Creighton, both of San Francisco, California; and Andrew T. Creighton and Timothy Creighton, both of Boulder, Colorado; to join the Creighton Family Control Group, a group acting in concert, to retain voting shares of High Plains Banking Group, Inc., and thereby indirectly retain voting shares of High Plains Bank, both of Flager, Colorado. Joseph P. Creighton was previously permitted by the Federal Reserve System to join the Creighton Family Group.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Associate Secretary of the Board.

[FR Doc. 2026–23904 Filed 11–23–26; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–27–0138; Docket No. CDC–2026–1816]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other federal agencies the opportunity to comment on a continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on an information collection project titled Pulmonary Function Testing Course Approval Program. The program consists of an application submitted by potential sponsors (universities, hospitals, and private consulting firms) who seek NIOSH approval to conduct courses, and if approved, notification to NIOSH of any course or faculty changes during the approval period, which is limited to five years.

DATES: CDC must receive written comments on or before January 25, 2027.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2026–1816 by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *www.regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*www.regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: *omb@cdc.gov.*

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. In addition, the PRA also requires federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. 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;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected;

4. 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 submissions of responses; and

5. Assess information collection costs.

Proposed Project

Pulmonary Function Testing Course Approval Program. (OMB Control No. 0920–0138, Exp. 4/30/2026)—Extension—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

NIOSH has the responsibility under the Occupational Safety and Health Administration's Cotton Dust Standard, 29 CFR 1920.1043, for approving courses to train technicians to perform pulmonary function testing in the cotton industry. Successful completion of a NIOSH-approved course is mandatory under this Standard. In addition, regulations at 42 CFR 37.95(a) specify that persons administering spirometry tests for the national Coal Workers' Health Surveillance Program must successfully complete a NIOSH-approved spirometry training course and maintain a valid certificate by periodically completing NIOSH-approved spirometry refresher training courses. Also, 29 CFR 1910.1053(i)(2)(iv), 29 CFR 1910.1053(i)(3), 29 CFR 1926.1153(h)(2)(iv) and 29 CFR 1926.1153(h)(3) specify that pulmonary function tests for initial and periodic examinations in general industry and construction performed under the Respirable Crystalline Silica Standard should be administered by a spirometry technician with a current certificate from a NIOSH-approved spirometry course.

To carry out its responsibility, NIOSH maintains a Pulmonary Function Testing Course Approval Program. The program consists of an application submitted by potential sponsors (universities, hospitals, and private consulting firms) who seek NIOSH approval to conduct courses, and if approved, notification to NIOSH of any course or faculty changes during the approval period, which is limited to five years. The application form and added materials, including an agenda, curriculum vitae, and course materials are reviewed by NIOSH to determine if the applicant has developed a program which adheres to the criteria required in the Standard. Following approval, any subsequent changes to the course are submitted by course sponsors via letter or email and reviewed by NIOSH staff

to assure that the changes in faculty or course content continue to meet course requirements. Course sponsors also voluntarily submit an annual report to inform NIOSH of their course activity level and any faculty changes. Sponsors who elect to have their approval renewed for an additional five-year period submit a renewal application and supporting documentation for review by NIOSH staff to ensure the course curriculum meets all current standard requirements. Approved courses that elect to offer NIOSH-Approved Spirometry Refresher Courses must submit a separate application and supporting documents for review by NIOSH staff. Institutions and organizations throughout the country voluntarily submit applications and materials to become approved course sponsors and administer training. Submissions are required for NIOSH to evaluate a course and determine whether it meets the criteria in the Standard and whether technicians will be adequately trained as mandated under the Standard.

CDC requests OMB approval for a total of 144 estimated annual burden hours. There is no cost to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Potential Sponsors	Initial Application	3	1	8	24
Approved Sponsors	Annual Report	33	1	30/60	17
Approved Sponsors	Report for Course Changes	24	1	30/60	12
Approved Sponsors	Renewal Application	10	1	6	60
Approved Sponsors	Refresher Course Application	3	1	8	24
Total					144

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2026–23909 Filed 11–23–26; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Office of Management and Budget #: 0970–0624]

Proposed Information Collection Activity; Voluntary Agencies Matching Grant Program Data Reporting

AGENCY: Office of Refugee Resettlement, Administration for Children and Families, U.S. Department of Health and Human Services.

SUMMARY: The Office of Refugee Resettlement (ORR) and the Administration for Children and Families (ACF) are requesting a 3-year extension of the Voluntary Agencies Matching Grant Data Reporting (Office of Management and Budget #: 0970–0624, expiration February 28, 2027). There are no changes requested.

DATES: Comments due January 25, 2027.

ADDRESSES: In compliance with the requirements of the Paperwork Reduction Act, ACF is soliciting public comment on the specific aspects of the information collection described above. 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: This Matching Grant Program data collection is designed to satisfy the statutory requirements of the Immigration and Nationality Act (INA). Section 412(a)(3) of INA (8 U.S.C. 1522(a)(3)) requires that the Director of ORR make a periodic assessment of the needs of refugees for assistance and services and the resources available to meet those needs. In addition, section 412(a)(7) of INA (8 U.S.C. 1522(a)(7)) requires ORR to develop a system to monitor the assistance provided under the Refugee Act that includes evaluation of the effectiveness of the program and data collection on the services provided and the results achieved. ORR proposes collecting both case-level and client-level data elements at multiple points in time, which will allow the ORR Director to better understand client demographics, services utilized, and the outcomes achieved by clients enrolled in the Matching Grant Program. Data elements include biographical information, progress made toward achieving self-sufficiency, and employment status. The data collected will inform evidence-based policy making and program design.

Respondents: National resettlement agencies participating in the Matching Grant Program.

ANNUAL BURDEN ESTIMATES

Information collection	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
Voluntary Agencies Matching Grant Program Data Reporting—Data System	7	1	60	420	140
Voluntary Agencies Matching Grant Program Data Reporting—Monthly Uploads	7	36	16	4,032	1,344
Estimated Total Annual Burden Hours					1,484