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SUPPLEMENTARY INFORMATION: In accordance with 41 CFR 102-3.150(b), less than 15 calendar days' notice is being given for this meeting due to the exceptional circumstances of the COVID-19 pandemic and rapidly evolving COVID-19 vaccine development and regulatory processes. The Secretary of Health and Human Services has determined that COVID-19 is a Public Health Emergency. A notice of this ACIP meeting has also been posted on CDC's ACIP website at: <http://www.cdc.gov/vaccines/acip/index.html>. In addition, CDC has sent notice of this ACIP meeting by email to those who subscribe to receive email updates about ACIP.

Purpose: The committee is charged with advising the Director, CDC, on the use of immunizing agents. In addition, under 42 U.S.C. 1396s, the committee is mandated to establish and periodically review and, as appropriate, revise the list of vaccines for administration to vaccine-eligible children through the Vaccines for Children (VFC) program, along with schedules regarding dosing interval, dosage, and contraindications to administration of vaccines. Further, under provisions of the Affordable Care Act, section 2713 of the Public Health Service Act, immunization recommendations of the ACIP that have been approved by the Director of the Centers for Disease Control and Prevention and appear on CDC immunization schedules must be covered by applicable health plans.

Matters To Be Considered: The agenda will include discussions on Janssen (Johnson & Johnson) COVID-19 Vaccine Safety. A recommendation vote(s) is scheduled. Agenda items are subject to change as priorities dictate. For more information on the meeting agenda visit <https://www.cdc.gov/vaccines/acip/meetings/meetings-info.html>.

Meeting Information: The meeting will be webcast live via the World Wide Web; for more information on ACIP please visit the ACIP website: <http://www.cdc.gov/vaccines/acip/index.html>.

Public Participation

Interested persons or organizations are invited to participate by submitting written views, recommendations, and data. Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to

public disclosure. Comments will be posted on <https://www.regulations.gov>. Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. CDC will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. CDC will carefully consider all comments submitted into the docket.

Written Public Comment: Written comments must be received on or before April 23, 2021.

Oral Public Comment: This meeting will include time for members of the public to make an oral comment. Oral public comment will occur before any scheduled votes including all votes relevant to the ACIP's Affordable Care Act and Vaccines for Children Program roles. Priority will be given to individuals who submit a request to make an oral public comment before the meeting according to the procedures below.

Procedure for Oral Public Comment: All persons interested in making an oral public comment at the April 23, 2021 ACIP meeting must submit a request at <http://www.cdc.gov/vaccines/acip/meetings/> no later than 11:59 p.m., EDT, April 21, 2021 according to the instructions provided.

If the number of persons requesting to speak is greater than can be reasonably accommodated during the scheduled time, CDC will conduct a lottery to determine the speakers for the scheduled public comment session. CDC staff will notify individuals regarding their request to speak by email by 12:00 p.m., EDT, April 22, 2021. To accommodate the significant interest in participation in the oral public comment session of ACIP meetings, each speaker will be limited to 3 minutes, and each speaker may only speak once per meeting.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and

Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Data Collection for the Engaging Fathers and Paternal Relatives: A Continuous Quality Improvement Approach in the Child Welfare System Project (New Collection)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Planning, Research, and Evaluation (OPRE) within the Administration for Children and Families (ACF) is proposing to conduct data collection activities for the Engaging Fathers and Paternal Relatives: A Continuous Quality Improvement Approach in the Child Welfare System (Fathers and Continuous Learning in Child Welfare [FCL]) Project. This evaluation is a descriptive study of child welfare agencies' use of a continuous quality improvement process called the Breakthrough Series Collaborative (BSC) to implement strategies to improve father and paternal relative engagement in the child welfare system. The project is designed to examine the use of the BSC methodology to strengthen fathers' and paternal relatives' engagement with children involved in child welfare and to add to the evidence base on engagement strategies for fathers and paternal relatives in child welfare.

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: Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing OPREinfocollection@acf.hhs.gov. Alternatively, copies can also be obtained by writing to the Administration for Children and

Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests, emailed or written should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The FCL evaluation has three equally important aims. The first is to describe promising strategies for engaging fathers and paternal relatives in the child welfare system. The second is to assess the promise of the BSC as a continuous quality improvement framework for addressing challenges in the child welfare system, including whether and to what extent the BSC has potential, and if so, how it may be applied to other child welfare challenges. The third is to assess the extent to which agencies experienced a

shift in organizational culture in terms of the importance of father engagement.

The descriptive evaluation will build on the findings of the pilot study conducted under the umbrella generic: Formative Data Collections for ACF Program Support (OMB #0970–0531). (Site selection for the pilot study was conducted under the umbrella generic: Formative Data Collections for ACF Research (OMB #0970–0356.) It will focus on organizational changes and network supports for father and paternal relative engagement, changes in staff attitudes and skills for engaging fathers and paternal relatives, and father and paternal relative engagement outcomes. This evaluation will explore the implementation of father and paternal relative engagement strategies by examining process outcomes. By examining process outcomes, the evaluation is designed to indicate

whether strategies developed in the BSC are likely to lead to placement stability and permanency outcomes.

Data collection will take place with stakeholders in as many as five child welfare agencies implementing the BSC. Data collection activities include discussions with participating agency staff and key partners during site visits, focus groups with fathers and paternal relatives with relatively recent experience with the focal child welfare agencies, and web surveys of participating agency staff.

Respondents: Child welfare agency leaders, child welfare agency program staff and key partner staff involved in implementing the engagement strategies, and father and paternal relative clients of the agencies. Program staff may include senior leaders, managers, and frontline staff.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Avg. burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Interview topic guide	180	1	1.5	270	90
Father and paternal relative focus group protocol	72	1	1.5	108	36
Staff survey	90	2	0.33	59	20

Estimated Total Annual Burden Hours: 146.

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 information shall have practical utility; (b) the accuracy of the agency’s estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority Sec. 403. [42 U.S.C. 603] and Sec. 426. [42 U.S.C. 626].

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–08025 Filed 4–19–21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: Updates to Uniform Standard for Waiver of the Ryan White HIV/AIDS Program Core Medical Services Expenditure Requirement, OMB No. 0906–XXXX–NEW

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

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than June 21, 2021.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Information Collection Clearance Officer, Room 14N39, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call Lisa Wright-Solomon, the HRSA Information Collection Clearance Officer at (301) 443–1984.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the information request collection title for reference.

Information Collection Request Title: Updates to Uniform Standard for Waiver of the Ryan White HIV/AIDS Program Core Medical Services Expenditure Requirement, OMB No. 0906–XXXX–NEW.

Abstract: In accordance with sections 2604(c), 2612(b), and 2651(c) of the Public Health Service Act, Ryan White HIV/AIDS Program (RWHAP) recipients are required to spend not less than 75 percent of grant funds on core medical