

January 19, 2023).

- * 8. Nho, SW, S-J. Kim, O. Kweon, et al. "Microbiological Survey of Commercial Tattoo and Permanent Makeup Inks Available in the United States." *Journal of Applied Microbiology*, 124: 1294–1302 (2018).
- * 9. Food and Drug Administration, "FDA Advises Consumers, Tattoo Artists, and Retailers to Avoid Using or Selling Certain Tattoo Inks Contaminated with Microorganisms"; available at <https://www.fda.gov/cosmetics/cosmetics-recalls-alerts/fda-advises-consumers-tattoo-artists-and-retailers-avoid-using-or-selling-certain-tattoo-inks> (accessed January 19, 2023).

Dated: June 6, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2023–12380 Filed 6–12–23; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection

Activities: Proposed Collection: Public Comment Request: Information Collection Request Title: Evaluation of the Maternal and Child Health Bureau's Autism CARES Act Initiative, OMB No. 0915–0335–Revision

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, HRSA 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. OMB may act on HRSA's ICR only after the 30-day comment period for this notice has closed.

DATES: Comments on this ICR should be received no later than July 13, 2023.

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 Review—Open for Public Comments" or by using the search function.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, contact Samantha Miller, the HRSA Information Collection Clearance Officer, at paperwork@hrsa.gov or call 301–443–3938.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the information request collection title for reference.

Information Collection Request Title: Evaluation of the Maternal and Child Health Bureau's Autism CARES Act Initiative, OMB No. 0915–0335–Revision.

Abstract: HRSA's Maternal and Child Health Bureau (MCHB) provides funds to support several programs related to autism, as authorized by 42 U.S.C. 280i–1 (title III, section 399BB of the Public Health Service Act), as amended by the Autism Collaboration, Accountability, Research, Education, and Support (CARES) Act of 2019 (Pub. L. 116–60). The Autism CARES Act of 2019 emphasizes improving health outcomes and the well-being of individuals with Autism Spectrum Disorder and Developmental Disabilities across the lifespan.

MCHB's programs related to autism fall within three distinct but complementary areas—research, state systems, and training. The awards advance research on early screening and interventions for autism and developmental disabilities; improve the capacity of state public health agencies to build and maintain coordinated systems of services for individuals with autism and developmental disabilities; and train the health care workforce to screen, refer, and provide services for children and youth with autism and developmental disabilities. MCHB currently funds 12 programs and 95 awardees. HRSA seeks to implement annual comprehensive evaluations of MCHB's Autism CARES Initiative investments.

This ICR is a revision to an existing package; this study is the fifth evaluation of HRSA's autism activities and employs similar data collection methodologies as the prior studies. Grantee interviews remain the primary form of data collection. Minor proposed revisions to the data collection process include modifications to the interview questions and grantee survey based on current legislation and HRSA's Notices of Funding Opportunity for programs authorized under the Autism CARES Act. In addition, the previous data collection compiled survey responses from all grantees, whereas this revised

data collection will only seek survey responses from the Research and State Systems grantees. The previous data collection also included a quantitative data collection form for the Research grantees that the current data collection will not collect. These changes result in fewer burden hours estimated across all primary data collection activities.

A 60-day notice published in the **Federal Register** on March 21, 2023, vol. 88, No. 54; pp. 16995–16996. There were no public comments.

Need and Proposed Use of the Information: The purpose of this data collection is to implement a comprehensive evaluation that describes the activities, accomplishments, outcomes, barriers, and challenges of the grant programs in implementing the provisions of the Autism CARES Act. The data will be used to (1) conduct performance monitoring of the programs; (2) provide credible and rigorous evidence of program effectiveness; (3) meet program needs for accountability, decision-making, and quality assurance; and (4) strengthen the evidence base for best practices.

Likely Respondents: The survey respondents will include Principal Investigators/Project Directors from the research programs and networks (Autism Intervention Research Network on Physical Health, Autism Intervention Research Network on Behavioral Health, MCHB Secondary Data Analysis Research Program, Autism Field-Initiated Innovative Research Studies Program, Autism Single Investigator Innovation Program, the Developmental-Behavioral Pediatrics Research Network, and the Healthy Weight Research Network for Children with Autism and Other Developmental Disabilities); and state systems programs (State Innovations) and coordinating center (State Public Health Coordinating Center for Autism). The respondents for the interviews will include Principal Investigators/Project Directors from the research and state systems programs above, and the training programs (Leadership Education in Neurodevelopmental and Related Disabilities program, the Developmental Behavioral Pediatrics program, and the National Interdisciplinary Training Resource Center).

