

Additionally, half the sample will receive additional content related to effects that EHRs have on clinical workflow, efficiencies, and may address issues of access, quality, and costs of associated with the delivery of health care. Subsequent years of the entire 2014 NEHRS may receive longitudinal follow-up to evaluate the effect of EHR on the delivery of health care over time.

Users of NAMCS NEHRS data include, but are not limited to,

Congressional offices, Federal agencies, state and local governments, schools of public health, colleges and universities, private industry, nonprofit foundations, professional associations, clinicians, researchers, administrators, and health planners.

NAMCS NEHRS will survey 10,302 physicians a year, for eligibility. It is expected that all physicians will participate in an interview annually. In 2014, one-half of the physicians will

receive the regular NAMCS NEHRS and one-half of the physicians will receive an expanded NAMCS NEHRS. All the 2014 eligible physicians (10,302) will be asked to take the follow-up NAMCS NEHRS in 2015 and 2016.

There are no costs to the respondents other than their time. The total estimated annual burden hours are 7,155.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Responses per respondent	Average burden per response (in hours)
Office-based physicians	Regular NAMCS NEHRS	8,585	1	20/60
	Expanded NAMCS NEHRS	1,717	1	30/60
	NAMCS NEHRS expansion (Follow-up)	6,868	1	30/60

Leroy A. Richardson,

Chief, Information Collection Review Office,
Office of Scientific Integrity, Office of the
Associate Director for Science, Office of the
Director, Center for Disease Control and
Prevention.

[FR Doc. 2014-02667 Filed 2-6-14; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-14-13YQ]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call (404) 639-7570 or send an email to omb@cdc.gov. Send written comments to 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

Institutional Awareness and
Commitment to Ensuring Safe, Stable,

and Nurturing Relationships and
Environments for Children and
Prevention Child Maltreatment—New—
National Center for Injury Prevention
and Control (NCIPC)—Centers for
Disease Control and Prevention (CDC).

Background and Brief Description

Safe, stable, nurturing relationships and environments set children on a positive trajectory for optimal child development and health, provide a buffer against the effects of adverse child experiences, are fundamental to healthy brain development and have a positive impact on a broad range of health problems across the life course. Promoting safe, stable, nurturing relationships and environments may also reduce child maltreatment which is a significant public health problem affecting physical and emotional health throughout the lifespan.

NCIPC has funded five state health departments in Fiscal Year 2014 to coordinate and manage existing and new partnerships with other sectors to promote safe, stable, nurturing relationships and environments for children; and work with partners to identify strategies across sectors that promote safe, stable, nurturing relationships and environments. CDC requests OMB approval for two years to collect information that will establish the baseline level of state health departments' and partners' awareness and commitment to ensuring safe,

stable, and nurturing relationships and environments for children and preventing child maltreatment.

Information will be collected over a 2-year period from 3 staff members from each of the 5 health departments (15 respondents), and 3 staff members from each of the 5 health departments' 10 partner organizations (150 respondents)—for a total of 165 respondents (83 respondents per year). Information will be collected once using SurveyMonkey®, an electronic web-based interface which is a *secure Web site that meets the Safe Harbor and European Union data protection requirements*. This ICR will only collect data pertaining to organizations. No individual identifiable information will be requested.

Each grantee will receive a personalized advance notification letter, followed by an email with a link to the SurveyMonkey® site. In turn, the grantee will send a personalized advance notification letter, followed by an email with a link to the SurveyMonkey® site to each new partner throughout the funding period. CDC will use this information to establish state health departments' and partners' level of awareness and commitment at the start of the funding period.

There are no costs to respondents other than their time. The total estimated annual burden hours are 39.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)
Health Departments	Institutional awareness and commitment survey	8	1	28/60
Partner Organizations	Institutional awareness and commitment survey	75	1	28/60

Leroy A. Richardson,

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[FR Doc. 2014-02670 Filed 2-6-14; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10499]

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: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by April 8, 2014.

ADDRESSES: When commenting, please reference the document identifier or

OMB control number (OCN). 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: 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-10499 Public Health Agency/Registry Readiness To Support Meaningful Use

Under the Paperwork Reduction Act (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 Collections

1. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* Public Health Agency/Registry Readiness to Support Meaningful Use; *Use:* The Medicare and Medicaid Electronic Health Record (EHR) Incentive Programs provide incentives for the meaningful use of Certified Electronic Health Record Technology (CEHRT). We defined meaningful use as a set of objectives and measures in either Stage 1 or Stage 2 depending on how long an eligible provider has participated in the program. Both Stage 1 (3 objectives) and Stage 2 (5 objectives) of meaningful use contain objectives and measures that require eligible providers to determine the readiness of public health agencies and registries to receive electronic data from CEHRT. Public comments on the notice of proposed rulemaking for Stage 2 of meaningful use (77 FR 13697) asserted that the burden for each individual eligible provider to determine the readiness of multiple public health agencies and registries could be nearly eliminated if we were to maintain a database on the readiness of public health agencies and registries. In the final rule for Stage 2 of meaningful use (77 FR 53967), we agreed that the burden on eligible providers, public health agencies and registries would be greatly reduced and established that we would create such a database and it would serve as the