSES: (Below specifics hotel where each meeting will be held:)

Bethesda North Marriott Hotel & Conference Center, 5701 Marinelli Road, Bethesda, Maryland 20852, (HEOR, HITR, HCRT, HSVR).

Hilton Washington DC/Rockville Hotel & Executive Meeting Center, 1750 Rockville Pike, Rockville, MD 20852, (HSQR).

FOR FURTHER INFORMATION CONTACT: (To obtain a roster of members, agenda or minutes of the non-confidential portions of the meetings.)

Jenny Griffith, Acting Committee Management Officer, Office of Extramural Research Education and Priority Populations, Agency for Healthcare Research and Quality (AHRQ), 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 427–1557.

SUPPLEMENTARY INFORMATION: In accordance with section 10 (a)(2) of the Federal Advisory Committee Act (5 U.S.C. App. 2), AHRQ announces meetings of the above-listed scientific peer review groups, which are subcommittees of AHRQ's Health Services Research Initial Review Group Committees. Each subcommittee meeting will commence in open session before closing to the public for the duration of the meeting. The subcommittee meetings will be closed to the public in accordance with the provisions set forth in 5 U.S.C. App. 2 section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). 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.

Agenda items for these meetings are subject to change as priorities dictate.

Virginia L. Mackay-Smith,

Associate Director.

[FR Doc. 2019–27263 Filed 12–17–19; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10108, CMS–10243, CMS–10383, CMS–10609, CMS–R–131 and CMS–10662]

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 January 17, 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:* Medicaid Managed Care Regulations; *Use:* The requirements contained in this information collection request implement regulations that allow states greater flexibility to implement mandatory managed care programs, implement new beneficiary protections, and eliminate certain requirements viewed by state agencies as impediments to the growth of managed care programs. Information collected includes information about managed care programs, grievances and appeals, enrollment broker contracts, and managed care organizational capacity to provide health care services. Medicaid enrollees use the information collected and reported to make informed choices regarding health care, including how to access health care services and the grievance and appeal system. States use the information collected and reported as part of its contracting process with managed care entities, as well as its compliance oversight role. We use the information collected and reported in an oversight role of state Medicaid managed care programs. *Form Number:*