ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Law enforcement officers	Actigraph fitting and return	60	3	10/60	30
Total					389

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. 2017–22201 Filed 10–12–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–17–17ND]

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. CDC previously published a ''Proposed Data Collection Submitted for Public Comment and Recommendations'' notice on February 10, 2017 to obtain comments from the public and affected agencies. CDC received one comment related to the previous notice. The purpose of this notice is to allow an additional 30 days for public comments.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. The Office of Management and Budget is particularly interested in comments that:

(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, 725 17th Street NW., Washington, DC 20503 or by fax to (202) 395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Annual Progress Report (APR) for Injury Control Research Centers (ICRC)—New—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Injury Control Research Centers (ICRCs) form a national network of ten comprehensive academic research centers that focus on three core functions: Research, training, and outreach. ICRCs are on the scientific front line conducting cutting-edge, multidisciplinary research on the causes, outcomes, and prevention of injuries and violence.

ICRC research focuses on issues of local and national importance including motor vehicle injuries; interpersonal violence and suicide; opioid overdoses; older adult falls; and traumatic brain injuries. ICRCs work with states and communities to ensure research is put into action to prevent injuries and violence. They provide technical assistance to disseminate and translate research findings which leads to increased awareness and influences action. ICRCs play a critical role training and developing the current and next generation of researchers and public health professionals. This helps ensure there is an adequate supply of qualified practitioners and researchers to advance prevention research, address new problems, and reach new populations across the nation.

The CDC seeks OMB approval for three years to collect Annual Progress Report (APR) information from 10 grantees funded under Grants for Injury Control Research Centers (ICRC). ICRC awardees will report activity information to CDC annually using three fillable electronic templates. The first Word-based template is the principal tool for the Indicators Data Collection (IDC), which is based on a set of program activity indicators and key ICRC evaluation questions. The second Word-based template collects information about non-CDC-funded studies, and the third template, which is Excel-based, collects information about ICRC personnel and publications. Information will be reported electronically to the NCIPC for program monitoring, and hard copies will be submitted to CDC's Office of Financial Resources (OFR). Together, the tools describe grantees' annual goals, objectives, progress, and performance towards overall cooperative agreement aims. The tools also describe how grantees implement and use evidence-based injury prevention and control strategies.

Information to be collected will provide crucial data for program performance monitoring, will allow CDC to analyze and synthesize information from grantees, help ensure consistency in documenting progress and technical assistance, enhance accountability of the use of federal funds, and provide timely reports as frequently requested by the Department of Health and Human Services, the White House, and Congress.

Submission of the Annual Progress Report is required for cooperative agreement grantees. The total estimated annualized burden hours are 500. There is no cost 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 (in hours)
Injury Control Research Center (ICRC) Grantees.	ICRC Indicators Data Collection	10	1	20
(ICRC) Grantees	ICRC Indicators Data Collection: Non-CDC Study Supplement.	10	1	10
	ICRC Personnel and Publication Excel Data Collection.	10	1	20

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 Identifier: CMS–10305]

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 December 12, 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 *http://www.cms.hhs.gov/PaperworkReductionActof1995.*

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–10305 Medicare Part C and Part D Data Validation (42 CFR 422.516(g) and 423.514(g))

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 Part C and Part D Data Validation (42 CFR 422.516(g) and 423.514(g)); *Use:* Medicare Part C and Part D sponsoring organizations (Medicare Advantage Organizations), must submit Medicare Part C, Medicare Part D, or Medicare Part C and Part D data (depending on the type of contracts they have in place with CMS). In order for the reported data to be useful for monitoring and performance measurement, the data must be reliable, valid, complete, and comparable among sponsoring organizations. To maintain the independence of the validation process, sponsoring organizations are responsible for hiring external, independent data validation contractors (DVCs) who meet a minimum set of qualifications and credentials. For the retrospective review in 2018, the DVCs will review data submitted by sponsoring organizations for CY2017. The main changes for the 2018 DV are to eliminate the Part C/D reporting section Sponsor Oversight of Agents and adding the Part D reporting section Improving Drug Utilization Review