Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a previously approved information collection requirement of GSA Form 1142, Release of Claims, regarding final payment under construction and building services contract.

DATES: Submit comments on or before: June 15, 2021.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to: *http://www.regulations.gov.* Submit comments via the Federal eRulemaking portal by searching for Information Collection 3090–0080. Select the link ''Comment Now'' that corresponds with ''Information Collection 3090–0080, Contract Financing Final Payment; GSA Form 1142, Release of Claims''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 3090–0080, Contract Financing Final Payment; GSA Form 1142, Release of Claims'' on your attached document. If your comment cannot be submitted using *https://www.regulations.gov,* call or email the points of contact in the **FOR FURTHER INFORMATION CONTACT** section of this document for alternate instructions.

Instructions: Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two-to-three days after submission to verify posting.

FOR FURTHER INFORMATION CONTACT: Bryon Boyer, Procurement Analyst, Office of Governmentwide Policy, by phone at 817–850–5580 or by email at *gsarpolicy@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration Acquisition Regulation (GSAR) clause 552.232–72 requires construction and building services contractors to submit a release of claims before final payment is made to ensure contractors are paid in accordance with their contract requirements and for work performed. GSA Form 1142, Release of Claims is used to achieve uniformity and consistency in the release of claims process.

B. Annual Reporting Burden

Respondents: 1,330.
Responses per Respondent: 1.
Annual Responses: 1,330.
Hours per Response: 0.10.
Total Burden Hours: 133.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate and based on valid assumptions and methodology; and ways to enhance the quality, utility, and clarity of the information to be collected.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the Regulatory Secretariat Division by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.*

Please cite OMB Control No. 3090–0080; Contract Financing Final Payment, GSA Form 1142, Release of Claims, in all correspondence.

Jeffrey A. Koses,

Senior Procurement Executive, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2021–07806 Filed 4–15–21; 8:45 am]

BILLING CODE 6820–61–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–21–21EK; Docket No. CDC–2021–0037]

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 proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled American Academy of Pediatrics (AAP) Neurodevelopmental Extension for Community Health Outcomes (ECHO)program on Children with Fetal Alcohol Spectrum Disorders (FASD). The purpose of this information collection is to monitor and evaluate the American Academy of Pediatrics (AAP) Neurodevelopmental Extension for Community Health Outcomes (ECHO) Program on Children with Fetal Alcohol Spectrum Disorders (FASD). The intent of the project is to improve practicing pediatrician capacity for identification and care of children with neurodevelopmental disorders, particularly prenatal exposure to alcohol, in the medical home.

DATES: CDC must receive written comments on or before June 15, 2021.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2021–0037 by any of the following methods:

- *Federal eRulemaking Portal: 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–D74, Atlanta, Georgia 30329.

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

Please note: Submit all comments through the Federal eRulemaking portal (*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–D74, Atlanta, Georgia 30329; phone: 404–639–7118; 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

American Academy of Pediatrics (AAP) Neurodevelopmental Extension for Community Health Outcomes (ECHO) Program on Children with Fetal Alcohol Spectrum Disorders (FASD)—New—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The purpose of this information collection is to monitor and evaluate the American Academy of Pediatrics (AAP) Neurodevelopmental Extension for Community Health Outcomes (ECHO) Program on Children with Fetal Alcohol Spectrum Disorders (FASD). The intent of the project is to improve practicing pediatrician capacity for identification and care of children with neurodevelopmental disorders, particularly prenatal exposure to alcohol, in the medical home.

Evaluation information will be used to monitor any incorporation of presented materials or suggestions from ECHO sessions into participating pediatric practices. Feedback also will inform any needed changes in topics, procedures, or other aspects of the program. The purpose and use of the session evaluation data will be to assure that specific information is conveyed and understood by participants for each monthly session, ongoing improvement in identification and referral by participating pediatricians, and to inform subsequent neurodevelopmental ECHO projects.

Data will be collected through secure email and will include monthly chart reviews, a monthly session evaluation survey, one overall program evaluation survey at the end of the project period, and one overall debriefing conference call at the end of the project. The target population is actively practicing pediatricians. Quantitative descriptive analyses are planned for the chart reviews. Qualitive data will be obtained from the session and program evaluation surveys, as well as the debriefing conference call. CDC requests approval for an estimated 496 annualized burden hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses	Burden per response (minutes)	Burden in hours
Pediatricians	Chart Review	15	160	12/60	480
Pediatricians	Session evaluation survey	15	8	5/60	10
Pediatricians	Program evaluation survey	15	1	5/60	1
Pediatricians	Debriefing conference call	15	1	60/60	15
Total					496

Jeffrey M. Zirger,
Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2021–07840 Filed 4–15–21; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–21–21EE; Docket No. CDC–2021–0033]

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 proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Integrated Viral Hepatitis Surveillance and Prevention Funding for Health Departments. This new data collection is for viral hepatitis (VH) case reporting data collected from the National Notifiable Diseases Surveillance System (NNDSS) which provides the primary population-based data used to describe the epidemiology of VH in the United States and for annual reporting of surveillance, prevention, and epidemiology performance measures via an Annual Performance Report.

DATES: CDC must receive written comments on or before June 15, 2021.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2021–0033 by any of the following methods:

- *Federal eRulemaking Portal: 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–D74, Atlanta, Georgia 30329.

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

Please note: Submit all comments through the Federal eRulemaking portal (*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