

quality and relevance of the information it produces. NCHS will conduct voluntary customer surveys to assess strengths in agency products and services and to evaluate how well it addresses the emerging needs of its data users. Results of these surveys will be used in future planning initiatives.

The data will be collected using a combination of methodologies appropriate to each survey. These may include: evaluation forms, mail surveys, focus groups, automated and electronic technology (e.g., email, web-based surveys), and telephone surveys. Systematic surveys of several groups will be folded into the program. Among these are Federal customers and policy

makers, state and local officials who rely on NCHS data, the broader educational, research, and public health community, and other data users. Respondents may include data users who register for and/or attend NCHS sponsored conferences; persons who access the NCHS website and the detailed data available through it; consultants; and others. Respondent data items may include (in broad categories) information regarding respondent's gender, age, occupation, affiliation, location, etc., to be used to characterize responses only. Other questions will attempt to obtain information that will characterize the

respondents' familiarity with and use of NCHS data, their assessment of data content and usefulness, general satisfaction with available services and products, and suggestions for improvement of surveys, services and products.

In order to capture feedback opportunities, this extension request allows for both respondents and time per response for a total estimated annual burden total of 4,000 hours over the three-year requested approval period. There is no cost to respondents other than their time to participate. The resulting information will be for NCHS internal use.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (hours)
Questionnaire for conference registrants/ attendees.	Public/private researchers, Consultants, and others.	6,000	1	15/60
Focus groups	Public/private researchers, Consultants, and others.	500	1	1
Web-based	Public/private researchers, Consultants, and others.	6,000	1	15/60
Other customer surveys	Public/private researchers, Consultants, and others.	2,000	1	15/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2020-13490 Filed 6-22-20; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-20-0006]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled Statement in Support of Application for Waiver of Inadmissibility Under Immigration and Nationality Act to the Office of Management and Budget (OMB) for review and approval. CDC previously published a "Proposed Data Collection Submitted for Public Comment and Recommendations" notice on January 22, 2020 to obtain comments from the public and affected agencies. CDC received two comments related to the

previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

- (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. 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. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Statement in Support of Application for Waiver of Inadmissibility Under Immigration and Nationality Act (OMB Control No.0920-0006, Exp. 06/30/2020)—Revision—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 212(a)(1) of the Immigration and Nationality Act states that aliens with specific health related conditions are ineligible for admission into the United States. The Attorney General may waive application of this inadmissibility on health-related grounds if an application for waiver is

filed and approved by the consular office considering the application for visa. CDC uses this application primarily to collect information to establish and maintain records of waiver applicants in order to notify the U.S. Citizenship and Immigration Services when terms, conditions and controls imposed by waiver are not met.

CDC is removing information collection form 4.422–1a in this revision, because CDC no longer uses this form to receive information about the evaluation report of an applicant who received a waiver. Since this form is no longer being used, the estimated annual total burden hours decreased by 67 hours, from 100 to 33 annual burden hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Physician	CDC 4.422–1	200	1	10/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2020–13489 Filed 6–22–20; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; National Child Abuse and Neglect Data System (OMB #0970–0424)

AGENCY: Children’s Bureau; Administration on Children, Youth and Families; Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) in the U.S. Department of Health and Human Services (HHS) is requesting a 3-year extension of the National Child Abuse and Neglect Data System (NCANDS) collection (OMB #0970–0424, expiration 02/28/2021). There are no changes requested to this data collection.

DATES: *Comments due within 30 days of publication.* OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

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.

SUPPLEMENTARY INFORMATION:

Description: The Child Abuse Prevention and Treatment Act (CAPTA) was amended in 1988 to direct the Secretary of HHS to establish a national data collection and analysis program, which would make available state child abuse and neglect reporting information. HHS responded by establishing NCANDS as a voluntary national reporting system.

In 1996, CAPTA was amended to require all states that receive funds from the Basic State Grant program to work with the Secretary of HHS to provide specific data elements, to the maximum extent practicable, about children who had been maltreated. Most of the required data elements were added to the NCANDS data collection. Subsequent CAPTA reauthorizations and amendments added required data elements. The current list of CAPTA-required data elements includes:

(1) The number of children who were reported to the state during the year as victims of child abuse or neglect.

(2) Of the number of children described in paragraph (1), the number with respect to whom such reports were—

- (a) Substantiated;
- (b) Unsubstantiated; or
- (c) Determined to be false.

(3) Of the number of children described in paragraph (2)—

(a) the number that did not receive services during the year under the state program funded under this section or an equivalent state program;

(b) the number that received services during the year under the state program funded under this section or an equivalent state program; and

(c) the number that were removed from their families during the year by disposition of the case.

(4) The number of families that received preventive services, including use of differential response, from the state during the year.

(5) The number of deaths in the state during the year resulting from child abuse or neglect.

(6) Of the number of children described in paragraph (5), the number of such children who were in foster care.

(7) (a) The number of child protective service personnel responsible for the— (i.) intake of reports filed in the previous year;

(ii.) screening of such reports; (iii.) assessment of such reports; and (iv.) investigation of such reports.

(b) The average caseload for the workers described in subparagraph (A).

(8) The agency response time with respect to each report pertaining to the initial investigation of child abuse or neglect.

(9) The response time with respect to the provision of services to families and children where an allegation of child abuse or neglect has been made.

(10) For child protective service personnel responsible for intake, screening, assessment, and investigation of child abuse and neglect reports in the state—

(a) information on the education, qualifications, and training requirements established by the state for child protective service professionals, including for entry and advancement in the profession, including advancement to supervisory positions;

(b) data of the education, qualifications, and training of such personnel;

(c) demographic information of the child protective service personnel; and