identifiable health information. In addition, your comment should not include any ''trade secret or any commercial or financial information which . . . is privileged or confidential''—as provided by Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2)—including in particular competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled ''Confidential,'' and must comply with FTC Rule 4.9(c). In particular, the written request for confidential treatment that accompanies the comment must include the factual 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 Web site—as legally required by FTC Rule 4.9(b)—we cannot redact or remove your comment from the FTC Web site, 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 Web site 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 October 10, 2017. 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 (''FTC'' or ''Commission'') has accepted, subject to final approval, a consent agreement applicable to Tru Communication, Inc. dba TCPrinting.net (''TCP'').

The proposed consent order has been placed on the public record for thirty (30) days for receipt of comments by interested persons. Comments received during this period will become part of the public record. After thirty 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 TCP made to consumers concerning its participation in the EU-U.S. Privacy Shield framework agreed upon by the U.S. and the European Union (''EU''). The EU-U.S. Privacy Shield framework allows U.S. companies to transfer data outside the EU consistent with EU law. To join the EU-U.S. Privacy Shield framework, a company must self-certify to the U.S. Department of Commerce (''Commerce'') that it complies with a set of principles and related requirements that have been deemed by the European Commission as providing ''adequate'' privacy protection. These principles include notice; choice; accountability for onward transfer; security; data integrity and purpose limitation; access; and recourse, enforcement, and liability. Commerce maintains a public Web site, *https://www.privacyshield.gov/list,* where it posts the names of companies that have self-certified to the EU-U.S. Privacy Shield framework. The listing of companies indicates whether their self-certification is current. Companies are required to re-certify every year in order to retain their status as current members of the EU-U.S. Privacy Shield framework.

TCP provides printing services such as copying, binding and scanning of documents. According to the Commission's complaint, TCP has set forth on its Web site, *www.tcprinting.net/info/lpi-privacy-policy.php,* privacy policies and statements about its practices, including statements related to its participation in the EU-U.S. Privacy Shield framework.

The Commission's complaint alleges that TCP falsely represented that it was certified to participate in the EU-U.S. Privacy Shield framework when, in fact, TCP never completed the necessary steps to finalize its application and thus, was not certified to participate in the EU-U.S. Privacy Shield framework.

Part I of the proposed order prohibits TCP 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.

Parts II through VI 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 TCP submit an initial compliance report to the FTC. Part IV requires TCP to retain documents relating to its compliance with the order for a five-year period.

Part V mandates that TCP make available to the FTC information or subsequent compliance reports, as requested. Part VI is a provision ''sunsetting'' the order after twenty (20) years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the proposed order. It is not intended to constitute an official interpretation of the proposed complaint or order or to modify the order's terms in any way.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2017–19619 Filed 9–14–17; 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. *Health System and Value Research (HSVR)*
   Date: October 18, 2017 (Open from 8:00 a.m. to 9:00 a.m. on October 18th and closed for remainder of the meeting)
2. *Healthcare Safety and Quality Improvement Research (HSQR)*
   Date: October 18–19, 2017 (Open from 7:30 a.m. to 8:30 a.m. on October 18th and closed for remainder of the meeting)
3. *Healthcare Effectiveness and Outcomes Research (HEOR)*
   Date: October 18–19, 2017 (Open from 8:30 a.m. to 9:30 a.m. on October 18th and closed for remainder of

the meeting)

4. *Health Care Research and Training (HCRT)*

Date: October 19–20, 2017 (Open from 8:00 a.m. to 9:00 a.m. on October 19th and closed for remainder of the meeting)

5. *Healthcare Information Technology Research (HITR)*

Date: October 26–27, 2017 (Open from 8:00 a.m. to 9:00 a.m. on October 26th and closed for remainder of the meeting)

ADDRESSES: (Below specifics where each hotel will be held) Hilton Rockville, 1750 Rockville Pike, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: (To obtain a roster of members, agenda or minutes of the non-confidential portions of the meetings.) Mrs. Bonnie Campbell, 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) 427–1554.

SUPPLEMENTARY INFORMATION: These meetings will be closed to the public in accordance with 5 U.S.C. App. 2 section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). 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.

Sharon B. Arnold,

Deputy Director.

[FR Doc. 2017–19643 Filed 9–14–17; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection: Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

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

ACTION: Notice.

SUMMARY: As part of a Federal Government-wide effort to streamline the process to seek feedback from the public on service delivery, AHRQ has submitted a Generic Information Collection Request (Generic ICR): ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.''

This proposed information collection was previously published in the **Federal Register** on May 30, 2017, and allowed 60 days for public comment. No substantive comments were received. The purpose of this notice is to allow an additional 30 days for public comment.

DATES: Comments on this notice must be received by October 16, 2017.

ADDRESSES: Written comments should be submitted to: AHRQ's OMB Desk Officer by fax at (202) 395–6974 (attention: AHRQ's desk officer) or by email at *OIRA_submission@omb.eop.gov* (attention: AHRQ's desk officer).

SUPPLEMENTARY INFORMATION:

Proposed Project

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

AHRQ has submitted a Generic Information Collection Request (Generic ICR): ''Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery'' to OMB for approval under the Paperwork Reduction Act (PRA) (44 U.S.C. 3501 *et. seq.*). The information collection activity will gather qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery.

Qualitative feedback is information that provides useful insights on perceptions and opinions, but is not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. The feedback will contribute directly to the improvement of program management. The current clearance was approved on November 11, 2014 (OMB Control Number 0935–0179) and will expire on November 30, 2017.

Below we provide AHRQ's projected average annual estimates for the next three years:

Current Actions: New collection of information.

Type of Review: New Collection.

Affected Public: Individuals and Households, Businesses and Organizations, State, Local or Tribal Government.

Average Expected Annual Number of activities: 10.

Respondents: 10,900.

Annual responses: 10,900.

Frequency of Response: Once per request.

The total number of respondents across all 10 activities in a given year is 10,900.

Average minutes per response: 19.

Burden hours: 3,452.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid Office of Management and Budget control number.

Request for Comments

In accordance with the Paperwork Reduction Act, comments on AHRQ's information collection are requested with regard to any of the following: (a) Whether the proposed collection of information is necessary for the proper performance of AHRQ healthcare research and healthcare information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent