helping agencies achieve a smooth and successful transition to eTS by assisting you in effectively determining your eTS strategy, selecting an eTS vendor and awarding a task order, and executing your agency-wide migration to eTS. Working together in a collaborative partnership, we can ensure timely success of this very important Presidential initiative.

Dated: February 17, 2005.

G. Martin Wagner,

Associate Administrator, Office of Governmentwide Policy.

[FR Doc. 05–3722 Filed 2–25–05; 8:45 am]

BILLING CODE 6820–14–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: OS–4040–0002]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Office of the Secretary, Grants.gov Program Management Office.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. Type of Information Collection Request: Regular;

Title of Information Collection: SF–424 Mandatory (M);

Form/OMB No.: OS–4040–0002.

Use: The SF–424(M) will become the government-wide data set for applications, plans, and related submissions under mandatory grant programs. Federal agencies and applicants/recipients under mandatory grant programs will use the standard data set and definitions for paper and electronic applications/plans/related submissions. At this time, the Federal agencies are proposing a set of data elements to be used as cover information. Additional standard data elements for other components of an application/plan, *e.g.*, a standard budget, may be proposed at a later date.

The proposed standard data set will replace numerous agency data sets and reduce the administrative burden placed on the grants community. Federal agencies will not be required to collect all of the information included in the proposed data set. The agency will identify the data that must be provided by applicants through instructions that will accompany the application package.

Frequency: Recordkeeping, Application, and on occasion;

Affected Public: Federal, State, local, or tribal governments, farms, and not for profit institutions;

Annual Number of Respondents: 1,161;

Total Annual Responses: 21,900;

Average Burden Per Response: 1 hour;

Total Annual Hours: 21,900.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access the HHS Web site address at *http://www.hhs.gov/oirm/infocollect/pending/* or e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *naomi.cook@hhs.gov,* or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the OS Paperwork Clearance Officer designated at the following address: Department of Health and Human Services, Office of the Secretary, Assistant Secretary for Budget, Technology, and Finance, Office of Information and Resource Management, Attention: Naomi Cook (4040–0002), Room 531–H, 200 Independence Avenue, SW., Washington DC 20201.

Dated: February 22, 2005.

Robert E. Polson,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. 05–3711 Filed 2–25–05; 8:45 am]

BILLING CODE 4168–17–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

State Health Fraud Task Force Grants; Availability of Funds; Request for Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

The Food and Drug Administration (FDA) is correcting notice document 04–14593 beginning on page 36091 in the issue of Monday, June 28, 2004, by making the following corrections:

On page 36091, in the first column, the second sentence under **SUMMARY** is corrected to read: ''Grant funds will be used to assist agencies in identifying and prosecuting perpetrators of health fraud and AIDS Health Fraud; obtain and disseminate information on the use of fraudulent drugs and therapies; disseminate information on approved drugs and therapies; and provide health fraud information obtained by the State Health Fraud Task Force to State health agencies, community based organizations, and FDA staff.''

On page 36091, in the first column, the **DATES** section is corrected to read: ''**DATES:** The application receipt date for new applications is April 30, 2005. The application receipt date for new applications for each subsequent year that this program is in effect will be April 30.''

On page 36091, in the first column, the **ADDRESSES** section is corrected to read:

''**ADDRESSES:** FDA is accepting new applications for this program electronically via Grants.gov. Applicants are strongly encouraged to apply electronically by visiting the Web site *http://www.grants.gov* and following instructions under 'APPLY.' The applicant must register in the Central Contractor Registration (CCR) database in order to be able to submit the application. Information about CCR is available at *http://www.grants.gov/CCRRegister.* The applicant must register with the Credential Provider for Grants.gov. Information about this requirement is available at *http://www.grants.gov/CredentialProvider.*

If applicants cannot submit applications through the electronic process, application forms are available from, and completed applications should be submitted to, Djuana Gibson, Division of Contracts and Grants Management (HFA–500), Food and Drug Administration, 5600 Fishers Lane, rm. 2131, Rockville, MD 20857, 301–827–

7177, e-mail: *dgibson@oc.fda.gov*. Application forms PHS 5161–1 are available via the Internet at: *http://www.hhs.gov/forms* (revised 7/00). Applications hand-carried or commercially delivered should be addressed to 5630 Fishers Lane (HFA–500), rm. 2131, Rockville, MD 20852. An application not received in time for orderly processing will be returned to the applicant without consideration.''

