Sherri Berger,

Chief of Staff, Centers for Disease Control and Prevention.

[FR Doc. 2021–24388 Filed 11–3–21; 4:15 pm]

BILLING CODE 4163–18–C

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB #0970–0529]

Proposed Information Collection Activity; Prevention Services Data Collection

AGENCY: Children's Bureau, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF), Children's Bureau is requesting a 3-year extension of the Prevention Services Data Collection (OMB #0970–0529, expiration 7/31/2022). There are no changes requested to the form.

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 *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: Section 471(e)(4)(E) of the Social Security Act (the Act) (42 U.S.C. 671), as amended by Public Law 115–123, requires state and tribal child welfare agencies to collect and report to ACF information on children receiving prevention and family services and programs. Title IV–E Agencies must report the following:

- The specific services or programs provided.
- The total expenditures for each of the services or programs provided.
- The duration of the services or programs provided, and
- If the child was identified in a prevention plan as a candidate for foster care:
  - The child's placement status at the beginning, and at the end, of the 12-month period that begins on the date the child was identified as a candidate for foster care in a prevention plan; and
  - Whether the child entered foster care during the initial 12-month period and during the subsequent 12-month period.

To date, approximately ¾ of the Title IV–E Agencies have chosen to provide these prevention services; however, it is believed that this number will continue to increase over time as states voluntarily opt-in to the program in order to utilize IV–E funding to provide prevention programs and services to children and families.

The data collected will continue to inform federal policy decisions, program management, and responses to Congressional and Departmental inquiries. Specifically, the data will provide information about the use and availability of prevention services to children to prevent the need for foster care placement. The data contains personally identifiable information (date of birth and race/ethnicity).

Respondents: Title IV–E Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
Prevention Services Data Collection	55	2	31	3,410	1,137

Estimated Total Annual Burden Hours: 1,137.

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: Section 471(e)(4)(E) of the Act (42 U.S.C. 671), as amended by Public Law 115–123.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–24224 Filed 11–4–21; 8:45 am]

BILLING CODE 4184–29–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Request for Certification of Adult Victims of Human Trafficking

AGENCY: Office on Trafficking in Persons, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF), Office on Trafficking in Persons (OTIP), is requesting a 3-year extension of the Request for Certification of Adult Victims of Human Trafficking (RFC) form (Office of Management and Budget (OMB) #: 0970–0454, expiration 2/28/22). Minor revisions have been made to the form, including the addition of a few fields that will enable OTIP to be more responsive to congressional inquiries, federal reporting requirements, and the needs of victims.

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.* All requests should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The U.S. Department of Health and Human Services (HHS) provides letters of certification to foreign national victims of severe forms of trafficking in persons under the authority of the Trafficking Victims Protection Act of 2000 (TVPA), as amended 22 U.S.C. Section 7105(b)(1)(C) and (E). HHS delegated this authority to OTIP. Certification is required for foreign national adult victims of human trafficking in the United States to apply for federally funded benefits and services.

OTIP developed a form for potential victims and their advocates, including case managers, attorneys, law enforcement officers, service providers, and other representatives to provide the required information for certification to HHS in accordance with the TVPA of 2000, as amended. The RFC form (formerly titled Trafficking Victims Tracking System) was renamed in order to create continuity between the RFC and Request for Assistance for Child Victims of Human Trafficking (RFA) forms (OMB Control Number 0970–0362).

Since the RFC form originally received clearance, OTIP modernized its request process and launched Shepherd, an online case management system, to process requests for certification and assistance on behalf of foreign national adult and minor victims of trafficking. The PDF version of the form should only be used in exceptional circumstances when the online case management system is inaccessible. If a requester encounters issues submitting a request through Shepherd, they may submit the RFC form to OTIP as a password protected PDF to *Trafficking@acf.hhs.gov.* The form asks the requester for their identifying information, identifying information for the foreign national adult in the event the form is submitted by a case manager, and information describing the victim's case management service needs. The minor revisions made to this form enable OTIP to better fulfill its mandate in accordance with the TVPA of 2000, as amended. These revisions also enable OTIP to be more responsive to congressional inquiries, federal reporting requirements, and the needs of victims, as the information provided will be factored into policy and program development efforts.

Respondents: Potential victims of a severe form of trafficking in persons and their advocates, including case managers, attorneys, law enforcement officers, service providers, and other representatives.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Annual burden hours
Request for Certification of Adult Victims of Human Trafficking	1,300	1	1	1,300	433

Estimated Total Annual Burden Hours: 433.

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: 22 U.S.C. 7105.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–24233 Filed 11–4–21; 8:45 am]

BILLING CODE 4184–47–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0515]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Postmarketing Adverse Experience Reporting and Recordkeeping for Drug and Biological Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by December 6, 2021.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under Review—Open for Public Comments'' or by using the search function. The OMB control number for this information collection is 0910–0230. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Postmarketing Adverse Experience Reporting and Recordkeeping for Drug and Biologics Products

OMB Control Number 0910–0230—Revision

This information collection supports statutory provisions set forth in the Federal Food, Drug, and Cosmetic Act (FD&C Act) regarding the monitoring of FDA-regulated products. Specifically,