

## ANNUAL BURDEN ESTIMATES

| Information collection instrument | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|-----------------------------------|-----------------------|------------------------------------|-----------------------------------|--------------------|
| Registration Screens .....        | 648                   | 1                                  | 0.15                              | 97.20              |

*Estimated Total Annual Burden Hours:* 97.20.

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chap 35), the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW, Washington, DC 20201. Attn: ACF Reports Clearance Officer. Email address: [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). All requests should be identified by the title of the information collection.

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.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2018–10888 Filed 5–21–18; 8:45 am]

**BILLING CODE 4184–41–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

[OMB No.: 0970–0204]

#### Submission for OMB Review; Comment Request; Grants to States for Access and Visitation

*Description:* On an annual basis, States must provide OCSE with data on programs that the Grants to States for Access and Visitation Program has funded. These program reporting requirements include, but are not limited to, the collection of data on the number of parents served, types of services delivered, program outcomes, client socio economic data, referrals sources, and other relevant data.

*Respondents:* State Child Access and Visitation Programs and State and/or Local Service Providers.

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| Instruments: state and local child access program survey | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|----------------------------------------------------------|-----------------------|------------------------------------|-----------------------------------|--------------------|
| Part I: 54 states/jurisdictions .....                    | 54                    | 1                                  | 16                                | 864                |
| Part II: 296 local service grantees (estimated) .....    | 296                   | 1                                  | 16                                | 4736               |

*Estimated Total Annual Burden Hours:* 5600.

#### *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 title of the information collection. Email address: [infocollection@acf.hhs.gov](mailto: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](mailto:OIRA_SUBMISSION@OMB.EOP.GOV).

Attn.: Desk Officer for the Administration for Children and Families

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2018–10820 Filed 5–21–18; 8:45 am]

**BILLING CODE 4184–41–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

#### Proposed Information Collection Activity; Comment Request

*Proposed Projects:*

Title: Federal Child Support Portal Registration.

OMB No.: 0970–0370.

*Description:* The federal Office of Child Support Enforcement (OCSE), Division of Federal Systems, maintains the Child Support Portal (CSP), through which authorized users may view, update, or upload information for child support purposes. To securely access the Child Support Portal, authorized users must complete a one-time CSP registration form. OCSE uses information collected at the time of registration to verify the user's authorization to access the portal and to establish a secure account.

The federal Child Support Portal Registration information collection activities are authorized by 42 U.S.C. 653(m)(2), which requires the Secretary to establish and implement safeguards to restrict access to confidential information in the Federal Parent Locator Service to authorized persons,

and to restrict use of such information to authorized purposes.

*Respondents:* Employers, Financial Institutions, Insurers, and Child Support Agencies

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Estimated Total Annual Burden Hours: 97.20

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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.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2018–10910 Filed 5–21–18; 8:45 am]

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#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

##### Food and Drug Administration

[Docket No. FDA–2018–N–1787]

##### Advisory Committee; Blood Products Advisory Committee; Renewal

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; renewal of advisory committee.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the renewal of the Blood Products Advisory Committee by the Commissioner of Food and Drugs (the Commissioner). The Commissioner has determined that it is in the public interest to renew the Blood Products Advisory Committee for an additional 2 years beyond the charter expiration date. The new charter will be in effect until May 13, 2020.

**DATES:** Authority for the Blood Products Advisory Committee will expire on May 13, 2020, unless the Commissioner formally determines that renewal is in the public interest.

**FOR FURTHER INFORMATION CONTACT:** Bryan Emery, Division of Scientific Advisors and Consultants, Center for Biologics Evaluation and Research, Food and Drug Administration, 10993 New Hampshire Ave., Bldg. 71, Rm. 6268, Silver Spring, MD 20993–0002, 240–402–8054, [Bryan.emery@fda.hhs.gov](mailto:Bryan.emery@fda.hhs.gov).

**SUPPLEMENTARY INFORMATION:** Pursuant to 41 CFR 102–3.65 and approval by the Department of Health and Human Services pursuant to 45 CFR part 11 and by the General Services Administration, FDA is announcing the renewal of the Blood Products Advisory Committee. The committee is a discretionary Federal advisory committee established to provide advice to the Commissioner.

The Blood Products Advisory Committee advises the Commissioner or designee in discharging responsibilities as they relate to helping to ensure safe and effective drugs for human use and,

as required, any other product for which FDA has regulatory responsibility.

The Committee shall consist of a core of 17 voting members including the Chair. Members and the Chair are selected by the Commissioner or designee from among authorities knowledgeable in the fields of clinical and administrative medicine, hematology, immunology, blood banking, surgery, internal medicine, biochemistry, engineering, biological and physical sciences, biotechnology, computer technology, statistics, epidemiology, sociology/ethics, and other related professions. Members will be invited to serve for overlapping terms of up to 4 years. Almost all non-Federal members of this committee serve as Special Government Employees. The core of voting members may include one technically qualified member, selected by the Commissioner or designee, who is identified with consumer interests and is recommended by either a consortium of consumer-oriented organizations or other interested persons. In addition to the voting members, the Committee may include one non-voting member who is identified with industry interests.

The Commissioner or designee shall have the authority to select members of other scientific and technical FDA advisory committees (normally not to exceed 10 members) to serve temporarily as voting members and to designate consultants to serve temporarily as voting members when: (1) Expertise is required that is not available among current voting standing members of the Committee (when additional voting members are added to the Committee to provide needed expertise, a quorum will be based on the combined total of regular and added members), or (2) to comprise a quorum when, because of unforeseen circumstances, a quorum is or will be lacking. Because of the size of the Committee and the variety in the types of issues that it will consider, FDA may, in connection with a particular committee meeting, specify a quorum that is less than a majority of the current