On page 36091, in the second column, **FOR FURTHER INFORMATION CONTACT** is corrected to read:

''**FOR FURTHER INFORMATION CONTACT**:

Regarding the administrative and financial management aspects of this notice: Djuana Gibson (see **ADDRESSES**).

Regarding the programmatic aspects of this notice: Stephen Toigo, Division of Federal-State Relations (DFSR), Office of Regulatory Affairs, Food and Drug Administration (HFC–150), 5600 Fishers Lane, rm. 12–07, Rockville, MD 20857, 301–827–6906, or access the Internet at: *http://www.fda.gov/ora/fed_state/default.htm*. For general ORA program information contact your Public Affairs Specialists at *http://www.fda.gov/ora/fed_state/DFSR_Activities/*.''

On page 36091, in the second column, under section I, the first paragraph is corrected to read: ''FDA will support projects covered by this notice under title XVII of the Public Health Service Act (42 U.S.C. 1702). FDA's project program is described in the Catalog of Federal Domestic Assistance, No. 93.245, and applicants are limited to States that have an existing AIDS Health Fraud Task Force or States that are in the process of developing a Health Fraud Task Force.''

On page 30692, in the first column, under section V.A, a sentence is added at the end of the paragraph that reads: ''A Current Listing of SPOCs can be found at *http://www.whitehouse.gov/omb/grants/spoc.html*.''

On page 36092, in the third column, under section VII, the paragraph is corrected to read: ''Applicants are encouraged to apply electronically (see **ADDRESSES**). If not, the original and two copies of the completed grant application Form PHS–5161–1 (revised 7/00) for State and local governments should be delivered to the Grants Management Office. The receipt date is April 30, 2005. No supplemental material or addenda will be accepted after the receipt date.''

On page 36092, in the third column, under section VIII.A, the section is corrected to read:

''*A. Submission Instructions*

Applications will be accepted during working hours, 8 a.m. to 4:30 p.m., Monday through Friday, on or before the established receipt date. Applications will be considered received on time if sent or mailed on or before the receipt date as evidenced by a legible U.S. Postal Service dated postmark or a legible date receipt from a commercial carrier, unless they arrive too late for orderly processing. Private metered postmarks shall not be acceptable as proof of timely mailing. Applications not received on time will not be considered for review and will be returned to the applicant. Applicants should note that the U.S. Postal Service does not uniformly provide dated postmarks. Before relying on this method, applicants should check with their local post office.

Do not send applications to the Center for Scientific Research, NIH. Any application sent to NIH that is then forwarded to FDA and not received in time for orderly processing will be deemed unresponsive and returned to the applicant. FDA is able to receive applications via the Internet.

The outside of the mailing package and item 2 of the application face page should be labeled 'Response to FDA–ORA–04–2.' You must submit only one application, an original and two copies, per package.''

Please note that the only change to section VIII.A is that FDA is now accepting applications via the Internet.

Dated: February 18, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–3710 Filed 2–25–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

General and Plastic Surgery 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: General and Plastic Surgery 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 April 11, 2005, from 8 a.m. to 5:30 p.m., on April 12, 2005, from 8 a.m. to 6 p.m., and on April 13, 2005, from 8 a.m. to 6 p.m.

Location: Hilton Washington DC North/Gaithersburg, Grand Ballroom, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: David Krause, Center for Devices and Radiological Health (HFZ–410), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–3090, ext. 141, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512519. Please call the Information Line or access the Internet address of *http://www.fda.gov/cdrh/panelmtg.html* for up-to-date information on this meeting.

Agenda: On April 11, 2005, the committee will hear oral presentations from the public. On April 12 and 13, 2005, the committee will discuss, make recommendations, and vote on two premarket approval applications for Silicone Gel-Filled Breast Prostheses. Background information, including the agenda and questions for the committee, will be available to the public on April 8, 2005, on the Internet at *http://www.fda.gov/cdrh/panelmtg.html*.

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 by March 28, 2005. Oral presentations from the public will be scheduled on April 11, 2005, between approximately 8 a.m. and 5:30 p.m., and on April 12 and 13, 2005, between approximately 2:45 p.m. and 3:45 p.m. Time allotted for each presentation is limited. Those desiring to make formal oral presentations should notify the contact person before March 28, 2005, and submit a brief statement of the general nature of the comments they wish to present, and the names and addresses of proposed participants.

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 AnnMarie Williams, Conference Management Staff, at 240–276–0450, ext. 113, at least 7 days in advance of the meeting.