promulgated in 42 CFR parts 70 and 71. Part 71 contains regulations to prevent the introduction, transmission, and spread of communicable diseases into the states and possessions of the United States.

Passenger and crewmember manifests are used to collect travelers' information from airlines and vessels after travel has been completed and when a disease is confirmed or there is a suspected exposure. Manifests include locating and contact information, as well as information concerning where passengers sat while aboard an airline or their location (*e.g.,* cabin numbers) and activities aboard a vessel. Manifests collect the following data elements:

- Full name (last, first, and, if available, middle or others);
- Date of birth;
- Sex;
- Country of residence;
- If a passport is required; passport number, passport country of issuance, and passport expiration date;
- If a travel document, other than a passport is required, travel document type, travel document number, travel document country of issuance and travel document expiration date;
- Address while in the United States (number and street, city, state, and zip code), except that U.S. citizens and lawful permanent residents will provide address of permanent residence in the U.S. (number and street, city, state, and zip code; as applicable);
- Primary contact phone number to include country code;
- Secondary contact phone number to include country code;
- Email address;
- Airline name;
- Flight number;
- City of departure;
- Departure date and time;
- City of arrival;
- Arrival date and time; and
- Seat number for all passengers.
- CDC also requests seat configuration for the requested contact area (example: AB/aisle/CDE/aisle/FG, bulkhead in front of row 9), identification on the manifest of the crew and what zone crew were assigned to, the identification of any babes-in-arms, and finally CDC requests the total number of passengers on board if measles is the cause of the investigation, due to the highly infectious nature of the disease.

CDC then uses this passenger and crew manifest information to coordinate with state and local health departments or International Health Regulation (IHR) National Focal Points (NFPs) so they can follow-up with residents who live or are currently located in their jurisdiction. In most cases, the manifests are issued for air travel and state and local health departments or IHR NFPs are responsible for the contact investigations; airlines and vessels may take responsibility for follow-up of crew members. In rare cases, CDC may use the manifest data to perform the contact investigation directly.

CDC requests OMB approval for an estimated 875 annual burden hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Airline Medical Officer or Equivalent/ Analysist/Travel Specialist/Manager Equivalent.	International Manifest Template/Informal Manifest Request Template.	350	1	150/60

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. 2023–13897 Filed 6–27–23; 11:15 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–R–266]

Agency Information Collection Activities: Proposed Collection; Comment Request

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

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send 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 August 28, 2023.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. 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: Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number:_____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 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/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

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

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–R–266 Medicaid Disproportionate Share Hospital (DSH) Annual Reporting Requirements

Under the 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. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires Federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Medicaid Disproportionate Share Hospital (DSH) Annual Reporting Requirements; *Use:* States are required to submit an annual report that identifies each disproportionate share hospital (DSH) that received a DSH payment under the state's Medicaid program in the preceding fiscal year and the amount of DSH payments paid to that hospital in the same year along with other information that the Secretary determines necessary to ensure the appropriateness of DSH payments; *Form Number:* CMS–R–266 (OMB control number: 0938–0746); *Frequency:* Yearly; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 51; *Total Annual Responses:* 51; *Total Annual Hours:* 2,142. (For policy questions regarding this collection contact Rich Cuno at 410–786–1111.)

Dated: June 26, 2023.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2023–13877 Filed 6–28–23; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Voluntary Acknowledgment of Paternity and Required Data Elements for Paternity Establishment Affidavits

AGENCY: Office of Child Support Services, Administration for Children and Families, United States Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Office of Child Support Services (OCSS), Administration for Children and Families (ACF), United States Department of Health and Human Services, is requesting a three-year extension of the Voluntary Acknowledgment of Paternity and Required Data Elements for Paternity Establishment Affidavits (OMB #0970–0171, expiration 1/31/2024). No changes are proposed.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: 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: Section 466(a)(5)(C) of the Social Security Act requires States to enact laws ensuring a simple civil process for voluntarily acknowledging paternity via an affidavit. The development and use of an affidavit for the voluntary acknowledgment of paternity would include the minimum requirements of the affidavit specified by the Secretary under section 452(a)(7) of the Social Security Act and give full faith and credit to such an affidavit signed in any other State according to its procedures. The State must provide that, before a mother and putative father can sign a voluntary acknowledgment of paternity, the mother and putative father must be given notice, orally, or through the use of video equipment, and in writing, of the alternatives to, the legal consequences of, and the rights (including any rights, if one parent is a minor, due to minority status) and responsibilities of acknowledging paternity. The affidavits will be used by hospitals, birth record agencies, and other entities participating in the voluntary paternity establishment program to collect information from the parents of nonmarital children.

Respondents: The parents of nonmarital children, State and Tribal agencies operating child support programs under Title IV–D of the Social Security Act, hospitals, birth record agencies, and other entities participating in the voluntary paternity establishment program.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Training	130,240	1	1	130,240
Paternity Acknowledgment Process	1,618,412	1	0.17	275,130
Data Elements	54	1	1	54
Ordering Brochures	2,604,802	1	.08	208,384

Estimated Total Annual Burden Hours: 613,808.

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the