

collects accounting data from the States on payables and receivables. *Form Number:* CMS–R–199 (OMB control number: 0938–0697); *Frequency:* Yearly; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 56; *Total Annual Responses:* 56; *Total Annual Hours:* 1,792. (For policy questions regarding this collection contact: Kimaada Battle at (410) 786–8042.)

William N. Parham, III,

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

[FR Doc. 2025–17203 Filed 9–5–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10398 #56]

Medicaid and Children’s Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance related to the “generic” clearance process. Generally, this is an expedited process by which agencies may obtain OMB’s approval of collection of information requests that are “usually voluntary, low-burden, and uncontroversial collections,” do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an overarching plan that defines the scope of the individual collections that would fall under its umbrella. This **Federal Register** notice seeks public comment on one or more of our collection of information requests that we believe are generic and fall within the scope of the umbrella. Interested persons are invited to submit comments regarding our burden estimates 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 must be received by September 22, 2025.

ADDRESSES: When commenting, please reference the applicable form number (CMS–10398 #56) and the OMB control number (0938–1148). To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for “Comment or Submission” or “More Search Options” to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: CMS–10398 #56/OMB control number: 0938–1148, Room C4–26–05, 7500 Security Boulevard, Baltimore, MD 21244–1850.

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/medicare/regulations-guidance/legislation/paperwork-reduction-act-1995/pralisting>.

FOR FURTHER INFORMATION CONTACT: William N. Parham at 410–786–4669.

SUPPLEMENTARY INFORMATION: Following is a summary of the use and burden associated with the subject information collection(s). More detailed information can be found in the collection’s supporting statement and associated materials (see **ADDRESSES**).

Generic Information Collection

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Section 1115 Demonstration: Budget Neutrality Monitoring Workbook; *Use:* To help CMS track if a state’s demonstration is budget neutral, the state must submit the Budget Neutrality Monitoring Workbook with their expenditure data (at least annually) that compares their actual and projected spending to the budget neutrality limits agreed to at approval of the demonstration. If a state is found to be close or over the limit of budget neutrality, a corrective action plan may be started to help the state achieve budget neutrality during the

demonstration period. *Form Number:* CMS–10398 #56 (OMB control number: 0938–1148); *Frequency:* Yearly and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 47; *Total Annual Responses:* 466; *Total Annual Hours:* 3,003. (For policy questions regarding this collection contact: Raven Smith at 410–786–3731.)

William N. Parham, III,

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

[FR Doc. 2025–17121 Filed 9–5–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Sickle Cell Disease Newborn Screening Follow-Up Program

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

ACTION: Notice of Supplemental awards.

SUMMARY: HRSA will provide \$65,500 in supplemental award funds to each of the 25 Sickle Cell Disease (SCD) Newborn Screening Follow-Up Program (NBS FP) grantees under HRSA–21–036 in fiscal year (FY) 2025, totaling \$1,637,500. The FY 2025 supplemental funds will support grant activities that are in scope of program goals and objectives. The supplemental funding can support activities related to: increasing access to medical and support services, particularly during the pediatric to adult transition period; improving data systems; increasing workforce training and knowledge; expanding quality improvement projects; or conducting outreach to help identify individuals living with SCD who have been lost to follow-up. These activities will strengthen the linkage of individuals with SCD to appropriate and ongoing care to improve health outcomes.

FOR FURTHER INFORMATION CONTACT: Hakim Fobia, Public Health Analyst and Project Officer, Genetic Services Branch, Division of Services for Children with Special Health Needs, HRSA, at scdprograms@hrsa.gov or 301–945–9842.

SUPPLEMENTARY INFORMATION:

Intended Recipients of the Award: 25 SCD NBS FP recipients listed in Table 1.

Amount of Non-Competitive Awards: 25 awards at \$65,500 (total: \$1,637,500).

Project Period: September 1, 2021, to August 31, 2026.
 Assistance Listing Number: 93.110.

Award Instrument: Non-competitive supplement.

Authority: 42 U.S.C. 701(a)(2) (§ 501(a)(2) of the Social Security Act).

