

ESTIMATED ANNUALIZED BURDEN HOURS

| Respondents | Form name | Number of respondents | Number of responses per respondent | Average burden per response (in hours) |
|----------------------|---|-----------------------|------------------------------------|--|
| Private Sector | Health Center Organizational Assessment | 21 | 1 | 2 |
| | Quarterly Health Center Performance Measure Reporting Tool. | 21 | 3 | 4 |
| | Annual Health Center Performance Measure Reporting Tool. | 21 | 1 | 6 |
| | Health Center Provider Survey | 126 | 1 | 20/60 |
| | Youth Serving Organization (YSO) Organizational Assessment. | 15 | 1 | 1 |
| | YSO Performance Measure Reporting Tool | 15 | 4 | 1 |
| | Youth Serving Organization (YSO) Staff Survey | 225 | 1 | 20/60 |
| | Awardee Training and Technical Assistance Tool | 3 | 12 | 2 |
| | Awardee Performance Measure Reporting Tool | 3 | 1 | 1 |
| | Health Center Youth Survey | 1050 | 1 | 10/60 |
| Individual | Health Center Organizational Assessment | 4 | 1 | 2 |
| | Quarterly Health Center Performance Measure Reporting Tool. | 4 | 3 | 4 |
| | Annual Health Center Performance Measure Reporting Tool. | 4 | 1 | 6 |
| | Health Center Provider Survey | 24 | 1 | 20/60 |
| | Youth Serving Organization (YSO) Organizational Assessment. | 20 | 1 | 1 |
| | YSO Performance Measure Reporting Tool | 20 | 4 | 1 |
| | Youth Serving Organization (YSO) Staff Survey | 300 | 1 | 20/60 |
| | State and Local Government | | | |

Jeffrey M. Zirger,

Acting Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Medicare & Medicaid Services, HHS**

[Document Identifiers: CMS-10286 and CMS-10488]

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 any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) 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 *August 11, 2016*:

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 <http://www.cms.hhs.gov/PaperworkReductionActof1995>.

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: Reports Clearance Office at (410) 786-1326.

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:* Revision of a currently approved information collection; *Title of Information Collection:* Notice of

Research Exception under the Genetic Information Nondiscrimination Act; *Use:* Under the Genetic Information Nondiscrimination Act of 2008 (GINA), a plan or issuer may request (but not require) a genetic test in connection with certain research activities so long as such activities comply with specific requirements, including: (i) The research complies with 45 CFR part 46 or equivalent federal regulations and applicable State or local law or regulations for the protection of human subjects in research; (ii) the request for the participant or beneficiary (or in the case of a minor child, the legal guardian of such beneficiary) is made in writing and clearly indicates that compliance with the request is voluntary and that non-compliance will have no effect on eligibility for benefits or premium or contribution amounts; and (iii) no genetic information collected or acquired will be used for underwriting purposes. The Secretary of Labor or the Secretary of Health and Human Services is required to be notified if a group health plan or health insurance issuer intends to claim the research exception permitted under Title I of GINA. Nonfederal governmental group health plans and issuers solely in the individual health insurance market or Medigap market will be required to file with the Centers for Medicare & Medicaid Services (CMS). The Notice of Research Exception under the Genetic Information Nondiscrimination Act is a model notice that can be completed by group health plans and health insurance issuers and filed with either the Department of Labor or CMS to comply with the notification requirement. *Form Number:* CMS-10286 (OMB Control Number 0938-1077); *Frequency:* Occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 2; *Total Annual Responses:* 2; *Total Annual Hours:* 1. (For policy questions regarding this collection contact Russell Tipps at 301-492-4371).

