information provided in the application, forms, and supporting documentation. Awards are made to applicants who demonstrate a high potential for providing quality primary health care services in HPSAs.

The program forms include the following: The NHSC Scholarship Program Application, Letter of Recommendation, the Authorization to Release Information, the Verification of Acceptance/Good Standing Report, the Receipt of Exceptional Financial Need Scholarship, and the Verification Regarding Disadvantaged Background.

The annual estimate of burden is as follows:

Instrument	Number of respondents	Responses/ respondent	Total responses	Hours per response	Total burden hours
NHSC Scholarship Program Application	1800	1	1800	2.0	3600
Letter of Recommendation	1800	2	3600	.50	1800
Authorization to Release Information	1800	1	1800	.10	180
Verification of Acceptance/Good Standing Report	1800	1	1800	.25	450
Receipt of Exceptional Financial Need Scholarship	100	1	100	.25	25
Verification Regarding Disadvantaged Background	300	1	300	.25	75
Total			9400		6130

E-mail comments to *paperwork@hrsa.gov* or mail the HRSA Reports Clearance Officer, Room 10–33, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: October 14, 2010.

Robert Hendricks,

Director, Division of Policy and Information Coordination.

[FR Doc. 2010–26329 Filed 10–19–10; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–11–0753]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Evaluation of the Centers for Disease Control and Prevention's Consumer Response Service Center, CDC INFO (OMB No. 0920–0753 exp. 10/31/2010) —Revision—Office for the Associate Director of Communication, Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In September 2005, the Centers for Disease Control and Prevention launched CDC–INFO, a consolidated, comprehensive effort to respond to consumer, provider and partner inquiries on a broad spectrum of public health topics by telephone or e-mail. More than 40 nationwide public health hotlines and warm lines were consolidated into one central phone number using a phased approach from 2005 to 2008. Management of CDC–INFO services is increasingly guided by a comprehensive evaluation that includes point-of-service and follow-up customer satisfaction surveys. These surveys provide the public with ongoing opportunity to express their level of satisfaction and report how they have used this information. All members of the public, health care providers and businesses can contact CDC–INFO by phone, e-mail, or postal mail to request health information or order CDC publications. CDC–INFO is a proactive, unified, and integrated approach to the delivery of public health information and is designed to contribute to improving the health and safety of the public. Customers are defined as any individual or group seeking health or public health information from CDC. This includes the public, media, medical and healthcare professionals, public health professionals, partner groups, businesses, researchers, and others.

The data collected since the approval of the original CDC–INFO study have been used for assessment of contact center performance and customer satisfaction.

This request is for a three year extension and revision of the existing data collection. Due to budget cuts, the following evaluation activities which were previously approved will be discontinued and are not included in the revised request: CDC–INFO Live Phone Follow-up Survey, Postcard Survey for Single Publication Orders, Postcard Survey for Bulk Mailing, Web Survey for Internet Publication Orders, Web Survey for E–Mailed Publication Orders, Customer Representative Survey, Special Outreach Surveys (General Public), Special Outreach Surveys (Professionals), Emergency Response Surveys (General Public), Emergency Response Surveys (Professionals). CDC–INFO will continue to offer two of the previously approved customer satisfaction surveys. The Interactive Voice Response Survey—offered in English and Spanish and the Web Survey for E–Mail Inquirers—offered in English and Spanish. Both surveys underwent minimal changes. The changes to the surveys will allow CDC–INFO to collect race/ethnicity data that is consistent with the Census form which gives participants the opportunity to identify as multi-racial.

Sample size, respondent burden, and intrusiveness have been minimized to be consistent with national evaluation objectives. There is no cost to the respondent, other than their time. The total estimated annual burden hours are 6,206.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number responses per respondent	Average burden per response (in hrs)
General Callers	Brief Interactive Voice Response Survey (English & Spanish).	92,000	1	4/60
Email Inquirers	Web Survey for E-mail Inquires (English & Spanish) ..	1,460	1	3/60

Dated: October 14, 2010.

Carol E. Walker,

Acting Reports Clear Officer, Centers for Disease Control and Prevention.

[FR Doc. 2010–26386 Filed 10–19–10; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–F–0537]

Arcadia Biosciences, Inc.; Filing of Food Additive Petition (Animal Use); Safflower Seed Meal

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Arcadia Biosciences, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of seed meal from a variety of bioengineered safflower in cattle and poultry feeds.

DATES: Submit either electronic or written comments on the petitioner's environmental assessment by November 19, 2010.

ADDRESSES: Submit electronic comments to: *http://www.regulations.gov.* Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Isabel W. Pocurull, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–453–6853, e-mail: *isabel.pocurull@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (section 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 2267) has been filed by Arcadia Biosciences, Inc., 202 Cousteau Pl., suite 105, Davis, CA 95618. The petition proposes to amend the food additive regulations in part 573 *Food Additives Permitted in Feed and Drinking Water of Animals* (21 CFR part 573) to provide for the safe use of seed meal from a variety of bioengineered safflower (*Carthamus tinctorius* L.) in cattle and poultry feeds. The safflower variety has been bioengineered to contain a gene from the water mold *Saprolegnia diclina* responsible for production of γ-linolenic acid in the seed oil. Seed meals are the ground residues obtained after processing seeds to extract their oil and are a common ingredient in livestock feed.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Division of Dockets Management (*see* **DATES** and **ADDRESSES**) for public review and comment.

Interested persons may submit to the Division of Dockets Management (*see* **ADDRESSES**) either electronic or written comments regarding this document. It is only necessary to send one set of comments. It is no longer necessary to send two copies of mailed comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register.** If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.51(b).

Dated: October 14, 2010.

William T. Flynn,

Acting Director, Center for Veterinary Medicine.

[FR Doc. 2010–26345 Filed 10–19–10; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Alcohol Abuse and Alcoholism; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 USC, as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable materials, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute on Alcohol Abuse and Alcoholism, Special Emphasis Panel, NIAAA Member Conflict Applications.

Date: October 26, 2010.

Time: 11 a.m. to 2 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 5635 Fishers Lane, Bethesda, MD 20892. (Telephone Conference Call)

Contact Person: Ranga Srinivas, PhD, Chief, Extramural Project Review Branch, EPRB, National Institute on Alcohol Abuse and Alcoholism, National Institutes of Health, 5635 Fishers Lane, Room 2085, Bethesda, MD 20892. 301–451–2067. *srinivar@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.271, Alcohol Research Career Development Awards for Scientists and Clinicians; 93.272, Alcohol National Research Service Awards for Research Training; 93.273, Alcohol Research Programs; 93.891, Alcohol Research Center Grants; 93.701, ARRA Related Biomedical Research