### ESTIMATED ANNUALIZED BURDEN HOURS—Continued

<table>
<thead>
<tr>
<th>Respondents</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
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<tr>
<td>Exploratory—HIV Communication and Awareness Focus Group Interview Guide.</td>
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<td>Consumer Concept Testing Focus Group Interview Guide.</td>
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<td>Consumer Message Testing Focus Group Interview Guide.</td>
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<td>Consumer Materials Testing Focus Group Interview Guide.</td>
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<td>HIV Prevention with Positives Survey.</td>
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<td>Intercept Interview Guide.</td>
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<td>1</td>
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</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. 2016–30408 Filed 12–16–16; 8:45 am]
BILLING CODE 4163–18–P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Centers for Disease Control and Prevention**  
[30Day–17–16AUE]

**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. Written comments and/or suggestions regarding the items contained in this notice should be directed 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**

Developing Effective Messages about Excessive Alcohol Consumption: Formative Focus Groups with Adult Drinkers and Abstainers—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

Excessive alcohol use, including binge drinking, is responsible for approximately 88,000 deaths in the U.S. annually—including one in 10 deaths among working-age adults ages 20–64. On average, for each death due to alcohol, an individual’s life is cut short by 30 years. Excessive alcohol use can also lead to motor vehicle crashes; intimate partner violence; and risky sexual behaviors, increasing the risk of HIV, other sexually transmitted infections, and unintended pregnancy. Over time, excessive alcohol use can lead to alcohol dependence, liver disease, high blood pressure, heart attack, stroke, and certain kinds of cancer. Furthermore, in 2010, excessive alcohol use cost the United States government $249 billion, or $2.05 per drink.

Binge drinking (defined as four or more drinks on an occasion for women or five or more drinks on an occasion for men) accounts for more than half of the deaths and three-quarters of the economic costs of excessive drinking. More than 38 million U.S. adults binge drink about four times a month, averaging eight drinks per binge. However, most (90%) binge drinkers are not alcohol dependent, presenting an opportunity for prevention through messages that improve voluntary compliance with recommended guidelines. States and communities can prevent binge drinking by supporting evidence-based strategies, such as those recommended by the Community Preventive Services Task Force; however, these strategies are underused.

Understanding the type of information and messages that the larger community—those who drink but not excessively or abstain from drinking in addition to those who engage in binge drinking—respond to will be essential in developing the communication strategy for future outreach.

CDC plans to collect information needed to improve understanding of current knowledge, perceptions, and attitudes related to excessive alcohol consumption. Respondents will be 72 adults ages 21–64 years who agree to participate in focus group discussions of about 1.5 hours each. A total of 12 focus groups are planned in three geographically diverse locations with appropriate facilities (four focus groups per location). Each focus group will involve six respondents and will be guided by a professional moderator.

Through an initial screening process, CDC will also collect the information needed to assess knowledge,
perceptions, and attitudes across various audience segments: those who engage in binge drinking, those who drink but not excessively, and those who abstain from drinking.

The focus group discussions will be analyzed using qualitative tools and leverage a structured approach to thematic analysis. Findings from this information collection will guide the CDC Alcohol Program in the development and refinement of targeted messages to effectively communicate the problem of excessive alcohol use, and encourage support for effective prevention strategies. The ultimate goal of the subsequent messaging is a reduction in binge drinking, which will in turn reduce alcohol-related injuries and deaths among adults.

OMB approval is requested for one year. Participation is voluntary and there are no costs to respondents other than their time. The total estimated annual burden hours are 132.

**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 hours)</th>
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<tbody>
<tr>
<td>Adults aged 21–64</td>
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<td>Focus Group</td>
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<tr>
<td>Total</td>
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<td></td>
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</table>

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10171]

Agency Information Collection Activities: Proposed Collection; 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 (the 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 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including any of the following subjects: the necessity and utility of the proposed information collection for the proper performance of the agency’s functions; the accuracy of the estimated burden; ways to enhance the quality, utility, and clarity of the information to be collected; and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by February 17, 2017.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. Electronically. You may send your comments electronically to http://www.regulations.gov. Follow the instructions for “Comment or Submission” or “More Search Options” to find the information collection document(s) that are accepting comments.

2. By regular mail. You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number ____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

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:


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:**

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection’s supporting statement and associated materials (see ADDRESSES).

CMS–10171 Collecting Benefit Coordination Data

Under the 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 requires federal agencies to publish a 60-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.

Information Collection

1. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Collecting Benefit Coordination Data; Use: This collection of information request coordinates Part D plan prescription drug coverage with other prescription drug coverage. The collected information will assist CMS, Part D