requests for approval are evaluated. The applications are submitted at will and the most reasonable prediction of respondents is the number from the most recent year, 63 in 2013. The decrease is likely due to random fluctuations and changes in business conditions. No survey was conducted to more thoroughly analyze the reasons for the change in number of respondents. Although there is no cost to respondents to submit other than their time to participate, respondents requesting respirator approval are required to submit fees for necessary testing as specified in 42 CFR Parts 84.20–22, 84.66, 84.258 and 84.1102. In calendar year 2013 $449,610.135 was accepted.

Applicants are required to provide test data that shows that the manufacturer is capable of ensuring that the respirator is capable of meeting the specified requirements in 42 CFR Part 84. The requirement for submitted test data is likely to be satisfied by standard testing performed by the manufacturer, and is not required to follow the relevant NIOSH Standard Test Procedures. As additional testing is not required, providing proof that an adequate test has been performed is limited to providing existing paperwork.

42 CFR Part 84 approvals offer corroboration that approved respirators are produced to certain quality standards. Although 42 CFR Part 84 Subpart E prescribes certain quality standards, it is not expected that requiring approved quality standards will impose an additional cost burden over similarly effective quality standards that are not approved under 42 CFR Part 84.

Manufacturers with current approvals are subject to site audits by the Institute or its agents. There is no fee associated with audits. Audits may occur periodically or as a result of a reported issue. Sixty site audits were scheduled for the 2013 calendar year.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Response type	Expected annual number of respondents	Average annual responses per respondent	Average burden hours per response	Total burden hours
Business or other for-profit	Standard Application for the Approval of Respirators Version 7 and Version 8.	63	7	229	100,989
Business or other for-profit	Audit	60	1	24	1,440
Total					102,429

Leroy Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–13–0729]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call (404) 639–7570 or send an email to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Customer Surveys Generic Clearance for the National Center for Health Statistics (0920–0729, Expiration 04/30/2014)—Revision—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C. 242k), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on ''the extent and nature of illness and disability of the population of the United States.'' This is a revision request for a generic approval from OMB to conduct customer surveys over the next three years.

As part of a comprehensive program, the National Center for Health Statistics (NCHS) plans to continue to assess its customers' satisfaction with the content, 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 Web site 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.

The resulting information will be for NCHS internal use. There is no cost to respondents other than their time to participate in the survey. The total

burden for three years of clearance is 2,040 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name/survey type	Number of respondents	Responses per respondent	Average burden per response (in hours)
Public/private researchers, Consultants, and others.	Questionnaire for conference registrants/ attendees.	4,500	1	10/60
Public/private researchers, Consultants, and others.	Focus groups	240	1	1
Public/private researchers, Consultants, and others.	Web-based	4,500	1	10/60
Public/private researchers, Consultants, and others.	Other customer surveys	1,200	1	15/60

LeRoy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Center for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: State Plan for Grants to States for Refugee Resettlement.

OMB No.: 0970–0351.

Description: A State Plan is required by 8 U.S.C. 1522 of the Immigration and Nationality Act (the Act) [Title IV, Sec. 412 of the Act] for each State agency requesting Federal funding for refugee resettlement under 8 U.S.C. 524 [Title IV, Sec. 414 of the Act], including Refugee Cash and Medical Assistance, Unaccompanied Minor Refugee Program, Refugee Social Services, Cuban/Haitian Entrant Program and Targeted Assistance program funding. The State Plan is a comprehensive narrative description of the nature and scope of a States programs and provides assurances that the programs will be administered in conformity with the specific requirements stipulated in 45 CFR 400.4–400.9. The State Plan must include all applicable State procedures, designations, and certifications for each requirement as well as supporting documentation. The plan assures ORR that the State is capable of administering refugee assistance and coordinating employment and other social services for eligible caseloads in conformity with specific requirements. Implementation of the Affordable Care Act has significant impacts on States' administration of Refugee Medical Assistance and requires information to ensure accountability and compliance with regulations. Also, Revised Medical Screening Guidelines for Newly Arriving Refugees policy (State Letter #12–09) requires assurances that medical screening is conducted in compliance with regulations and policies. The increasing complexity of the Unaccompanied Refugee Minor program, impacted by changes in federal child welfare legislation as well as state child welfare statutes, regulations and IV–B and IV–E plans, necessitates information and assurances for review of State Plans for URM programs against requirements and mandatory standards under 45 CFR Part 400, subpart H and associated State Letters and ORR guidance. Information and assurances address administrative structure and state oversight, legal responsibility, eligibility, services and case review/ planning, and interstate movement.

States must use a pre-print format for required components of State Plans for ORR-funded refugee resettlement services and benefits prepared by the Office of Refugee Resettlement (ORR) of the Administration for Children and Families (ACF).

States must submit by August 15 each year new or amended State Plan for the next Federal fiscal year. For previously approved plan, States must certify no later than October 31 each year that the approved State plan is current and continues in effect.

Respondents: State Agencies, Replacement Designees under 45 CFR 400.301(c), and Wilson-Fish Grantees (State 2 Agencies) administering or supervising the administration of programs under Title IV of the Act.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Title IV State Plan	50	1	15	750

Estimated Total Annual Burden Hours: 750.

In compliance with the requirements of Section 506(c) (2) (A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary