SUMMARY OF ANNUAL BURDEN

Information collection description	Type of burden	Obligation to respond	Estimated number of respondents	Estimated frequency of responses	Estimated time per response (hours)	Estimated annual burden (hours)
Interagency Notice of Change in Director or Executive Officer.	Reporting	Mandatory	107	On Occasion	2	214

Total Estimated Annual Burden: 214 hours.

General Description of Collection: Section 32 of the FDIA (12 U.S.C. 1831i) requires an insured depository institution or depository institution holding company under certain circumstances to notify the appropriate federal banking agency of the proposed addition of any individual to the board of directors or the employment of any individual as a senior executive officer of such institution at least 30 days before such addition or employment becomes effective. Section 32 of the FDIA also provides that the FDIC may disapprove an individual's service as a director or senior executive officer of certain state nonmember banks or state savings associations if, upon assessing the individual's competence, experience, character, and integrity, it is determined that the individual's service would not be in the best interest of the depositors of the institution or the public. The Interagency Notice of Change in Director or Senior Executive Officer, with the information contained in the Interagency Biographical and Financial Report (described above) as an attachment, is used by the FDIC to collect information relevant to assess the individual's competence, experience, character, and integrity.

Request for Comment

Comments are invited on: (a) Whether the collection of information is necessary for the proper performance of the FDIC's functions, including whether the information has practical utility; (b) the accuracy of the estimates of the burden of the information collection, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. All comments will become a matter of public record.

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on April 7, 2021.

James P. Sheesley,

Assistant Executive Secretary.

[FR Doc. 2021–07498 Filed 4–12–21; 8:45 am]

BILLING CODE 6714–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended, and the Determination of the Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—PAR 18–812, NIOSH Member Conflict Review.

Date: June 24, 2021.

Time: 1:00 p.m.–4:00 p.m., EDT.

Place: Teleconference.

Agenda: To review and evaluate grant applications.

For Further Information Contact: Michael Goldcamp, Ph.D., Scientific Review Officer, Office of Extramural Programs, National Institute for Occupational Safety and Health, CDC, 1095 Willowdale Road, Morgantown, West Virginia 26506, Telephone (304) 285–5951; *MGoldcamp@cdc.gov.*

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2021–07487 Filed 4–12–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–718–721, CMS–724, CMS–2088–17 and CMS–1763]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 (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 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 on the collection(s) of information must be received by the OMB desk officer by *May 13, 2021.*

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

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' website address at website address at: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

FOR FURTHER INFORMATION CONTACT: William Parham at (410) 786–4669.

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:* Revision of a currently approved collection; *Title of Information Collection:* Business Proposal Forms for Quality Improvement Organizations (QIOs); *Use:* The submission of proposal information by current quality improvement associations (QIOs) and other bidders, on the appropriate forms, will satisfy our need for meaningful, consistent, and verifiable data with which to evaluate contract proposals. We use the data collected on the forms associated with this information collection request to negotiate QIO contracts. We will be able to compare the costs reported by the QIOs on the cost reports to the proposed costs noted on the business proposal forms. Subsequent contract and modification negotiations will be based on historic cost data. The business proposal forms will be one element of the historical cost data from which we can analyze future proposed costs. In addition, the business proposal format will standardize the cost proposing and pricing process among all QIOs. With well-defined cost centers and line items, proposals can be compared among QIOs for reasonableness and appropriateness. *Form Number:* CMS–718–721 (OMB control number: 0938–0579); *Frequency:* Annually; *Affected Public:* Business or other for-profits and Not-for-profit institutions; *Number of Respondents:* 58; *Total Annual Responses:* 58; *Total Annual Hours:* 2,320. (For policy questions regarding this collection contact Benjamin Bernstein at 410–786–6570.)

2. *Type of Information Collection Request:* Reinstatement without change of a previously approved collection; *Title of Information Collection:* Medicare/Medicaid Psychiatric Hospital Survey Data and Supporting Regulations; *Use:* The CMS–724 form is used to collect data that assists us in program planning and evaluation and in maintaining an accurate database on providers participating in the psychiatric hospital program. Specifically, we use the information collected on this form in evaluating the Medicare psychiatric hospital program. The form is also used for audit purposes; determining patient population and characteristics of the hospital; and survey term composition. *Form Number:* CMS–724 (OMB control number: 0938–0378); *Frequency:* Annually; *Affected Public:* Business or other for-profits and Not-for-profit institutions; *Number of Respondents:* 19; *Total Annual Responses:* 191; *Total Annual Hours:* 96. (For policy questions regarding this collection contact Caroline Gallaher at 410–786–8705.)

3. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Community Mental Health Center Cost Report *Use:* CMS requires the Form CMS–2088–17 to determine a provider's reasonable cost incurred in furnishing medical services to Medicare beneficiaries and reimbursement due to or from a provider. In addition, CMHCs may receive reimbursement through the cost report for Medicare reimbursable bad debts. CMS uses the Form CMS–2088–17 for rate setting; payment refinement activities, including market basket analysis; Medicare Trust Fund projections; and to support program operations. The primary function of the cost report is to determine provider reimbursement for services rendered to Medicare beneficiaries. Each CMHC submits the cost report to its contractor for reimbursement determination.

Section 1874A of the Act describes the functions of the contractor. CMHCs must follow the principles of cost reimbursement, which require they maintain sufficient financial records and statistical data for proper determination of costs. The S series of worksheets collects the provider's location, CBSA, date of certification, operations, and unduplicated census days. The A series of worksheets collects the provider's trial balance of expenses for overhead costs, direct patient care services, and non-revenue generating cost centers. The B series of worksheets allocates the overhead costs to the direct patient care and non-revenue generating cost centers using functional statistical bases. The Worksheet C computes the apportionment of costs between Medicare beneficiaries and other patients. The D series of worksheets are Medicare specific and calculate the reimbursement settlement for services rendered to Medicare beneficiaries. The Worksheet F collects the provider's revenues and expenses data from the provider's income statement. *Form Number:* CMS–2088–17 (OMB control number: 0938–0378); *Frequency:* Annually; *Affected Public:* Private Sector, Business or other for-profits, Not-for-profits institutions; *Number of Respondents:* 184; *Total Annual Responses:* 184; *Total Annual Hours:* 16,560. (For policy questions regarding this collection contact Jill Keplinger at 410–786–4550.)

4. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Request for Termination of Premium-Hospital and or Supplementary Medical Insurance; *Use:* Form CMS–1763 provides the necessary information to process the enrollee's request for termination of Part B and/or premium Part A coverage.

Sections 1818(c)(5), 1818A(c)(2)(B) and 1838(b)(1) of the Act and corresponding regulations at 42 CFR 406.28(a) and 407.27(c) require that a Medicare enrollee wishing to voluntarily terminate Part B and/or premium Part A coverage file a written request with CMS or SSA. The statute and regulations also specify when coverage ends based upon the date the request for termination is filed.

Form CMS–1763 collects the information necessary to process Medicare enrollment terminations. The Request for Termination of Premium

Hospital and/or Supplementary Medical Insurance (Form CMS–1763) provides a standardized means to satisfy the requirements of law, as well as allow both agencies to protect the individual from an inappropriate decision. *Form Number:* CMS–1763 (OMB control number: 0938–0025); *Frequency:* Annually; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 114,215; *Total Annual Responses:* 114,215; *Total Annual Hours:* 19,074. (For policy questions regarding this collection contact Carla Patterson at 410–786–1000.)

Dated: April 7, 2021.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2021–07478 Filed 4–12–21; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10757]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 June 14, 2021.

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: CMS–P–0015A, 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' website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

FOR FURTHER INFORMATION CONTACT: William N. 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–10757 CLIA Collection of Information Requirements Related to SARS–CoV–2 Test Results Reporting

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:* Extension of a currently approved information collection; *Title of Information Collection:* CLIA Collection of Information Requirements Related to SARS–CoV–2 Test Results Reporting; *Use:* In order to be in compliance with the new CLIA mandatory SARS–CoV–2 test results reporting requirements, laboratories will need to develop a mechanism to track, collect, and report test results as well as update policies and procedures. In addition, Accreditation Organizations (AOs) and Exempt States (ESs) will need to update laboratory standards to reflect the reporting requirements and update policies and procedures related to reporting laboratories that do not report test results as required.

The CDC has an information collection request (OMB Control Number 0920–1299) in order to collect laboratory data related to the COVID–19 Pandemic Response. The CMS package (ICR) is for laboratory implementation and CMS monitoring of compliance with the CMS–3401–IFC CLIA-certified laboratory reporting requirements.

The information collected by the Centers for Medicare and Medicaid Services (CMS) or its designee, such as a CMS agent or CMS approved laboratory accreditation organization, when conducting inspections will be used to determine a laboratory's compliance with the CLIA SARS–CoV–2 test result reporting requirements. During an on-site survey, the Condition-level laboratory requirement at 42 CFR 493.41and 493.1100(a) are assessed for compliance. The information is used by CMS in determining appropriate Civil Money Penalties (CMPs) when laboratories fail to report as required. *Form Number:* CMS–10757 (OMB control number: 0938–1391); *Frequency:* Daily; *Affected Public:* Private Sector Not-for-profit institutions and State, Local and Tribal Governments; *Number of Respondents:* 77,033; *Total Annual Responses:* 308,114; *Total Annual Hours:* 1,386,873 (For policy questions regarding this collection contact Sarah Bennett at 410–786–3354.)

Dated: April 8, 2021.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2021–07559 Filed 4–12–21; 8:45 am]

BILLING CODE 4120–01–P