TABLE 1—RECIPIENT(S) AND AWARD AMOUNT(S)

Grant No.	Award recipient name	City, state	Award amount
D1JMC42466	Sickle Cell Disease Association of Illinois	Chicago, IL	\$65,500
D1JMC42474	Hemostasis and Thrombosis Center of Nevada	Las Vegas, NV	65,500
D1JMC42475	Ohio Sickle Cell & Health Association, Inc	Columbus, OH	65,500
D1JMC42467	Martin Center, Inc	Indianapolis, IN	65,500
D1JMC42468	Baton Rouge Sickle Cell Anemia Foundation, Inc	Baton Rouge, LA	65,500
D1JMC42463	Cayenne Wellness Center and Children's Foundation, Inc	Burbank, CA	65,500
D1JMC42472	Children's Sickle Cell Foundation, Inc	Pittsburgh, PA	65,500
D1JMC42473	James R. Clark Memorial Sickle Cell Foundation	Columbia, SC	65,500
D1JMC42471	Piedmont Health Services and Sickle Cell Agency	Greensboro, NC	65,500
D1JMC42476	Sickle Cell Anemia Foundation of Oregon, Inc	Portland, OR	65,500
D1JMC42470	Sickle Cell Association	Florissant, MO	65,500
D1JMC42477	The Sickle Cell Association of New Jersey, Inc	Newark, NJ	65,500
D1JMC42469	Sickle Cell Disease Association of America Michigan Chapter, Inc	Detroit, MI	65,500
D1JMC42478	Sickle Cell Disease Association of America Philadelphia-Delaware Valley Chapter.	Philadelphia, PA	65,500
D1JMC42462	Sickle Cell Foundation of Arizona, Inc	Vail, AZ	65,500
D1JMC42465	Sickle Cell Foundation of Georgia, Inc	Atlanta, GA	65,500
D1JMC42479	Sickle Cell Association of Texas Marc Thomas Foundation	Austin, TX	65,500
D1JMC42480	Sickle Cell/Thalassemia Patients Networks, Inc	Brooklyn, NY	65,500
D1JMC42481	Supporters of Families with Sickle Cell Disease, Inc	Tulsa, OK	65,500
D1JMC42464	University of Miami	Miami, FL	65,500
D1JMC46837	Sickle Cell Disease Foundation	Ontario, CA	65,500
D1JMC46836	Sickle Cell Disease Association of America Mobile Chapter	Mobile, AL	65,500
D1JMC46834	Crescent Foundation, Inc.	Philadelphia, PA	65,500
D1JMC46835	Metropolitan Seattle Sickle Cell Task Force	Seattle, WA	65,500
D1JMC46838	Sickle Cell Foundation of Tennessee	Memphis, TN	65,500

Justification: In year 4 of 5 of this program, funding is being made available for FY 2025 to 25 current funding recipients under HRSA–21–036 to facilitate access to quality SCD care by conducting outreach and working with individuals living with SCD and families from the time a newborn screen identifies a child with possible SCD through diagnosis, treatment, and follow-up across the lifespan. Such funding helps bridge access to services by supporting community-based organizations in reaching individuals in the communities where they reside and providing a critical link to health care and other social services to improve the lives of individuals living with SCD. SCD NBS FP grantees will use the supplemental funding to improve appropriate health care and social services utilization by targeting direct and indirect barriers to accessing health care such as transition support, social service linkage, data system development, workforce educational support and training, expanding quality improvement projects, and other services. The above activities are within

the scope of the NBS FP program as described in HRSA–21–036.

Thomas J. Engels,
Administrator.

[FR Doc. 2025–17215 Filed 9–5–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Vaccine Injury Compensation Program; List of Petitions Received

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

ACTION: Notice.

SUMMARY: HRSA is publishing this notice of petitions received under the National Vaccine Injury Compensation Program (the Program), as required by the Public Health Service (PHS) Act, as amended. While the Secretary of HHS is named as the respondent in all proceedings brought by the filing of petitions for compensation under the Program, the United States Court of Federal Claims is charged by statute

with responsibility for considering and acting upon the petitions.

FOR FURTHER INFORMATION CONTACT: For information about requirements for filing petitions, and the Program in general, contact Lisa L. Reyes, Clerk of Court, United States Court of Federal Claims, 717 Madison Place NW, Washington, DC 20005, (202) 357–6400. For information on HRSA’s role in the Program, contact the Director, Division of Injury Compensation Programs, 5600 Fishers Lane, Room 14W–18, Rockville, Maryland 20857; 1–800–338–2382, or visit our website at: <https://www.hrsa.gov/vaccine-compensation>.

SUPPLEMENTARY INFORMATION: The Program provides a system of no-fault compensation for certain individuals who have been injured by specified childhood vaccines. Subtitle 2 of Title XXI of the PHS Act, 42 U.S.C. 300aa–10 *et seq.*, provides that those seeking compensation are to file a petition with the United States Court of Federal Claims and to serve a copy of the petition to the Secretary of HHS, who is named as the respondent in each proceeding. The Secretary has delegated this responsibility under the Program to HRSA. The Court is directed by statute to appoint special masters who take evidence, conduct hearings as appropriate, and make initial decisions