55. Michael Frevola, New York, New York, Court of Federal Claims No: 21–1871V
56. Joyce Scratchard, Burleson, Texas, Court of Federal Claims No: 21–1875V
57. Michael Maxwell, Oviedo, Florida, Court of Federal Claims No: 21–1877V
58. Briana Watson, Kansas City, Missouri, Court of Federal Claims No: 21–1879V
59. Donna Spaid, Winchester, Virginia, Court of Federal Claims No: 21–1880V
60. Amanda Deluca, St. Louis, Missouri, Court of Federal Claims No: 21–1881V
61. Melanie Stanek, Wellesley Hills, Massachusetts, Court of Federal Claims No: 21–1882V
62. Bi Ying Gao, Houston, Texas, Court of Federal Claims No: 21–1884V
63. Terry Yormark, Chicago, Illinois, Court of Federal Claims No: 21–1886V
64. Richard Trudell, Boise, Idaho, Court of Federal Claims No: 21–1887V
65. Angela Mosley, Oklahoma City, Oklahoma, Court of Federal Claims No: 21–1889V
66. Andrea P. Giguere, Woonsocket, Rhode Island, Court of Federal Claims No: 21–1890V
67. John R. Greene, Jr., Arlington, Virginia, Court of Federal Claims No: 21–1891V
68. Rosemary Harville, Jefferson City, Tennessee, Court of Federal Claims No: 21–1893V
69. Ann Petrea Crawford, Cedar Falls, Iowa, Court of Federal Claims No: 21–1896V
70. Maryetta Spells, Deceased, Sacramento, California, Court of Federal Claims No: 21–1898V
71. George Grace, Tonawanda, New York, Court of Federal Claims No: 21–1901V
72. Janel Trepiccione, New Cumberland, Pennsylvania, Court of Federal Claims No: 21–1902V
73. Rachel Luginsky, St. Charles, Illinois, Court of Federal Claims No: 21–1903V
74. William R. Eddington, M.D., Marmaduke, Arkansas, Court of Federal Claims No: 21–1906V
75. Sheryl Askins, Philadelphia, Pennsylvania, Court of Federal Claims No: 21–1907V
76. Genevieve Arsenault, Southbridge, Massachusetts, Court of Federal Claims No: 21–1908V
77. Yesica Valdovinos Valle on behalf of M. V., Greensboro, North Carolina, Court of Federal Claims No: 21–1909V
78. John Michael Smith, Jr., Mayville, New York, Court of Federal Claims No: 21–1911V
79. Kimberly Beth Garrett, Grand Rapids, Michigan, Court of Federal Claims No: 21–1913V
80. Kevin Slayton, Annapolis, Maryland, Court of Federal Claims No: 21–1914V
81. Anna Howle on behalf of M. L. L., Washington, District of Columbia, Court of Federal Claims No: 21–1915V
82. Limor Mazlin, Washington, District of Columbia, Court of Federal Claims No: 21–1916V
83. Jo Ellen Jackman on behalf of Kenneth W. Jackman, Clarkston, Washington, Court of Federal Claims No: 21–1917V
84. Dana Ball, Wellesley Hills, Massachusetts, Court of Federal Claims No: 21–1920V
85. Janis Rockey, Hendersonville, Tennessee, Court of Federal Claims No: 21–1921V
86. Marjorie Putnam, Washington, District of Columbia, Court of Federal Claims No: 21–1923V
87. Mary Richardson, Steubenville, Ohio, Court of Federal Claims No: 21–1924V
88. Christopher Hurtte, Granite City, Illinois, Court of Federal Claims No: 21–1925V
89. Emma M. Grim, Sacramento, California, Court of Federal Claims No: 21–1927V
90. Rebecca Egan, Palmdale, California, Court of Federal Claims No: 21–1928V
91. Robert Elliott, Sioux Falls, Iowa, Court of Federal Claims No: 21–1929V
92. Linda Slafer, Orlando, Florida, Court of Federal Claims No: 21–1930V
93. Rebeka Spengler, Des Moines, Iowa, Court of Federal Claims No: 21–1932V
94. Dianne Harvanek, Lakewood, Colorado, Court of Federal Claims No: 21–1933V
95. Tamara Bernadine on behalf of E. S., Phoenix, Arizona, Court of Federal Claims No: 21–1935V
96. Tammy Macklin, Englewood, New Jersey, Court of Federal Claims No: 21–1936V
97. Michael Johnson, Englewood, New Jersey, Court of Federal Claims No: 21–1937V
98. Bonnie McKirdy, Philadelphia, Pennsylvania, Court of Federal Claims No: 21–1938V

