

questionnaire is referred to as the Standardized National Hypothesis Generating Questionnaire (SNHGQ). However, not all of the data elements in the SNHGQ are relevant to the parasite *Cyclospora* (e.g., questions about consumption of meat and dairy products); on the other hand, additional data elements (besides those in the SNHGQ) are needed to capture information pertinent to *Cyclospora* and to fresh produce vehicles of infection. Therefore, the Cyclosporiasis National Hypothesis Generating Questionnaire (CNHGQ) has been developed, by using core data elements from the SNHGQ and incorporating modifications pertinent to *Cyclospora*.

The core data elements from the SNHGQ were developed by a series of working groups comprised of local,

state, and federal public health partners. Subject matter experts at CDC have developed the CNHGQ, by modifying the SNHGQ to include and focus on data elements pertinent to *Cyclospora*/cyclosporiasis. Input also was solicited from state public health partners. Because relatively few data elements in the SNHGQ needed to be modified, a full vetting process was determined not to be necessary. The CNHGQ has been designed for administration over the telephone by public health officials, to collect data elements from case-patients or their proxies. The data that is collected will be pooled and analyzed at CDC, to generate hypotheses about potential vehicles/sources of infection.

CDC requests OMB approval to collect information via the CNHGQ from persons who have developed

symptomatic cases of *Cyclospora* infection during periods in which increased numbers of such cases are reported (typically, during spring and summer months). In part because molecular typing methods are not yet available for *C. cayetanensis*, it is important to interview all case-patients identified during periods of increased reporting, to help determine if their cases could be part of an outbreak(s).

The CNHGQ is not expected to entail substantial burden for respondents. The estimated total annualized burden associated with administering the CNHGQ is 750 hours (approximately 1,000 individuals interviewed x 45 minutes/response). There will be no costs to 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 (hours)
Individuals	Cyclosporiasis National Hypothesis Generating Questionnaire	1,000	1	45/60

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.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10147, CMS-10203, CMS-R-21, and CMS-R-148]

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 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 June 5, 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:

1. Access CMS' Web site address at <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html>.

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: William Parham at (410) 786-4669.

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-10147 Medicare Prescription Drug Coverage and Your Rights
- CMS-10203 Medicare Health Outcomes Survey (HOS)
- CMS-R-21 Withholding Medicare Payments to Recover Medicaid

Overpayments and Supporting Regulations in 42 CFR 447.31
 CMS–R–148 Limitations on Provider Related Donations and Health Care Related Taxes; Limitation on Payment to Disproportionate Share Hospitals; Medicaid and Supporting Regulations in 42 CFR 433.68, 433.74 and 447.272
 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:* Revision of a currently approved collection; *Title of Information Collection:* Medicare Prescription Drug Coverage and Your Rights; *Use:* Through the delivery of this standardized notice, Part D plan sponsors’ network pharmacies are in the best position to inform enrollees (at the point of sale) about how to contact their Part D plan if their prescription cannot be filled and how to request an exception to the Part D plan’s formulary. The notice restates certain rights and protections related to the enrollees Medicare prescription drug benefits, including the right to receive a written explanation from the drug plan about why a prescription drug is not covered; *Form Number:* CMS–10147 (OMB control number: 0938–0975); *Frequency:* Occasionally; *Affected Public:* Private sector (business or other for-profits); *Number of Respondents:* 62,000; *Total Annual Responses:* 40,100,000; *Total Annual Hours:* 668,066. (For policy questions regarding this collection contact Sabrina Sparkman at 410–786–3209.)

2. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Medicare Health Outcomes Survey (HOS); *Use:* The collection is necessary to hold Medicare managed care contracts accountable for the quality of care they deliver to

beneficiaries. This reporting requirement allows us to obtain the information necessary for proper oversight of the Medicare Advantage program. It is critical to our mission that we collect and disseminate valid and reliable information that can be used to improve quality of care through identification of quality improvement opportunities, assist us in carrying out our oversight responsibilities, and help beneficiaries make an informed choice among health plans. *Form Number:* CMS–10203 (OMB control number: 0938–0701); *Frequency:* Yearly; *Affected Public:* Individuals and households; *Number of Respondents:* 739,959; *Total Annual Responses:* 554,895; *Total Annual Hours:* 183,115. (For policy questions regarding this collection contact Kimberly DeMichele at 410–786–4286.)

3. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Withholding Medicare Payments to Recover Medicaid Overpayments and Supporting Regulations in 42 CFR 447.31; *Use:* Certain Medicaid providers that are subject to offsets for the collection of Medicaid overpayments may terminate or substantially reduce their participation in Medicaid, leaving the state Medicaid agency unable to recover the amounts due. Recovery procedures allow for determining the amount of overpayments and offsetting the overpayments by withholding the provider’s Medicare payments. To effectuate the withholding, the state agency must provide their respective CMS regional office with certain documentation that identifies the provider and the Medicaid overpayment amount. The agency must also demonstrate that the provider was notified of the overpayment and that demand for the overpayment was made. An opportunity to appeal the overpayment determination must be afforded to the provider by the Medicaid state agency. Lastly, Medicaid state agencies must notify CMS when to terminate the withholding; *Form Number:* CMS–R–21 (OMB control number: 0938–0287); *Frequency:* Occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 54; *Total Annual Responses:* 27; *Total Annual Hours:* 81. (For policy questions regarding this collection contact Stuart Goldstein at 410–786–0694.)

4. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Limitations on Provider Related Donations and Health

Care Related Taxes; Limitation on Payment to Disproportionate Share Hospitals; Medicaid and Supporting Regulations in 42 CFR 433.68, 433.74 and 447.272; *Use:* States may request a waiver of the broad based and uniformity tax program requirements. Each state must demonstrate that its tax program(s) do not violate the hold harmless provision. Additionally, state Medicaid agencies must report (quarterly) on health care related taxes collected and the source of provider related donations received by the state or unit of local government. Each state must maintain, in readily reviewable form, supporting documentation that provides a detailed description of each donation and tax program being reported, as well as the source and use of all donations received and collected. Without this information, the amount of Federal financial participation payable to a state cannot be determined; *Form Number:* CMS–R–148 (OMB control number: 0938–0618); *Frequency:* Quarterly and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 50; *Total Annual Responses:* 40; *Total Annual Hours:* 3,200. (For policy questions regarding this collection contact Stuart Goldstein at 410–786–0694.)

Dated: April 3, 2017.

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

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10501 and CMS–10635]

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