2. *Type of Information Collection Request:* Revision; *Title of Information Collection:* Consumer Experience Survey Data Collection; *Use:* Section 1311(c)(4) of the Affordable Care Act requires the Department of Health and Human Services (HHS) to develop an enrollee satisfaction survey system that assesses consumer experience with qualified health plans (QHPs) offered through an Exchange. It also requires public display of enrollee satisfaction information by the Exchange to allow individuals to easily compare enrollee satisfaction levels between comparable plans. HHS established the QHP

Enrollee Experience Survey (QHP Enrollee Survey) to assess consumer experience with the QHPs offered through the Marketplaces. The survey includes topics to assess consumer experience with the health care system such as communication skills of providers and ease of access to health care services. CMS developed the survey using the Consumer Assessment of Health Providers and Systems (CAHPS®) principles (<http://www.cahps.ahrq.gov/about.htm>) and established an application and approval process for survey vendors who want to participate in collecting QHP enrollee experience data.

The QHP Enrollee Survey, which is based on the CAHPS® Health Plan Survey, will (1) help consumers choose among competing health plans, (2) provide actionable information that the QHPs can use to improve performance, (3) provide information that regulatory and accreditation organizations can use to regulate and accredit plans, and (4) provide a longitudinal database for consumer research. CMS completed two rounds of developmental testing including 2014 psychometric testing and 2015 beta testing of the QHP Enrollee Survey. The psychometric testing helped determine psychometric properties and provided an initial measure of performance for Marketplaces and QHPs to use for quality improvement. Based on psychometric test results, CMS further refined the questionnaire and sampling design to conduct the 2015 beta test of the QHP Enrollee Survey. CMS obtained clearance for the national implementation of the QHP Enrollee Survey which is currently being conducted in 2016. At this time, CMS is requesting approval of adding six disability status items required by section 4302 of the Affordable Care Act and that were tested during the 2014 psychometric testing of the QHP Enrollee Survey. With the addition of these six questions, the revised total estimated annual burden hours of national implementation of the QHP Enrollee Survey is 37,823 hours with 105,015 responses. The revised total annualized burden over three years for this requested information collection is 113,469 hours and the total average annualized number of responses is 315,045 responses. *Form Number:* CMS-10488 (OMB control number 0938-1221). *Frequency:* Annually; *Affected Public:* Public Sector (Individuals and Household), Private Sector (business or other for-profit and not-for-profit institutions); *Number of Respondents:* 105,015; *Total Annual*

Responses: 105,015; *Total Annual Hours:* 37,823. (For policy questions regarding this collection contact Nidhi Singh Shah at 301-492-5110.)

Dated: July 7, 2016.

William N. Parham, III,
Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the National Preparedness and Response Science Board

AGENCY: Office of the Secretary, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) is hereby giving notice that the National Preparedness and Response Science Board (NPRSB) will be holding a public teleconference.

DATES: The NPRSB will hold a public meeting on July 29, 2016, from 4:00 p.m. to 5:00 p.m. EST. The agenda is subject to change as priorities dictate.

ADDRESSES: Individuals who wish to participate should send an email to NPRSB@HHS.GOV with "NPRSB Registration" in the subject line. The meeting will occur by teleconference. To attend via teleconference and for further instructions, please visit the NPRSB Web site at [HTTP://WWW.PHE.GOV/NPRSB](http://WWW.PHE.GOV/NPRSB).

FOR FURTHER INFORMATION CONTACT: Please submit an inquiry via the NPRSB Contact Form located at: <http://www.phe.gov/Preparedness/legal/boards/nprsb/Pages/RFNBSBComments.aspx>.

SUPPLEMENTARY INFORMATION: Pursuant to section 319M of the Public Health Service Act (42 U.S.C. 247d-7f) and section 222 of the Public Health Service Act (42 U.S.C. 217a), HHS established the NPRSB. The Board shall provide expert advice and guidance to the Secretary on scientific, technical, and other matters of special interest to HHS regarding current and future chemical, biological, nuclear, and radiological agents, whether naturally occurring, accidental, or deliberate. The NPRSB may also provide advice and guidance to the Secretary and/or the Assistant Secretary for Preparedness and Response on other matters related to