[FR Doc. 2021–22598 Filed 10–15–21; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request; the Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment Surveys, OMB No. 0906–0014, Revision

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

ACTION: Notice.

SUMMARY: In compliance with of the Paperwork Reduction Act of 1995, HRSA has 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 November 17, 2021.

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, email Samantha Miller, the acting HRSA Information Collection Clearance Officer at *paperwork@hrsa.gov* or call (301) 443–9094.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: The Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment Surveys. OMB No. 0906–0014—Revision.

Abstract: The purpose of the Public Health System Assessment Surveys is to inform the Advisory Committee on Heritable Disorders in Newborns and Children (Committee) on states' ability to add newborn screening for particular conditions, including the feasibility, readiness and overall capacity to screen for a new condition.

The Committee was established under the Public Health Service Act, 42 U.S.C. 217a: Advisory councils or committees (PDF—215 KB), and Title XI § 1111 (42 U.S.C. 300b–10). The purpose of the Committee is to provide the Secretary with recommendations, advice, and technical information regarding the most appropriate application of technologies, policies, guidelines, and standards for: (a) Effectively reducing morbidity and mortality in newborns and children having, or at risk for, heritable disorders; and (b) enhancing the ability of state and local health agencies to provide for newborn and child screening, counseling, and health care services for newborns and children having, or at risk for, heritable disorders. Specifically, the Committee makes systematic evidence-based recommendations on newborn screening for conditions that have the potential to change the health outcomes for newborns.

The Committee tasks an external workgroup to conduct systematic evidence-based reviews for conditions being considered for addition to the Recommended Uniform Screening Panel, and their corresponding newborn screening test(s), confirmatory test(s), and treatment(s). Reviews also include an analysis of the benefits and harms of newborn screening for a selected condition at a population level and an assessment of state public health newborn screening programs' ability to implement the screening of a new condition.

A 60-day notice published in the **Federal Register**, 86 FR 38726 (July 22, 2021).

There were no public comments.

Need and Proposed Use of the Information: The surveys are administered by the Committee's Evidence Review Group to collect data from state newborn screening programs in the United States. The surveys have been developed to capture the following: (1) Readiness of state public health newborn screening programs to expand newborn screening to include the target condition, (2) specific requirements of screening for a condition that could hinder or facilitate implementation in each state, and (3) estimated timeframes needed for each state to complete major milestones toward full implementation of newborn screening for the condition.

The following is a summary of proposed changes to the Committee's Public Health System

Assessment Surveys

Proposed changes to the ''INITIAL Survey of the Secretary's Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment'':

- Survey title:
  - *Current title:* ''INITIAL Survey of the Secretary's Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment''.
  - *Proposed change:* (strike ''Secretary's) ''INITIAL Survey of the Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment''
  - *Rationale:* Per the charter signed November 10, 2020, the ACHDNC is the correct name for the Committee.
- Introductory paragraph:
  - *Current introductory paragraph:* ''The purpose of this survey is to inform the Secretary of Health and Human Services Advisory Committee on Heritable Disorders in Newborns and Children (Committee) about states' ability to add newborn screening (NBS) for [condition x] using information gathered from most of the state and territorial NBS programs in the U.S. . . .''
  - *Proposed change:* (strike ''Secretary of Health and Human Services'') ''The purpose of this survey is to inform the Advisory Committee on Heritable Disorders in Newborns and Children (Committee) about states' ability to add newborn screening (NBS) for [condition x] using information gathered from most of the state and territorial NBS programs in the U.S. . . .''

