

Cancer Prevention and Control Research Network. The survey was found to be useful by CDC and the grantees (which received feedback reports). For example, after the each survey administration, CDC was able to tailor sessions at the Program Director's meeting to the needs of grantees that had been expressed during last year's information collection. DCPC has decided to continue the data collection, and is being supported through the National Association of Chronic Disease Directors. CDC's proposed survey builds on previous information collections

conducted from 2011–2013 through the CPRN.

Questions are of various types including dichotomous and multiple response. All information is to be collected electronically through the web-based survey. The estimated burden per response is 75 minutes.

This assessment will enable CDC to gauge its progress in meeting CRCCP program goals, identify implementation activities, monitor program transition to efforts aimed at impacting population-based screening, identify technical assistance needs of state, tribe and territorial health department cancer

control programs, and identify implementation models with potential to expand and transition to new settings to increase program impact and reach. The assessment will identify successful activities that should be maintained, replicated, or expanded as well as provide insight into areas that need improvement.

OMB approval is requested for three years. Participation is voluntary for CRCCP awardees and there are no costs to respondents other than their time. The total estimated annualized burden hours are 36.

ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondents | Form name | Number of respondents | Number of responses per respondent | Average burden per response (in hrs.) |
|--|---|-----------------------|------------------------------------|---------------------------------------|
| Colorectal Cancer Control Program Directors or Managers. | Colorectal Cancer Control Program (CRCCP) Grantee Survey of Program Implementation. | 29 | 1 | 75/60 |

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.

[FR Doc. 2015-05211 Filed 3-5-15; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Health Resources and Services Administration (HRSA) has submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period.

DATES: Comments on this ICR should be received no later than April 6, 2015.

ADDRESSES: Submit your comments, including the Information Collection Request Title, to the desk officer for HRSA, either by email to OIRA_submission@omb.eop.gov or by fax to 202-395-5806.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email the HRSA Information Collection Clearance Officer at paperwork@hrsa.gov or call (301) 443-1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Voluntary Partner Surveys to Implement Executive Order 12862 in the Health Resources and Services Administration.

OMB No. 0915-0212—Extension.

Abstract: In response to Executive Order 12862, the Health Resources and Services Administration (HRSA) proposes to conduct voluntary customer surveys of its partners to assess strengths and weaknesses in program services and processes. HRSA partners are typically state or local governments, health care facilities, health care consortia, health care providers, and researchers. HRSA is requesting a generic approval from OMB to conduct the partner surveys.

Partner surveys to be conducted by HRSA might include, for example, online or telephone surveys of grantees to determine satisfaction with grant processes or technical assistance provided by a contractor, or in-class evaluation forms completed by providers who receive training from

HRSA grantees, to measure satisfaction with the training experience. Results of these surveys will be used to plan and redirect resources and efforts as needed to improve services and processes.

Focus groups may also be used to gain partner input into the design of mail and telephone surveys. Focus groups, in-class evaluation forms, mail surveys, and telephone surveys are expected to be the preferred data collection methods.

A generic approval allows HRSA to conduct a limited number of partner surveys without a full-scale OMB review of each survey. If generic approval is approved, information on each individual partner survey will not be published in the **Federal Register**.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

| Form name | Number of respondents | Number of responses per respondent | Total responses | Average burden per response (in hours) | Total burden hours |
|-------------------------------------|-----------------------|------------------------------------|-----------------|--|--------------------|
| In-class evaluations | 40,000 | 1 | 40,000 | .05 | 2,000 |
| Mail/Telephone/Online Surveys | 12,000 | 1 | 12,000 | .25 | 3,000 |
| Focus groups | 250 | 1 | 250 | 1.5 | 375 |
| Total | 52,250 | 1 | 52,250 | .103 | 5,375 |

Jackie Painter,

Director, Division of the Executive Secretariat.

[FR Doc. 2015-05187 Filed 3-5-15; 8:45 am]

BILLING CODE 4165-15-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Medicare & Medicaid Services****[Document Identifier: CMS-1557, CMS-10531 and CMS-10535]****Agency Information Collection Activities: Submission for OMB Review; Comment Request****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 extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer April 6, 2015.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured

consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395-5806 or Email: OIRA_submission@omb.eop.gov.

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 the 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: 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. 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 (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-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 that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Survey Report

Form for Clinical Laboratory Improvement Amendments (CLIA) and Supporting Regulations; *Use:* The form is used to report surveyor findings during a CLIA survey. For each type of survey conducted (*i.e.*, initial certification, recertification, validation, complaint, addition/deletion of specialty/subspecialty, transfusion fatality investigation, or revisit inspections) the Survey Report Form incorporates the requirements specified in the CLIA regulations. *Form Number:* CMS-1557 (OMB control number: 0938-0544); *Frequency:* Biennially; *Affected Public:* Private sector (Business or other for-profit and Not-for-profit institutions, State, Local or Tribal Governments and Federal Government); *Number of Respondents:* 19,051; *Total Annual Responses:* 9,526; *Total Annual Hours:* 4,763. (For policy questions regarding this collection contact Kathleen Todd at 410-786-3385).

2. *Type of Information Collection Request:* New collection (Request for a new OMB control number); *Title of Information Collection:* Transcatheter Mitral Valve Repair (TMVR) National Coverage Decision (NCD); *Use:* The data collection is required by the Centers for Medicare and Medicaid Services (CMS) National Coverage Determination (NCD) entitled, "Transcatheter Mitral Valve Repair (TMVR)". The TMVR device is only covered when specific conditions are met including that the heart team and hospital are submitting data in a prospective, national, audited registry. The data includes patient, practitioner and facility level variables that predict outcomes such as all-cause mortality and quality of life.

We find that the Society of Thoracic Surgery/American College of Cardiology Transcatheter Valve Therapy (STS/ACC TVT) Registry, one registry overseen by the National Cardiovascular Data Registry, meets the requirements specified in the NCD on TMVR. The TVT Registry will support a national surveillance system to monitor the safety and efficacy of the TMVR technologies for the treatment of mitral regurgitation (MR). The data will also