

implemented. The study population for the primary care provider post-discharge questionnaire will be Primary Care Providers (PCP) associated with the same Medical Center who care for older adult study patients discharged each month. Four questionnaires will be administered. (1) The Pre-discharge patient questionnaire will be used to survey older adults in the hospital (before discharge). (2) The Post-

discharge patient questionnaire will be used to survey the older adults that completed the pre-discharge survey three additional times (at 14, 30 and 60 days) after being discharged from the Medical Center. (3) The Clinical staff evaluation questionnaire will be used to survey clinical staff at the Medical Center. (4) The Primary Care Provider (PCP) post-discharge questionnaire will be used to survey primary care

providers involved in the care of patients discharged. The open-ended questions will be analyzed to identify themes, and results will be presented by theme. Frequencies, cross-tabs, and regression analysis will be used for categorical questions.

The total estimated annualized burden hours is 622. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours (in hours)
Older adult Patients	Survey correspondence to patients and consent form for patients.	2,299	1	2/60	77
	Pre-discharge Patient	800	1	10/60	133
	Post-discharge Patient	800	3	10/60	400
Clinical staff	Survey correspondence to clinical staff.	100	1	1/60	2
	Clinical staff evaluation Questionnaire.	50	1	5/60	4
Primary care providers (PCP)	Survey correspondence to primary care providers.	100	1	1/60	2
	PCP post discharge survey	50	1	5/60	4
Total	622

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60-Day-19-191J; Docket No. CDC-2018-0118]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS)

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a

proposed information collection project titled Improving Performance Measurement and Monitoring by CDC programs. The purpose of this project is to evaluate the progress of CDC partners that receive awards distributed via cooperative agreements from the Office of Grants Services (OGS)

DATES: CDC must receive written comments on or before April 8, 2019.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2018-0118 by any of the following methods:

- **Federal eRulemaking Portal:** *Regulations.gov.* Follow the instructions for submitting comments.
- **Mail:** Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road, NE, MS-D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *Regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office,

Centers for Disease Control and Prevention, 1600 Clifton Road, NE, MS-D74, Atlanta, Georgia 30329; phone: 404-639-7570; Email: omb@cdc.gov.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. 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;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information,

including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected; and

4. 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 submissions of responses.

5. Assess information collection costs.

Proposed Project

Improving Performance Measurement and Monitoring by CDC programs—New—Office of Grant Services (OGS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Each year, 75% of CDC funding goes to extramural organizations, including state and local partners, via contracts, grants, and, most commonly, cooperative agreements. A cooperative agreement is an award mechanism used when there will be substantial Federal programmatic involvement, meaning that the CDC program staff will collaborate or participate in project or program activities. These funds are distributed from the Office of Grant Services (OGS) to partners throughout the world to promote health, prevent disease, injury and disability and prepare for new health threats. OGS is responsible for the stewardship of these

funds while providing excellent, professional services to our partners and stakeholders.

Currently, CDC uses the PPMR (OMB Control Number- 0920–1132, Expiration Date: 08/31/2019), a progress report form adapted from an information collection owned by the Administration for Children and Families (ACF). This tool may be used to collect information periodically from recipients of CDC funds regarding the progress made on CDC funded projects.

The Improving Performance Measurement and Monitoring by CDC Programs project will work with up to 25 CDC programs developing cooperative agreements to address the challenges they face with performance planning, measurement and monitoring. Each cooperative agreement will provide funding to an average of 35 local entities, for a total of up to 875 locally funded entities.

Through participation in this Project, CDC programs and recipients of cooperative agreement funds will: (1) Develop strong performance measurement systems and practices; (2) define and operationalize priority performance measures tailored to a specific cooperative agreement; and (3) establish common data collection and reporting expectations across all recipients for a specific cooperative agreement. The Project focuses on addressing these issues during the early stages of cooperative agreement development and implementation.

The Project proposes a generic clearance adapted from a previously

approved generic clearance (OMB approval number: 0970–0490, expiration date 1/31/2020) owned by ACF. This ACF generic clearance replaces the information collection that is the basis of CDC’s current PPMR. Project participants will customize a sample information collection to meet program-specific needs.

The information collected will enable the accurate, reliable, uniform and timely submission to CDC of each recipient’s progress and performance measures. The information collected by the generic information collection is designed to align with, and support the goals outlined for each of the CDC recipients. Collection and reporting of the information will occur in an efficient, standardized, and user-friendly manner that will generate a variety of routine and customizable reports. The generic information collection will allow each recipient to summarize activities and progress towards meeting performance measures and goals over a specified time period specific to each award. CDC will also have the capacity to generate reports that describe activities across multiple recipients. In addition, CDC will use the information collected to respond to inquiries from HHS, Congress and other stakeholder inquiries about program activities and their impact. CDC requests OMB approval for three years. The total estimated burden is 35,000 hours. 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 per year	Average burden per response (in hours)	Total burden (in hours)
CDC Award Recipients ..	Performance Measuring and Monitoring Project Information Collection Tool.	875	1	40	35,000
Total	35,000

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30-Day–19–18AVU]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled Assessment of

Outcomes Associated with the Preventive Health and Health Services Block Grant to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on September 6, 2018 to obtain comments from the public and affected agencies. CDC did not receive comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.