Rationale: Per the charter signed November 10, 2020, the ACHDNC is the correct name of the Committee.

- Instructions for question 3 (grammatical edit):
  - *Current instructions:* ''. . . The following question asks you to consider, in general, how much the following factors would be an issue in considering adding [condition x] to your NBS panel.''
  - *Proposed change:* (strike ''in'', replace with ''when''), ''. . . The following question asks you to consider, in general, how much the following factors would be an issue when considering adding [condition x] to your NBS panel.''
  - *Rationale:* Correction of grammatical error.

Proposed changes to the ''FOLLOW-UP Survey of the Secretary's Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment''

- Survey title:
  - *Current title:* ''INITIAL Survey of the Secretary's Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment''.
  - *Proposed change:* (strike ''Secretary's) ''INITIAL Survey of the Advisory Committee on Heritable Disorders in Newborns and Children's Pub.
  - *Rationale:* Per the charter signed November 10, 2020, the ACHDNC is the correct name of the Committee.
- Question 9 (grammatical edits):
  - *Current question:* ''Have you developed a follow up protocol and/or educational materials for [condition x]? If so please describe the steps for short-term follow and how the plan was developed.''
  - *Proposed change:* (insert hyphen in ''follow-up'', insert ''-up'' in the phrase ''short-term follow'') ''Have you developed a follow-up protocol and/or educational materials for [condition x]? If so, please describe the steps for short-term follow-up and how the plan was developed.
  - *Rationale:* Correction of grammatical errors.

The data gathered informs the Committee on the following: (1) Feasibility of implementing population-based screening for the target condition, (2) readiness of state newborn screening programs to adopt screening for the condition, (3) gaps or limitations related to the feasibility or readiness of states to screen for a condition, and (4) areas of technical assistance and resources needed to facilitate screening for conditions with low feasibility or readiness.

Likely Respondents: The respondents to the survey will be state and territorial newborn screening programs.

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

[It is anticipated that the proposed revisions will not impact the estimated annualized burden hours]

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
INITIAL Survey of the Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment	[1] 59	[3] 2	118	10.0	1,180

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS—Continued

[It is anticipated that the proposed revisions will not impact the estimated annualized burden hours]

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
FOLLOW-UP Survey of the Advisory Committee on Heritable Disorders in Newborns and Children's Public Health System Assessment	[2] 30	[3] 2	60	2.0	120
Total	89		178		1,300

[1] The respondents to the survey will be state and territorial newborn screening programs.

[2] Up to 30 states and/or territories will be asked to complete a follow-up survey.

[3] Up to two conditions may be reviewed per year.

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. 2021–22619 Filed 10–15–21; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–4040–0009]

Agency Information Collection Request; 60-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before December 17, 2021.

ADDRESSES: Submit your comments to *sagal.musa@hhs.gov* or by calling (202) 205–2634.

FOR FURTHER INFORMATION CONTACT: When submitting comments or requesting information, please include the document identifier 4040–0009–New–60D and project title for reference, to Sagal Musa, email: *sagal.musa@hhs.gov,* or call (202) 205–2634 the Reports Clearance Officer.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (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.

Title of the Collection: Assurances for Construction Programs (SF–424D).

Type of Collection: Renewal.

OMB No.: 4040–0009.

Abstract

Assurances for Construction Programs (SF–424D) is used by applicants to apply for Federal financial assistance. The Assurances for Construction Programs (SF–424D) form allows the applicants to provide specific assurances as part of their grant proposals. This form is evaluated by Federal agencies as part of the overall grant application. This IC expires on February 28, 2022. *Grants.gov* seeks a three-year clearance of these collections.

ANNUALIZED BURDEN HOUR TABLE

Forms (if necessary)	Respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
Assurances for Construction Programs (SF–424D).	Grant-seeking organizations	353	1	0.5	176.5
Total			1		176.5

Sherrette A. Funn,

Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.

[FR Doc. 2021–22649 Filed 10–15–21; 8:45 am]

BILLING CODE 4151–AE–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–4040–0019]

Agency Information Collection Request; 60-Day Public Comment Request

AGENCY: Office of the Secretary, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.