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.

[FR Doc. 2016–14725 Filed 6–21–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Healthcare Infection Control Practices Advisory Committee (HICPAC)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announce the following meeting for the aforementioned committee:

Times and Dates: 9:00 a.m.–5:00 p.m., EDT, July 14, 2016; 9:00 a.m.–12:00 p.m., EDT, July 15, 2016.

Place: Centers for Disease Control and Prevention, Global Communications Center, Building 19, Auditorium B, 1600 Clifton Road NE., Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 300 people. Please register for the meeting at *www.cdc.gov/hicpac.*

Purpose: The Committee is charged with providing advice and guidance to the Director, Division of Healthcare Quality Promotion, the Director, National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), the Director, CDC, the Secretary, Health and Human Services regarding (1) the practice of healthcare infection prevention and control; (2) strategies for surveillance, prevention, and control of infections, antimicrobial resistance, and related events in settings where healthcare is provided; and (3) periodic updating of CDC guidelines and other policy statements regarding prevention of healthcare-associated infections and healthcare-related conditions.

Matters for Discussion: The agenda will include updates on CDC's activities for prevention of healthcare associated infections (HAIs), updates on antimicrobial stewardship, an update on Draft Guideline for Prevention of Infections in Healthcare Personnel, and an update from the workgroup for considerations on endoscope reprocessing.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Erin Stone, M.S., HICPAC, Division of Healthcare Quality Promotion, NCEZID, CDC, 1600 Clifton Road NE., Mailstop A–07, Atlanta, Georgia 30333 Telephone (404) 639–4045, Email: *hicpac@cdc.gov.*

The Director, Management Analysis and Services Office, 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.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2016–14788 Filed 6–21–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–16–16TL]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Health Risks from Using Private Wells for Drinking Water—New — National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Safe Drinking Water Act of 1974 (SDWA) ensures that most Americans are provided access to water that meets established public health standards. However, for over 38 million Americans who rely on private wells or other drinking water not protected by the SDWA (herein referred to as private wells), that is not the case. There is no comprehensive knowledge about the locations of private wells, the populations served by these sources, potential contaminants that might be present in private well water in specific areas of the country, or the potential health risks associated with drinking water from these sources.

The purpose of this new generic clearance information collection request is to assess the health risks associated with exposure to contaminants in drinking water from private wells across varied geographic areas of the United States in partnership with the requesting agency (state, territorial, local, or tribal health department). The information obtained from these investigations will be used to describe health risks from exposure to contaminants in drinking water from private wells within a defined time period and geographic distribution. This information will be used to inform public health protection activities conducted by the requesting agencies.

The respondents are defined as adults at least 18 years old, who use private wells for drinking water, who are willing to receive and return a tap water sampling kit and urine specimen kit or to provide a blood specimen, and who are willing to answer survey questions. They will be recruited from geographic areas of interest as defined by the requesting agency.

Based on our historical activities, we estimate that CDC will conduct up to 10

investigations per year. Each investigation will involve on average 200 respondents. The total time burden is 2,084 hours. There will be no cost to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
Adult at least 18 years old using a private well for tap water.	Screening Form	2,500	1	6/60
	Questionnaire	2,000	1	35/60
	Urine Specimen and Tap Water Sample Collection.	2,000	1	20/60

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.

[FR Doc. 2016–14724 Filed 6–21–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–D–1543]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on Nonproprietary Naming of Biological Products; Withdrawal

AGENCY: Food and Drug Administration, HHS.

ACTION: Withdrawal of notice.

SUMMARY: This document withdraws a Food and Drug Administration (FDA) notice that published in the **Federal Register** of June 2, 2016 (81 FR 35367).

DATES: This notice is withdrawn on June 22, 2016.

FOR FURTHER INFORMATION CONTACT: Howard Muller, Center for Drug Evaluation and Research (CDER), 10903 New Hampshire Ave., Bldg. 51, Rm. 6234, Silver Spring, MD 20993–0002, 301–796–3474.

SUPPLEMENTARY INFORMATION: FDA published a notice in the **Federal Register** of June 2, 2016, informing interested parties that the proposed collection of information entitled ''Guidance for Industry on Nonproprietary Naming of Biological Products'' had been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 and inviting the public to submit comments on the proposed collection to OMB. FDA is withdrawing the proposed collection of information that published on June 2, 2016, at this time.

Dated: June 16, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–14722 Filed 6–21–16; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995), the Health Resources and Services Administration (HRSA) announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than August 22, 2016.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Information Collection Clearance Officer, Room 14N–39, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call the HRSA Information Collection Clearance Officer at (301) 443–1984.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the information request collection title for reference.

Information Collection Request Title: Data Collection Tool for State Offices of Rural Health Grant Program

OMB No. 0915–0322—Extension

Abstract: The mission of the Federal Office of Rural Health Policy (FORHP) is to sustain and improve access to quality care services for rural communities. In its authorizing language (section 711 of the Social Security Act [42 U.S.C. 912]), Congress charged FORHP with administering grants, cooperative agreements, and contracts to provide technical assistance and other activities as necessary to support activities related to improving health care in rural areas. In accordance with the Public Health Service Act, section 338J (42 U.S.C. 254r), HRSA proposes to continue the State Offices of Rural Health (SORH) Grant Program—Funding Opportunity Announcement (FOA) and Forms for the Application. The FOA is used by 50 states in preparing applications for grants under the SORH Grant Program of the Public Health Service Act, and in preparing the required report.

Need and Proposed Use of the Information: FORHP seeks to continue gathering information from grantees on their efforts to provide technical assistance to clients within their states. SORH grantees submit a Technical Assistance Report that includes: (1) The total number of technical assistance encounters provided directly by the grantee, and (2) the total number of unduplicated clients that received direct technical assistance from the grantee. The Technical Assistance Report is submitted via the HRSA Electronic