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

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

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]  
BILLING CODE 4163–18–P

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DATES: This notice is withdrawn on June 22, 2016.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 338 (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