

Each PRC is housed within an accredited school of public health or an accredited school of medicine or osteopathy with a preventive medicine residency program. The PRCs conduct outcomes-oriented, applied prevention research on a broad range of topics using a multi-disciplinary and community-engaged approach. Each PRC receives funding from the CDC to establish its core infrastructure and functions and support a core research project. In addition to core research projects, most PRCs are awarded funding to complete special interest projects (SIPs) and conduct other research projects.

The DP14-001 program announcement included language that was used to develop and operationalize a set of 25 PRC Program evaluation indicators. The PRC Program logic model identifies program inputs, activities, outputs, and outcomes. The list of indicators was revised to better reflect program needs and capture PRCs' center and research activities, outputs, and outcomes.

The CDC is currently approved to collect information from the PRCs through a structured telephone interview and a web-based survey hosted by a third-party. The web-based survey is designed to collect information on the PRCs' collaborations with health departments; formal training programs and other training activities; and other-funded research

projects conducted separate from their core projects or SIP research. Structured telephone interviews with key PRC informants allow PRC Program staff to collect indicator data that do not lend themselves to a survey-based methodology and require a qualitative approach.

CDC requests OMB approval to revise the information collection plan as follows:

(1) The content of the web-based survey will be updated to more closely align with revised evaluation indicators. In addition, the web-based survey will be migrated from a third party platform to a web-based data collection system hosted on CDC servers. Although the estimated burden per response will increase, the revised data collection system will be comprehensive and will reduce the need for follow-up clarification by PRC Program awardees.

(2) CDC will discontinue telephone interviews and conduct key informant interviews (KII) every other year to capture qualitative information about PRC Network formation and cohesion.

CDC will continue to use the information reported by PRCs to identify training and technical assistance needs, respond to requests for information from Congress and other sources, monitor grantees' compliance with cooperative agreement requirements, evaluate progress made in achieving goals and objectives, and describe the impact and effectiveness of the PRC Program.

The CDC currently funds 26 PRCs. Each PRC will annually report the required information to the CDC. The annualized estimated burden is expected to increase. This increase equates to an estimated weekly burden of one hour per respondent and more fully accounts for the burden of preparing responses, as well as the burden of reporting responses. Web-based data collection will occur on an annual basis. The KIIs will take place in 2016 and 2018. This equates to two PRC Network KIIs per PRC Program awardee during the three year OMB approval period. Responses are annualized in the burden table below.

The proposed web-based data collection system will allow data entry during the entire year, which will enable respondents to distribute burden throughout each funding year. Response burden is expected to decrease in funding years 2 through 5, since the web-based data collection system will replicate a number of data elements from year-to-year, and respondents will only need to enter changes.

OMB approval is requested for three years. CDC plans to implement revised reporting requirements in March 2016. PRC Program awardees are required to participate in information collection. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1,299.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
Prevention Research Center ..	Web-based Data Collection	26	1	48
	Key Informant Interview: PRCs Network	17	1	3

Leroy A. Richardson,
*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

Administration for Children and Families

[OMB No. 0970-0410]

Proposed Information Collection Activity; Comment Request

Title: Tribal PREP Implementation Plan.

Description: This request to collect information for the Tribal PREP Implementation Plan, is due by July 1,

2017. This plan will contain the description of how the grantee intends to structure, measure and evaluate the implementation of the project. Information contained in this Implementation Plan will enable the Program Office to provide the necessary technical assistance to help ensure that grantees are structuring Tribal PREP projects within the framework of PREP design guidance, including mandated adult preparation subjects, Positive Youth Development and evidence-based programming.

Respondents:

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Tribal PREP Plan	10	1	40	400

Estimated Total Annual Burden Hours: 400.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: infocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Robert Sargis,
Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2015-N-4809]

Patient and Medical Professional Perspectives on the Return of Genetic Test Results and Interpretations; Public Workshop; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is announcing the following public workshop entitled "Patient and Medical Professional Perspectives on the Return of Genetic Test Results." The purpose of this public workshop is to understand patient and provider perspectives on receiving potentially medically relevant genetic test results. The topic(s) to be discussed will focus on better defining the specific information patients and providers prefer to receive, with an emphasis on the type(s) and amount of evidence available to interpret the results for medical purposes, how those results should be returned, and what information is needed to understand the results in the event that they could effectively aid in medical decision making.

DATES: The public workshop will be held on March 2, 2016, from 8 a.m. to 4 p.m. Submit either electronic or written comments on the public workshop by March 31, 2016.

ADDRESSES: The public workshop will be held at FDA's White Oak Campus, 10903 New Hampshire Ave., Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993. Entrance for the public meeting participants (non-FDA employees) is through Building 1, where routine security check procedures will be performed. For parking and security information, please refer to <http://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/ucm241740.htm>.

You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- Federal eRulemaking Portal: <http://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <http://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a

third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <http://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA-2015-N-4809 for "Patient and Medical Professional Perspectives on the Return of Genetic Test Results; Public Workshop; Request for Comments." Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <http://www.regulations.gov> or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION". The Agency will review this copy, including