Estimated Total Annual Burden Hours: 627.

Authority: 42 U.S.C. 652(a)(7); 42 U.S.C. 666(f); 45 CFR 301.1; 45 CFR 303.7; and 45 CFR 309.120.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–24919 Filed 11–15–21; 8:45 am]

BILLING CODE 4184–41–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Head Start Family and Child Experiences Survey (FACES) (OMB #0970–0151)

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Planning, Research, and Evaluation (OPRE), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is proposing to collect data for a new wave of the Head Start Family and Child Experiences Survey (FACES) as well as a follow-up to a special data collection fielded in the fall of 2021.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about 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.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

SUPPLEMENTARY INFORMATION:

Description: The purpose of the FACES data collection is to support the 2007 reauthorization of the Head Start program (Pub. L. 110–134), which calls for periodic assessments of Head Start's quality and effectiveness.

FACES 2019 focuses on Head Start Regions I through X (which are geographically based); AIAN (American Indian and Alaska Native) FACES 2019 focuses on Region XI (which funds Head Start programs that serve federally recognized American Indian and Alaska Native tribes). Both studies will provide data on a set of key indicators for Head Start programs. Information about the Head Start program recruitment and center selection processes and on the fall 2019, spring 2020, and fall 2021 data collection activities for both FACES and AIAN FACES can be found here: *https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202005-0970-009.*

This 30-day notice describes:

- The spring 2022 round of FACES program- and classroom-level data collection.
- A follow-up in spring 2022 of the fall 2021 FACES and AIAN FACES child-level data collection.

FACES spring 2022 data collection will take place in 180 Head Start programs nationwide. Of the 180 programs, 60 will have participated in fall data collection and 120 will be added to participate in classroom- and program-data collection only. AIAN FACES will continue in the same 22 programs that participated in 2019, 2020, and 2021 data collection. Data collection activities will include teacher sampling (for the 120 FACES programs not part of fall 2021), parent surveys, teacher child reports, staff surveys, and, for FACES, classroom observations.

In the additional 120 programs added to FACES in spring 2022, data collection will begin with sampling of FACES teachers in 240 Head Start centers. Study team members will request a list of all teachers working with Head Start-funded children.

As in fall 2021, for the spring 2022 follow-up data collection, FACES will survey the parents of 2,400 Head Start children in Regions I–X (FACES 2019) and 800 children in Region XI (AIAN FACES 2019) and ask their Head Start teachers to rate children's learning skills and social and emotional skills. Parents of sampled children (2,400 for FACES and 800 for AIAN FACES) will complete surveys on the web or by telephone about their children and family. In all 202 programs (180 for FACES and 22 for AIAN FACES), Head Start teachers will rate each sampled child (approximately 10 children per teacher) using the web or paper-and-pencil forms. Teachers, program directors, and center directors will also complete a survey, also using the web or paper-and-pencil forms, about themselves and the services and instruction at Head Start.

Respondents: Parents of Head Start children; Head Start staff.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Average burden per response (in hours)	Total/annual burden (in hours)
FACES 2019 spring 2022 special teacher sampling form from Head Start staff	240	1	.17	41
FACES 2019 special Head Start parent survey	2,400	1	.58	1,392
FACES 2019 special Head Start teacher child report	240	10	.17	408
FACES 2019 Head Start teacher survey	720	1	.67	482
FACES 2019 Head Start center director survey	360	1	.58	209
FACES 2019 Head Start program director survey	180	1	.67	121
AIAN FACES 2019 special Head Start parent survey	800	1	.58	464
AIAN FACES 2019 special Head Start teacher child report	90	9	.17	138
AIAN FACES 2019 Head Start teacher survey	90	1	.58	52
AIAN FACES 2019 Head Start center director survey	42	1	.50	21
AIAN FACES 2019 Head Start program director survey	22	1	.50	11

Estimated Total Annual Burden Hours: 3,339.

Authority: Section 640(a)(2)(D) and section 649 of the Improving Head Start for School Readiness Act of 2007.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–24951 Filed 11–15–21; 8:45 am]

BILLING CODE 4184–22–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Public Comment Request; Evidence Based Program Fidelity Surveys [OMB #0985–New]

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living is announcing that the proposed collection of information listed above has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under section 506(c)(2)(A) of the Paperwork Reduction Act of 1995. This 30-Day notice collects comments on the information collection requirements related to the information collection requirements for the Evidence Based Program Fidelity Surveys [OMB #0985–New].

DATES: Submit written comments on the collection of information by December 16, 2021.

ADDRESSES: Submit written comments and recommendations for the proposed information collection within 30 days of publication of this notice online at *www.reginfo.gov/public/do/PRAMain.* Find the information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. Written comments can also be submitted By mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

FOR FURTHER INFORMATION CONTACT: Susan Jenkins, Administration for Community Living, Washington, DC 20201, 202–795–7369 or by email: *Susan.Jenkins@acl.hhs.gov.*

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

The Administration for Community Living (ACL) is requesting approval to collect data for the Evidence Based Program Fidelity Surveys [OMB #0985–New]. The Evidence Based Program Fidelity Surveys will be used by ACL to evaluate the fidelity with which ACL's grantee organizations, under the Older Americans Act, implement the required evidence-based programs. States that receive Older Americans Act funds under Title III–D are required to spend those funds on evidence-based programs to improve the health and well-being of their clients and to reduce disease and injury. Since 2003, the aging services network has been steadily moving towards wider implementation of disease prevention and health promotion programs that are based on scientific evidence and demonstrated to improve the health of older adults. The FY 2012 Congressional appropriations law included, for the first time, an evidence-based requirement related to Title III–D funds.

The results of this information collection will be used by ACL/AoA to:

- Effectively report its results to the President, to Congress, to the Department of Health and Human Services and to the public.
- Assess the effectiveness of ACL and its grantees in monitoring program fidelity.
- Aid in program refinement and continuous improvement.

Comments in Response to the 60-Day Federal Register Notice

A notice published in the **Federal Register** on, July 12, 2021 in 86 FR 13720. There were 0 public comments received during the 60-day FRN.

Estimated Program Burden:

ACL estimates the burden associated with this collection of information as follows:

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Annual burden hours
Grantee: Program selection process and survey	103	1	2.00	206
Local Implementation Organization Survey	412	1	0.58	239
Total:	515	1	0.86	445

Dated: November 9, 2021.

Alison Barkoff,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2021–24923 Filed 11–15–21; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–1992–N–0011]

Sanyasi Raju Kalidindi; Grant of Special Termination

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has issued an order under the Federal Food, Drug, and Cosmetic Act (FD&C Act) granting special termination of the debarment of Sanyasi Raju Kalidindi. FDA based the order on a finding that Dr. Kalidindi provided substantial assistance in the investigations or prosecutions of offenses relating to a matter under FDA's jurisdiction and that terminating Dr. Kalidindi's debarment served the interest of justice and protected the integrity of the drug approval process.

DATES: The order became effective September 15, 2021.

ADDRESSES: Comments should reference Docket No. FDA–1992–N–0011 and be sent to the Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Karena Cooper, Office of Scientific Integrity, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 1, Rm. 4218, Silver Spring, MD 20993, 301 796–1612.

SUPPLEMENTARY INFORMATION: In a **Federal Register** notice dated April 21, 1993 (58 FR 21470), FDA debarred Dr. Kalidindi from providing services in any capacity to a person with an approved or pending drug product