

PLACEMENT AND TRANSFER OF UNACCOMPANIED ALIEN CHILDREN INTO ORR CARE PROVIDER FACILITIES  
[OMB #0970-0554]

Instrument	Annual total number of respondents	Annual total number of responses per respondent	Average burden minutes per response	Annual total burden hours
Family Group Entity .....	16	188	5	251
Influx Transfer Manifest .....	3	12	20	12
Influx Transfer Manual and Prescreen Criteria Review .....	216	43,333	30	4,679,964
Estimated Annual Burden Hours Total .....				4,680,227

ADMINISTRATION AND OVERSIGHT OF THE UNACCOMPANIED ALIEN CHILDREN PROGRAM  
[OMB #0970-0547]

Instrument	Annual total number of respondents	Annual total number of responses per respondent	Average burden minutes per response	Annual total burden hours
Notification of Concern .....	301	15	15	1,129
Estimated Annual Burden Hours Total .....				1,129

*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.

**Authority:** 6 U.S.C. 279; 8 U.S.C. 1232; *Flores v. Reno Settlement Agreement*, No. CV85-4544-RJK (C.D. Cal. 1996).

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020-25477 Filed 11-18-20; 8:45 am]

**BILLING CODE 4184-45-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

#### Expedited OMB Review and Public Comment: Community Services Block Grant (CSBG) Annual Report (OMB #0970-0492)

**AGENCY:** Office of Community Services, Administration for Children and

Families, Department of Health and Human Services.

**ACTION:** Request for public comment.

**SUMMARY:** The Office of Community Services (OCS), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is requesting expedited review of an information collection request from the Office of Management and Budget (OMB) and is inviting public comments on the proposed collection of data for the new Community Services Block Grant (CSBG) CARES Act Supplemental and CSBG Disaster Supplemental funding. This information will be collected through modified versions of the currently approved CSBG Annual Report (OMB #0970-0492, expiration 2/28/2023).

**DATES:** *Comments due within 60 days of publication.* In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described in this notice.

**ADDRESSES:** Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). Alternatively, copies can also be obtained by writing to the ACF, Office of Planning, Research, and Evaluation (OPRE), 330 C Street SW, Washington, DC 20201, Attn: ACF Reports Clearance Officer. All requests,

emailed or written, should be identified by the title of the information collection.

**SUPPLEMENTARY INFORMATION:**

*Description:* ACF is requesting that OMB grant a 180-day approval for this request under procedures for expedited processing. A request for review under normal procedures will be submitted within 180 days of the approval for this request. Any edits resulting from public comment will be incorporated into the submission under normal procedures. The CSBG Supplemental Annual Reports include modified versions of Modules 1, 2, and 4. Module 1 is modified to align with CSBG Disaster Supplemental and CSBG CARES State Plans and to help reduce the burden to the states. OCS modified Modules 2 and 4 to collect specific data for the supplemental funding and to reduce burden, including the removal of questions that were not pertinent to the data collection for the Supplemental Reports. OCS made additional technical modifications including minor wording, headings, and numbering revisions. Respondents are only expected to submit Module 3 once through the current CSBG Annual Report; OCS made technical revisions to allow respondents to confirm which funding source they are using—CSBG, CARES, or Disaster.

*Respondents:* State governments, including the District of Columbia and the Commonwealth of Puerto Rico, and U.S. territories and CSBG eligible entities (Community Action 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
CSBG Annual Report (States) .....	52	3	198	30,088	10,029
CSBG Annual Report (Eligible Entities) .....	1,009	3	697	2,109,819	703,273
CSBG CARES Supplemental Annual Report (States) .....	52	3	107	16,692	5,564
CSBG CARES Annual Report (Eligible Entities) .....	1,009	3	493	1,492,311	497,437
CSBG Disaster Supplemental Annual Report (States) .....	15	3	95	4,275	1,425
CSBG Disaster Supplemental Annual Report (Eligible Entities) .....	50	3	476	71,400	23,800

*Estimated Total Annual Burden Hours: 1,241,528.*

**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:** 112 Stat. 2729; 42 U.S.C. 9902(2).

**Mary B. Jones,**

*ACF/OPRE Certifying Officer.*

[FR Doc. 2020-25479 Filed 11-18-20; 8:45 am]

**BILLING CODE 4184-27-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2007-D-0369]

#### Product-Specific Guidances; Draft and Revised Draft Guidances for Industry; Availability

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

**ACTION:** Notice of availability.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) is announcing the availability of additional draft and revised draft product-specific guidances. The guidances provide product-specific recommendations on, among other things, the design of bioequivalence (BE) studies to support abbreviated new

drug applications (ANDAs). In the **Federal Register** of June 11, 2010, FDA announced the availability of a guidance for industry entitled "Bioequivalence Recommendations for Specific Products" that explained the process that would be used to make product-specific guidances available to the public on FDA's website. The guidances identified in this notice were developed using the process described in that guidance.

**DATES:** Submit either electronic or written comments on the draft guidances by January 19, 2021 to ensure that the Agency considers your comment on these draft guidances before it begins work on the final versions of the guidances.

**ADDRESSES:** You may submit comments on any guidance at any time as follows:

#### Electronic Submissions

Submit electronic comments in the following way:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the

manner detailed (see "Written/Paper Submissions" and "Instructions").

#### Written/Paper Submissions

Submit written/paper submissions as follows:

- **Mail/Hand Delivery/Courier (for written/paper submissions):** Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

**Instructions:** All submissions received must include the Docket No. FDA-2007-D-0369 for "Product-Specific Guidances; Draft and Revised Draft Guidances for Industry." Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <https://www.regulations.gov> or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240-402-7500.

- **Confidential Submissions—**To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this