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

Grant program/instrument	Number of respondents	Average number of responses per respondent	Total responses	Average burden per response (in hours)	Total hour burden
Research: Survey for individual grantees	12	1	12	0.5	6.0
Research: Survey for research networks	4	1	4	0.5	2.0
Research: Interview guide for individual grantees	12	1	12	1.5	18.0
Research: Interview guide for research networks	4	1	4	1.5	6.0
State Systems: Survey for state innovation grants	5	1	5	0.5	2.5
State Systems: Interview guide for the state innovation grants	5	1	5	1.5	7.5
State Systems: Interview guide for the state coordinating center	1	1	1	1.5	1.5
Training: Interview guide for the individual training grantees	72	1	72	1.5	108.0
Training: Interview Guide for the Resource Center	1	1	2	1.5	3.0
Total	116	116	154.5

HRSA specifically requests comments on (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.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2023-12608 Filed 6-12-23; 8:45 am]

BILLING CODE 4165-15-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Committee on Vital and Health Statistics; Meeting and Request for Information

AGENCY: Centers for Disease Control and Prevention, HHS.

ACTION: Notice of meeting. Notice of request for information (RFI).

SUMMARY: Pursuant to the Federal Advisory Committee Act, the Department of Health and Human Services (HHS) announces the following advisory committee meeting and related Request for Information (RFI). The meeting is open to the public. The public is invited and welcome to obtain the link to attend this meeting by following the instructions posted on the Committee website: <https://ncvhs.hhs.gov/meetings-meeting/>.

DATES: The meeting will be held Thursday, August 3, 2023: 10:00 a.m.–5:30 p.m. EDT.

To submit comments in response to the RFI, please send by close of business June 30, 2023, to NCVHSmal@cdc.gov, and include on the subject line:

Response from [your organization or name] regarding ICD-11 RFI.

ADDRESSES: Virtual open meeting.

FOR FURTHER INFORMATION CONTACT:

Substantive program information may be obtained from Rebecca Hines, MHS, Executive Secretary, NCVHS, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Hyattsville, Maryland 20782, via electronic mail to vgh4@cdc.gov; or by telephone (301) 458-4715. Summaries of meetings and a roster of Committee members are available on the home page of the NCVHS website <https://ncvhs.hhs.gov/>, where further information including an agenda and instructions to access the broadcast of the meeting will be posted.

Should you require reasonable accommodation, please telephone the CDC Office of Equal Employment Opportunity at (770) 488-3210 as soon as possible.

SUPPLEMENTARY INFORMATION:

Name: National Committee on Vital and Health Statistics (NCVHS).

Meeting of the Workgroup on Timely and Strategic Action to Inform ICD-11 Policy.

The National Committee on Vital and Health Statistics was established by Congress to serve as the statutory [42 U.S.C. 242k(k)] advisory body to the Secretary of Health and Human Services

for health data, statistics, privacy and national health information policy and the Health Insurance Portability and Accountability Act (HIPAA).¹ In that capacity, the Committee provides advice and assistance to the Department and serves as a forum for interaction with interested private sector groups on health data issues. It fulfills important review and advisory functions regarding health data and administrative standards of national and international scope, conducts studies of prevailing current topics, and makes recommendations for improvement of the Nation's health statistics and information systems.

Purpose: The purpose of the International Classification of Diseases (ICD-11) expert roundtable meeting is to gather information and identify gaps in currently available information and research essential for analysis and policy decisions on the U.S. approach to support adoption and implementation of ICD-11 for morbidity. A supplemental goal is to enable coordination of public and private entities that may affect ICD-11 integration into U.S. health information environments by obtaining broad stakeholder input on studies or assessments HHS should undertake to inform the transition and on what timeline. Together with comments received in response to the RFI, the input received at the roundtable will inform the Workgroup's findings to be provided to the full Committee in

¹Public Law 104-191, 110 Stat. 1936 (Aug 21, 1996), available at: <https://www.congress.gov/104/plaws/publ191/PLAW-104publ191.pdf>.