Component 2 funding only). Clinic-level data collection assessed CRCCP's primary outcome of interest—CRC screening rates within partner health systems—by measuring: (1) Partner health system, clinic, and patient population characteristics, (2) reporting period (for screening rates), (3) Chart review screening rate data, (4) Electronic Health Record (EHR) screening rate, and (5) Priority evidence-based EBIs and SAs.

Based on feedback from grantees and internal subject matter experts, CDC proposes use of updated data collection instruments. Specifically, CDC plans to implement a revised CRCCP grantee survey that eliminates questions related to EBI and SA implementation as these data are more accurately reported at the clinic level. Conversely, CDC will implement a revised CRCCP clinic-level data collection template with additional data variables related to EBI and SA implementation, as well as monitoring and evaluation activities, at the clinic level.

Redesigned data elements will enable CDC to better gauge progress in meeting CRCCP program goals and monitor implementation activities, evaluate outcomes, and identify grantee technical assistance needs. In addition, data collected will inform program improvement and help identify successful activities that need to be maintained, replicated, or expanded.

OMB approval is requested for three years. The total estimated annualized burden hours have decreased from 210 to 204 hours. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs)
CRCCP Grantees	CRCCP Annual Grantee Survey	30	1	24/60
	CRCCP Clinic-level Information Collection Template.	30	12	32/60
Total				

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2017–08490 Filed 4–26–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Voluntary Acknowledgement of Paternity and Required Data Elements for Paternity Establishment Affidavits.

OMB No.: 0970–0171.

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) 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 acknowledgement of paternity, the mother and putative father must be given notice, orally 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 and State and Tribal IV–D agencies, hospitals, birth record agencies and other entities participating in the voluntary paternity establishment program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents/ partner	Number of responses per respondent/ partner	Average burden hours per response	Total burden hours
Training	130,330	1	1	130,300
Paternity Acknowledgment Process	2,606,596	1	0.17	443,121
Data Elements	54	1	1	54
Data Elements	2,606,596	1	.08	208,528

Estimated Total Annual Burden Hours: 782,003

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of

having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–08510 Filed 4–26–17; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: State Plan Child Support Collection and Establishment of Paternity Title IV–D, OCSE–100.

OMB No.: 0970–0017.

Description: The Office of Child Support Enforcement has approved a IV–D state plan for each state. Federal regulations require states to amend their state plans only when necessary to reflect new or revised federal statutes or regulations or material change in any state laws, regulations, policies, or IV–D agency procedures. The requirement for submission of a state plan and plan amendments for the Child Support Enforcement program is found in sections 452, 454, and 466 of the Social Security Act.

Respondents: State IV–D Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response (hours)	Total burden hours
State Plan (OCSE–100)	54	5	.5	135
State Plan Transmittal (OCSE–21–U4)	54	5	.25	67.5

Estimated Total Annual Burden Hours: 202.5 hours.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*. Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–08506 Filed 4–26–17; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health and Human Development; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel; Chemical Synthesis Facility.

Date: June 20, 2017.

Time: 1:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6710 B Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Sathasiva B. Kandasamy, Ph.D., Scientific Review Administrator, Division of Scientific Review, National Institute of Child Health and Human Development, 6710B Rockledge Drive, Bethesda, MD 20892, (301) 435–6680, *skandasa@mail.nih.gov.*

Name of Committee: National Institute of Child Health and Human Development Special Emphasis Panel; NICHD Member Conflicts Teleconference Review.

Date: June 20, 2017.

Time: 3:30 p.m. to 5:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6710B Rockledge Drive, Bethesda, MD 20817 (Telephone Conference Call).

Contact Person: Helen Huang, Ph.D., Scientific Review Officer, Division of Scientific Review, OD, Eunice Kennedy Shriver National Institute of Child Health and Human Development, NIH, DHHS, 6710B Rockledge Drive, Bethesda, MD 20892, 301–435–8207, *helen.huang@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.864, Population Research; 93.865, Research for Mothers and Children; 93.929, Center for Medical Rehabilitation Research; 93.209, Contraception and Infertility Loan Repayment Program, National Institutes of Health, HHS)

Dated: April 21, 2017.

Michelle Trout,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–08458 Filed 4–26–17; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meeting.