to publish a 30-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 that summarizes the following proposed collection(s) of information for public comment:

1. *Title of Information Collection:* Medicaid Managed Care Quality including Supporting Regulations; *Type of Information Collection Request:* Extension of a currently approved collection; *Use:* Medicaid beneficiaries and stakeholders use the information collected and reported to understand the state's quality improvement goals and objectives, and to understand how the state is measuring progress on its goals. States use this information to help monitor and assess the performance of their Medicaid managed care programs. This information may assist states in comparing the outcomes of quality improvement efforts and can assist them in identifying future performance improvement subjects. CMS uses this information as a part of its oversight of Medicaid programs. *Form Number:* CMS–10553 (OMB control number: 0938–1281); *Frequency:* Annually; *Affected Public:* Private Sector, Business or other for-profits and State, Local or Tribal Governments; *Number of Respondents:* 376; *Number of Responses:* 2,655; *Total Annual Hours:* 36,010. (For questions regarding this collection contact Jennifer Maslowski at 312–886–2567.)

2. *Title of Information Collection:* External Quality Review (EQR) of Medicaid and Children's Health Insurance Program (CHIP) Managed Care, EQR Protocols, and Supporting Regulations; *Type of Information Collection Request:* Revision of a currently approved collection; *Use:* This 2022 information collection request proposes to revise the active external quality review (EQR) protocols (which were last revised in 2019). The revisions would: (1) align the existing protocols, appendices, and worksheets with the 2020 Medicaid managed care final rule, and (2) add a new protocol, Validation of Network Adequacy (RIN 0938–AS25, CMS–2480–F). A summary of these changes includes, but is not limited to, adding three elements to 42 CFR 438.358(b)(1)(iii) to include a review of elements 438.56, 438.100, and 438.114; establishing the first protocol for the new mandatory activity described in 438.358(b)(1)(iv) for network adequacy validation for managed care organizations (MCOs), prepaid inpatient health plans (PIHPs), and prepaid ambulatory health plans (PAHPs); and other formatting changes. *Form Number:* CMS–R–305 (OMB control number: 0938–0786); *Frequency:* Annually; *Affected Public:* Private Sector, Business or other for-profits and State, Local or Tribal Governments; *Number of Respondents:* 603; *Number of Responses:* 5,945; *Total Annual Hours:* 413,310. (For questions regarding this collection contact Jennifer Maslowski at 312–886–2567.)

3. *Type of Information Collection Request:* Reinstatement without change of a previously approved collection; *Title of Information Collection:* Data Submission for the Federally-facilitated Exchange User Fee Adjustment; *Use:* Section 2713 of the Public Health Service Act requires coverage without cost sharing of certain preventive health services, including certain contraceptive services, in non-exempt, non-grandfathered group health plans and health insurance coverage. The final regulations establish rules under which the third party administrator of the plan would provide or arrange for a third party to provide separate contraceptive coverage to plan participants and beneficiaries without cost sharing, premium, fee, or other charge to plan participants or beneficiaries or to the eligible organization or its plan. Eligible organizations are required to self-certify that they are eligible for this accommodation and provide a copy of such self-certification to their third party administrators. The final rules also set forth processes and standards to fund the payments for the contraceptive services that are provided for participants and beneficiaries in self-insured plans of eligible organizations under the accommodation described previously, through an adjustment in the FFE user fee payable by an issuer participating in an FFE.

CMS will use the data collections from participating issuers and third party administrators to verify the total dollar amount for such payments for contraceptive services provided under this accommodation for the purpose of determining a participating issuer's user fee adjustment. The attestation that the payments for contraceptive services were made in compliance with 26 CFR 54.9815–2713A(b)(2) or 29 CFR 2590.715–2713A(b)(2) will help ensure that the user fee adjustment is being utilized to provide contraceptive services for the self-insured plans in accordance with the previously noted accommodation. *Form Number:* CMS–10492 (OMB control number: 0938–1285); *Frequency:* Annually; *Affected Public:* Private sector (Business or other for-profits and Not-for-profit institutions); *Number of Respondents:* 861; *Total Annual Responses:* 861; *Total Annual Hours:* 12,930. (For policy questions regarding this collection contact Jacqueline Wilson at *jacqueline.wilson1@cms.hhs.gov.*)

Dated: August 2, 2022.

William N. Parham, III,

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

[FR Doc. 2022–16864 Filed 8–4–22; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No. 0970–0577]

Proposed Information Collection Activity; Evaluation of LifeSet

AGENCY: Office of Planning, Research, and Evaluation; Administration for Children and Families; Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families (ACF) at the U.S. Department of Health and Human Services is proposing additional information collection activities to assess the implementation of LifeSet, a program that provides services and supports to young adults ages 17 to 21 with previous child welfare involvement. Current data collection activities were approved under this same Office of Management and Budget (OMB) #: 0970–0577.

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 *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The proposed information collection activities are part of the second phase of a study that intends to assess the impact and implementation of LifeSet, a program that provides services and supports to young adults ages 17 to 21 with previous child welfare involvement. The program aims to support young

adults in their transition from foster care to independent living in the areas of education, employment and earnings, housing and economic well-being, social support, well-being, health and safety, and criminal involvement. It focuses on helping young adults identify and achieve their goals while developing the skills necessary for independent living.

The evaluation is part of a larger project to help ACF build the evidence base in child welfare through rigorous evaluation of programs, practices, and policies. The activities and products from this project will contribute to evidence building in child welfare and help to determine the effectiveness of a program for youth formerly in foster care on young adult outcomes.

The implementation study will collect information through video conferences and site visits to the participating program and child welfare agency. Data collection activities for the implementation study began, as previously approved by OMB. Additional protocols are proposed as part of the implementation study. Proposed information collection activities include interviews and focus groups with administrators and staff from the program developer, child welfare agency, and program providers; online survey of program staff; interviews with youth who participated in the program; and focus groups with youth who participated in the program and who received services as usual.

Respondents: Program participants, young adults receiving services as usual, agency and program administrators and staff, other program stakeholders.

ANNUAL BURDEN ESTIMATES

Instrument	Respondents	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Burden for previously approved, ongoing data collection						
Site Visit 2 Focus Group Guide for Staff.	LifeSet Specialists	12	1	1.5	18	9
LifeSet Team Supervisors						
Baseline Youth Survey	Youth Formerly in Foster Care.	470	1	0.6	282	141
Administrative data file	Agency and Program Staff	12	1	5	60	30
Burden for newly requested information collection						
Site Visit 3 Interview Guide for Administrators.	Child Welfare Agency Administrators. Licensed LifeSet Experts Provider Agency Administrators LifeSet Developer Administrators Provider Agency Administrators	22	1	1	22	11
Site Visit 3 Focus Group Guide for Staff.	LifeSet Specialists ..... LifeSet Team Supervisors Child Welfare Agency Caseworkers	28	1	1.5	42	21
LifeSet Specialist Survey	LifeSet Specialists	16	1	.3	5	3
Interview Guide for Youth ...	LifeSet Program Youth	12	1	1	12	6
Focus Group Guide for Youth.	LifeSet Program Youth ..... Services As Usual Youth	64	1	1.5	96	48

Estimated Total Annual Burden Hours: 269.

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: 42 U.S.C. 677.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–16791 Filed 8–4–22; 8:45 am]

BILLING CODE 4184–25–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Findings of Research Misconduct

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: Findings of research misconduct have been made against Janina Jiang, M.D., Ph.D. (Respondent), former Assistant Researcher in the Department of Pathology & Laboratory Medicine at the David Geffen School of