arrearage by attaching and seizing assets of the obligor held in financial institutions; (2) 42 U.S.C. 666(a)(17), which requires state child support agencies to establish procedures under which the state child support agencies shall enter into agreements with financial institutions doing business in the state to develop and operate, in coordination with financial institutions and the Federal Parent Locator Service (in the case of financial institutions doing business in two or more states), a data match system, using automated data exchanges to the maximum extent feasible, in which a financial institution is required to quarterly provide information pertaining to a noncustodial parent owing past-due support who maintains an account at the institution; and (ii) in response to a notice of lien or levy, encumber or surrender, assets held; (3) 42 U.S.C. 652(a)(7), which requires OCSE to provide technical assistance to state child support enforcement agencies to help them establish effective systems for collecting child and spousal support; and (4) 45 CFR 303.7(a)(5), which requires state child support agencies to transmit requests for information and provide requested information electronically to the greatest extent possible.

Respondents: Multistate Financial Institutions and State Child Support Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Financial Data Match Result File	112	4	.33	147.84
Election Form	30	1	0.5	15
FAST Levy Response Withhold Record Specifications: State Child Support Enforcement Agencies	7	1	317.5	2,222.5
FAST Levy Response Withhold Record Specifications: Multistate Financial Institutions	5	1	317.5	1,587.5

Estimated Total Annual Burden Hours: 3,972.84.

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer.

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, 725 17th Street NW., Washington, DC 20503, Attn: Desk Officer for ACF.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 2014–10788 Filed 5–9–14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Case Plan Requirement, Title IV–E of the Social Security Act.

OMB No.: 0970–0428.

Description: Under section 471(a)(16) of title IV–E of the Social Security Act (the Act), to be eligible for payments, states and tribes must have an approved title IV–E plan that provides for the development of a case plan for each child for whom the State or Tribe receives foster care maintenance payments and that provides a case review system that meets the requirements in section 475(5) and 475(6) of the Act.

The case review system assures that each child has a case plan designed to achieve placement in a safe setting that is the least restrictive (most family-like) setting available and in close proximity to the child's parental home, consistent with the best interest and special needs of the child. Through these requirements, States and Tribes also comply, in part, with title IV–B section 422(b) of the Act, which assures certain protections for children in foster care.

The case plan is a written document that provides a narrative description of the child-specific program of care. Federal regulations at 45 CFR 1356.21(g) and section 475(1) of the Act delineate the specific information that should be addressed in the case plan. The Administration for Children and Families (ACF) does not specify a recordkeeping format for the case plan nor does ACF require submission of the document to the Federal government. Case plan information is recorded in a format developed and maintained by the State or Tribal child welfare agency.

Respondents: State and Tribe title IV–B and title IV–E agencies

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Case Plan	511,915	1	4.00	2,054,390

Estimated Total Annual Burden Hours: 2,054,390.

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, 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, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: *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 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. 2014–10791 Filed 5–9–14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0001]

Ophthalmic Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Ophthalmic Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on June 6, 2014, from 8 a.m. to 6 p.m.

Location: Hilton Washington, DC North/Gaithersburg, salons A, B, C, and D, 620 Perry Pkwy., Gaithersburg, MD 20877. The hotel telephone number is 301–977–8900.

Contact Person: Natasha Facey, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1552, Silver Spring, MD 20993, 301–796–5290, *Natasha.Facey@fda.hhs.gov,* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site at *http://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

Agenda: On June 6, 2014, the committee will discuss, make recommendations, and vote on information related to the premarket approval application for the KAMRA Inlay (Model ACI 7000) (KAMRA Inlay) submitted by applicant AcuFocus, Inc. The KAMRA Inlay is a permanent corneal implant that is placed intrastromally in a corneal pocket or under a corneal flap. The opaque annulus of the inlay reduces the aperture of the eye, which improves near vision by providing an increased depth of focus in the implanted eye. The proposed indication for use states that the KAMRA Inlay is indicated for the improvement of near and intermediate vision in presbyopic patients who require near or intermediate correction. The KAMRA Inlay is intended to be placed intrastromally in the cornea, on the visual axis, by way of a femtosecond laser-created pocket using a spot/line separation of 6x6 microns (μ) or less. The KAMRA Inlay should be placed at a depth equal to or greater than 180 μ.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before May 30, 2014. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. on June 6, 2014. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before May 21, 2014. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by May 27, 2014.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact James Clark at *James.Clark@fda.hhs.gov* or 301–796–5293 at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: May 6, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–10803 Filed 5–9–14; 8:45 am]

BILLING CODE 4160–01–P