

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)—RFA-IP-24-045, Network of Community Cohorts for Monitoring Changes in Respiratory Virus Epidemiology (Pandemic Preparedness Cohorts).

Dates: April 25–26, 2024.

Times: 10 a.m.–5 p.m., EDT.

Place: Videoconference.

Agenda: To review and evaluate grant applications.

For Further Information Contact:

Gregory Anderson, MS, M.P.H., Scientific Review Officer, National Center for HIV, Viral Hepatitis, STD, and TB Prevention, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H24-6, Atlanta, Georgia 30329-4027. Telephone: (404) 718-8833; Email: GAnderson@cdc.gov.

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10346]

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 February 20, 2024.

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, please access the CMS PRA website by copying and pasting the following web address into your web browser: <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing>.

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 without change of a currently approved collection; *Title of Information Collection:* Quality Bonus Payment Appeals; *Use:* Section 1853(o) of the Act requires CMS to make QBPs to MA organizations that achieve performance rating scores of at least 4 stars under a five-star rating system. While CMS has applied a Star Rating system to MA organizations for a number of years, prior to the QBP program these Star Ratings were used only to provide additional information for beneficiaries to consider in making their Part C and D plan elections. Beginning in 2012, the Star Ratings CMS assigns for purposes of QBPs directly affected the monthly payment amount MA organizations receive from CMS under their contracts. Additionally, section 1854(b)(1)(C)(v) of the Act, as added by the Affordable Care Act, also requires CMS to change the share of savings that MA organizations must provide to enrollees as the beneficiary rebate specified at § 422.266(a) based on the level of a sponsor's Star Rating for quality performance.

The information collected on the Request for Reconsideration form from MA organizations is considered by the reconsideration official and potentially the hearing officer to review CMS's determination of the organization's eligibility for a QBP. The form asks MA organizations to select the Star Ratings measure(s) they believe was miscalculated or used incorrect data and describe what they believe is the issue. Under § 422.260(c)(3)(ii) these are the only bases for appeals. In conducting the reconsideration, the reconsideration official will review the QBP determination, the evidence and findings upon which it was based, and any other written evidence submitted by the organization with their Request for Reconsideration or by CMS before the reconsideration determination is made. *Form Number:* CMS-10346 (OMB Control Number 0938-1129); *Frequency:* Yearly; *Affected Public:* Private Sector; *Number of Respondents:* 20; *Total Annual Responses:* 20; *Total Annual Hours:* 160. (For policy questions regarding this collection contact, Sarah Gaillot at 410-786-4637.)

Dated: January 12, 2024.

William N. Parham, III

Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2024-00897 Filed 1-17-24; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Runaway and Homeless Youth—Homeless Management Information System (RHY-HMIS; Office of Management and Budget #0970-0573)

AGENCY: Family and Youth Services Bureau, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Family and Youth Services Bureau’s Runaway and Homeless Youth (RHY) Program is requesting a 3-year extension of the Runaway and Homeless Youth—Homeless Management Information System (RHY-HMIS) data collection efforts (OMB #0970-0573, expiration 07/31/2024). There are no changes requested to the data elements.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: You can obtain copies of the proposed collection of information and submit comments by emailing infocollection@acf.hhs.gov. Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The RHY Program has a requirement to collect information from all youth who receive shelter and supportive services with RHY funding. In April 2015, the Administration on Children, Youth and Families, through a formal Memorandum of Understanding, integrated the RHY data collection with the U.S. Department of Housing and Urban Development’s (HUD) HMIS and HUD’s data standards along with other federal partners. HUD has OMB approval for HUD’s data standards and ACF has approval under a separate OMB number for the RHY data elements. The data collection effort includes universal data elements that are collected by all federal partners and RHY program specific elements, which are tailored to the RHY Program using HUD’s HMIS.

Respondents: Youth who receive emergency and longer-term shelter and supportive services under RHY funding.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
RHY-HMIS: Basic Center Program (Intake)	123,000	1	0.38	46,740	15,580
RHY-HMIS: Basic Center Program (Exit)	123,000	1	0.33	40,590	13,530
RHY-HMIS: Transitional Living Program (including Maternity Group Home program and TLP Demonstration Programs; Intake)	24,000	1	0.38	9,120	3,040
RHY-HMIS: Transitional Living Program (including Maternity Group Home program and TLP Demonstration Programs; Exit)	24,000	1	0.33	7,920	2,640
RHY-HMIS: Street Outreach Program (Contact)	108,000	1	0.5	54,000	18,000
RHY-HMIS: Street Outreach Program (Engagement)	30,000	1	0.28	8,400	2,800
RHY Funded Grantees (data entry)	308,225	2	0.36	221,922	73,974
RHY Funded Grantees (data submission)—FY24	675	2	0.16	216	72
RHY Funded Grantees (data submission)—FY25 & FY26	675	8	0.16	864	288
Estimated Total Annual Burden Hours	129,924

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency’s estimate of the burden of the proposed collection of information; (c) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: Reconnecting Homeless Youth Act of 2008 (Public Law 110-378) through Fiscal Year (FY) 2013 and reauthorized by the Juvenile Justice Reform Act through FY 2019.

Mary B. Jones,

ACF/OPRE Certifying Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2018-N-2727]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Institutional Review Board Waiver or Alteration of Informed Consent for Minimal Risk Clinical Investigations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing that a collection of information entitled “Institutional