of lading, the shipment was ''impounded in Algiers, Algeria for approximately four months.'' Complainant alleges that this error resulted in costs for which complainant would not have otherwise been responsible. Complainant alleges that it is ''subject to injury as a result of the violations by respondent of sections 46 U.S.C. code § 41104 and more specifically paragraphs 4 and 5.''

Complainant seeks reparations in the amount of $21,086.70, and other relief. The full text of the complaint can be found in the Commission's Electronic Reading Room at *www.fmc.gov/17-06/.*

This proceeding has been assigned to the Office of Administrative Law Judges. The initial decision of the presiding officer in this proceeding shall be issued by June 8, 2018, and the final decision of the Commission shall be issued by December 21, 2018.

Rachel E. Dickon,

Assistant Secretary.

[FR Doc. 2017–12296 Filed 6–13–17; 8:45 am]

BILLING CODE 6731–AA–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–17–1015]

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. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (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, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

The National Electronic Health Records Survey (NEHRS) (OMB Control No. 0920–1015, Expires 04/30/2017)—Reinstatement with Change—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C. 242k), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on ''utilization of health care'' in the United States. NEHRS was originally designed as a mail supplement to the National Ambulatory Medical Care Survey (NAMCS). Questions in NEHRS have been asked in NAMCS starting in 2001.

The purpose of NEHRS is to measure progress toward goals for electronic health records (EHRs) adoption. NEHRS target universe consists of all non-Federal office-based physicians (excluding those in the specialties of anesthesiology, radiology, and pathology) who are engaged in direct patient care.

NEHRS is the principal source of data on national and state-level EHR adoption in the United States. In 2008 and 2009, the sample size was 2,000 physicians annually. Starting in 2010, the annual sample size was increased five-fold, from 2,000 physicians to 10,302 physicians. The increased sample size allows for more reliable national estimates as well as state-level estimates on EHR adoption without having to be combined with NAMCS. For these reasons, in 2012 NEHRS became an independent survey, not as a supplement under NAMCS.

NEHRS collects information on characteristics of physician practices, the capabilities of EHRs in those practices, and intent to apply for meaningful use incentive payments. These data, together with trend data, may be used to monitor the adoption of EHR as well as accessing factors associated with EHR adoption. In 2017, a set of follow-up questionnaires will be incorporated into the survey that focuses on content related to physician attitudes on using EHRs.

Users of 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. There is no cost to the respondents other than their time. The total estimated annualized burden hours are 6,295.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (hours)
Office-based physicians	NEHRS	10,302	1	30/60
Office-based physicians	Follow-up NEHRS Elec Resp	858	1	20/60
Office-based physicians	Follow-up NEHRS Non-Elec Resp	859	1	20/60
Office-based physicians	Follow-up NEHRS Nonresp	1,717	1	20/60

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. 2017–12272 Filed 6–13–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10652]

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 August 14, 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–10652 Virtual Groups for Merit-Based Incentive Payment System (MIPS)

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:* New collection of information request; *Title of Information Collection:* Virtual Groups for Merit-Based Incentive Payment System (MIPS); *Use:* CMS acknowledges the unique challenges that small practices and practices in rural areas may face with the implementation of the Quality Payment Program. To help support these practices and provide them with additional flexibility, CMS has created a virtual group reporting option starting with the 2018 MIPS performance period. CMS held webinars and small, interactive feedback sessions to gain insight from clinicians as we developed our policies on virtual groups. During these sessions, participants expressed a strong interest in virtual groups, and indicated that the right policies could minimize clinician burden and bolster clinician success.

This information collection request is related to the statutorily required virtual group election process proposed in the CY 2018 Quality Payment Program proposed rule. A virtual group is a combination of Tax Identification Numbers (TINs), which would include at least two separate TINs associated with a solo practitioner TIN and National Provider Identifier (TIN/NPI) or group with 10 or fewer MIPS eligible clinicians and another solo practitioner (TIN/NPI) or group with 10 or fewer MIPS eligible clinicians.

Section 1848(q)(5)(I) of the Act requires that CMS establish and have in place a process to allow an individual MIPS eligible clinician or group consisting of not more than 10 MIPS eligible clinicians to elect, with respect to a performance period for a year to be in a virtual group with at least one other such individual MIPS eligible clinician or group. The Act also provides for the use of voluntary virtual groups for certain assessment purposes, including the election of practices to be a virtual group and the requirements for the election process.

Section 1848(q)(5)(I)(i) of the Act also provides that MIPS eligible clinicians electing to be a virtual group must: (1) Have their performance assessed for the quality and cost performance categories in a manner that applies the combined performance of all the MIPS eligible clinicians in the virtual group to each MIPS eligible clinician in the virtual group for the applicable performance period; and (2) be scored for the quality and cost performance categories based on such assessment.

CMS will use the data collected from virtual group representatives to determine eligibility to participate in a virtual group, approve the formation of that virtual group, based on determination of each TIN size, and assign a virtual group identifier to the virtual group. The data collected will