Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

### Inspection by Accredited Persons Program Under the Medical Device User Fee and Modernization Act of 2002—OMB Control Number 0910– 0510—Extension

The Medical Device User Fee and Modernization Act of 2002 (MDUFMA)

TABLE	1—ESTIMATED	ANNUAI	REPORTING	BURDEN <sup>1</sup>
		ANNOAL		DONDLIN

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
Request for accreditation	1	1	1	80	80

<sup>1</sup>There are no capital costs or operating and maintenance costs associated with this collection of information.

Dated: October 15, 2015. Leslie Kux, Associate Commissioner for Policy. [FR Doc. 2015–26641 Filed 10–20–15; 8:45 am] BILLING CODE 4164–01–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Meeting Notice for the President's Advisory Council on Faith-based and Neighborhood Partnerships

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the President's Advisory Council on Faith-based and Neighborhood Partnerships announces the following meetings:

Name: President's Advisory Council on Faith-based and Neighborhood Partnerships Council Meetings

Time and Date: Thursday, November 5th, 2015 1:00 p.m.–4:30 p.m. (EST) and Friday, November 6th, 2015 9:30 a.m.– 12:30 p.m. (EST)

Place: Meeting will be held at a location to be determined in the White House complex, 1600 Pennsylvania Ave. NW., Washington, DC. Space is extremely limited. Photo ID and RSVP are required to attend the event. Please RSVP to Ben O'Dell at *partnerships*@ *hhs.gov* 

The meeting will be available to the public through a conference call line. Register to participate in the conference call on Thursday, November 5th at the Web site *https://*  attendee.gotowebinar.com/register/ 7500158409923624193. Register to participate in the conference call on Friday, November 6th at the Web site https://attendee.gotowebinar.com/ register/8566024981889767937.

Status: Open to the public, limited only by space available. Conference call limited only by lines available.

Purpose: The Council brings together leaders and experts in fields related to the work of faith-based and neighborhood organizations in order to: Identify best practices and successful modes of delivering social services; evaluate the need for improvements in the implementation and coordination of public policies relating to faith-based and other neighborhood organizations; and make recommendations for changes in policies, programs, and practices.

Contact Person for Additional Information: Please contact Ben O'Dell for any additional information about the President's Advisory Council meeting at *partnerships@hhs.gov* 

Agenda: More information for the agenda for the meeting will be provided to those who register to attend in person or by conference call.

Public Comment: There will be an opportunity for public comment at the end of the meeting. Comments and questions can be sent in advance to *partnerships@hhs.gov.*  Dated: October 9, 2015. Ben O'Dell, Associate Director.

[FR Doc. 2015–26407 Filed 10–20–15; 8:45 am] BILLING CODE 4154–07–P

(Pub. L. 107-250) was signed into law

MDUFMA added a new paragraph (g) to

section 704 of the Federal, Food, Drug,

directing FDA to accredit third parties

inspections of eligible manufacturers of

'Implementation of the Inspection by

Accredited Persons Program Under the

participating in this voluntary program.

FDA estimates the burden of this

collection of information as follows:

on October 26, 2002. Section 201 of

and Cosmetic Act (21 U.S.C. 374),

(accredited persons) to conduct

class II or class III devices. FDA's guidance document entitled

Medical Device User Fee and

Accreditation Criteria" provides

information for those interested in

Modernization Act of 2002;

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### National Institutes of Health

## National Cancer Institute; Notice of Closed Meetings

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 meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, 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 Cancer Institute Special Emphasis Panel; Collaborative Research in Integrative Cancer Biology (U01).

*Date:* November 19, 2015. *Time:* 11:00 a.m. to 6:00 p.m.

*Agenda:* To review and evaluate grant applications.