open to the public, limited only by room seating available. The public is also welcome to listen to the meeting by 866-836-4010, passcode: 18307719, and, and 100 teleconference lines are available.

DATES: The meeting will be held on November 8, 2017, 9:00 a.m. to 5:00 p.m., EST and November 9, 2017, 9:00 a.m. to 12:00 p.m., EST.

ADDRESSES: Centers for Disease Control and Prevention, Global Communications Center, Building 19, Auditorium B, 1600 Clifton Road NE., Atlanta, Georgia

FOR FURTHER INFORMATION CONTACT: Erin Stone, M.A., HICPAC, Division of Healthcare Quality Promotion, NCEZID, CDC, l600 Clifton Road NE., Mailstop A-07, Atlanta, Georgia 30329, Telephone (404) 639-4045, Email: hicpac@cdc.gov.

SUPPLEMENTARY INFORMATION:

Purpose: The Committee is charged with providing advice and guidance to the Director, Division of Healthcare Quality Promotion (DHQP), the Director, National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), the Director, CDC, the Secretary, Health and Human Services regarding (1) the practice of healthcare infection prevention and control; (2) strategies for surveillance, prevention, and control of infections, antimicrobial resistance, and related events in settings where healthcare is provided; and (3) periodic updating of CDC guidelines and other policy statements regarding prevention of healthcare-associated infections and healthcare-related conditions.

Matters to be Considered: The agenda will include updates on CDC's activities for prevention of healthcare associated infections (HAIs), and DHQP's modeling efforts. It will also include updates from the following HICPAC workgroups: The Guideline for Prevention of Infection in Neonatal Intensive Care Unit (NICU) Patients; the Guideline for Prevention of Infection in Healthcare Personnel; the workgroup on updating the CDC recommendation categorization scheme; the workgroup on developing CDC recommendations for products and practices; and the National Healthcare Safety Network (NHSN) Surveillance Workgroup.

Agenda items are subject to change as priorities dictate.

Time will be available for public comment. The public is welcome to submit written comments in advance of the meeting. Comments should be submitted in writing by email to the contact person listed below. The deadline for receipt is October 25, 2017. All requests must contain the name, address, and organizational affiliation of the speaker, as well as the topic being addressed. Written comments should not exceed one single-spaced typed page in length and delivered in 3 minutes or less. Members of the public who wish to provide public comments should plan to attend the public comment session at the start time listed. Please note that the public comment period may end before the time indicated, following the last call for comments. Written comments received in advance of the meeting will be included in the official record of the meeting. Registration is required to attend in person or on the phone. Interested parties may register at www.cdc.gov/ hicpac.

The Director, Management Analysis and Services Office, 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.

Claudette Grant.

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid **Services**

[Document Identifiers: CMS-10292 and CMS-R-148]

Agency Information Collection Activities: Submission for OMB Review; 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 (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 October 18, 2017. 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 following:

1. Access CMS' Web site address at Web site address at https:// www.cms.gov/Regulations-and-Guidance/Legislation/Paperwork ReductionActof1995/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: 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: State Medicaid HIT Plan, Planning Advance Planning Document, and Implementation Advance Planning Document for Section 4201 of the Recovery Act; Use: To assess the appropriateness of state requests for the administrative Federal financial participation for expenditures under their Medicaid Electronic Health Record Incentive Program related to health information exchange, our staff will review the submitted information and documentation to make an approval determination of the state advance planning document. Form Number: CMS-10292 (OMB control number: 0938-1088); Frequency: Once and occasionally; Affected Public: State, Local, and Tribal Governments; Number of Respondents: 56; Total Annual Responses: 56; Total Annual Hours: 896. (For policy questions regarding this collection contact Marty Rice at 410-786-2417.)
- 2. Type of Information Collection Request: Extension of a currently approved collection; Title of *Information Collection:* Limitations on Provider Related Donations and Health Care Related Taxes; Limitation on Payment to Disproportionate Share Hospitals; Medicaid and Supporting Regulations in 42 CFR 433.68, 433.74 and 447.272; Use: States may request a waiver of the broad based and uniformity tax program requirements. Each state must demonstrate that its tax program(s) do not violate the hold harmless provision. Additionally, state Medicaid agencies must report (quarterly) on health care related taxes collected and the source of provider related donations received by the state or unit of local government. Each state must maintain, in readily reviewable form, supporting documentation that provides a detailed description of each donation and tax program being reported, as well as the source and use of all donations received and collected. Without this information, the amount of Federal financial participation payable to a state cannot be determined; Form Number: CMS-R-148 (OMB control number: 0938–0618); *Frequency:* Quarterly and occasionally; Affected Public: State, Local, or Tribal Governments; Number of Respondents: 50; Total Annual Responses: 40; Total Annual Hours: 3,200. (For policy questions regarding this collection contact Stuart Goldstein at 410-786-0694.)

Dated: September 13, 2017.

William N. Parham, III,

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

[FR Doc. 2017–19787 Filed 9–15–17; 8:45 am]

BILLING CODE 4120-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-437 and CMS-10515]

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 ČMS' 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 November 17, 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 http://www.cms.hhs.gov/Paperwork ReductionActof1995.

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–437 Psychiatric Unit Criteria Work Sheet

CMS-10515 Payment Collections Operations Contingency Plan

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: Reinstatement with Change of a currently approved collection; Title of Information Collection: Psychiatric Unit Criteria Work Sheet; Use: Certain