signed by an authorized Grants Management Officer, and mailed to the recipient fiscal officer identified in the application.

Unsuccessful applicants will receive notification of the results of the application review by mail.

VI.2. Administrative and National Policy Requirements

45 CFR Part 74 and Part 92

For more information on the Code of Federal Regulations, see the National Archives and Records Administration at the following Internet address: *http://www.access.gpo.gov/nara/cfr/cfr-table-search.html.*

The following additional requirements apply to this project:

- AR–7 Executive Order 12372.
- AR–10 Smoke-Free Workplace Requirements.
- AR–11 Healthy People 2010.
- AR–12 Lobbying Restrictions.
- AR–14 Accounting System Requirements.
- AR–15 Proof of Non-Profit Status.

Additional information on these requirements can be found on the CDC Web site at the following Internet address: *http://www.cdc.gov/od/pgo/funding/ARs.htm.*

VI.3. Reporting Requirements

You must provide CDC with an original, plus two hard copies of the following reports:

1. Interim progress report, no less than 90 days before the end of the budget period. The progress report will serve as your non-competing continuation application, and must contain the following elements:

a. Description of progress made during the current budget period on program activities and objectives.

b. Current Budget Period Financial Progress.

c. New Budget Period Proposed Program Activities and measurable Objectives.

d. Budget.

e. Additional Requested Information.

f. Measures of Effectiveness.

2. Financial status report, no more than 90 days after the end of the budget period.

3. Final financial and performance reports, no more than 90 days after the end of the project period.

These reports must be mailed to the Grants Management or Contract Specialist listed in the ''Agency Contacts'' section of this announcement.

VII. Agency Contacts

For general questions about this announcement, contact: Technical Information Management Section, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341, Telephone: 770–488–2700.

For program technical assistance, contact: Kari Sapsis, Project Officer, 1600 Clifton Rd., MS E–05, Atlanta, GA 30333, Telephone: 404–639–8837, E-mail: *ksapsis@cdc.gov.*

For financial, grants management, or budget assistance, contact: Peaches Brown, Grants Management Specialist, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341, Telephone: 770–488–2738, E-mail: *prb0@cdc.gov.*

Dated: February 27, 2004.

Edward Schultz,

Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 04–4808 Filed 3–3–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

National Mammography Quality Assurance 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: National Mammography Quality Assurance 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 19, 2004, from 9 a.m. to 6 p.m.

Location: Holiday Inn, Walker/Whetstone Rooms, Two Montgomery Village Ave., Gaithersburg, MD.

Contact Person: Charles Finder, Center for Devices and Radiological Health (HFZ–240), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301–594–3332, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512397. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss the following issues:

(1) Mechanisms to reduce the regulatory and inspection burden on facilities;

(2) Whether mammographic images obtained from reconstructed compressed digital data (lossless or lossy data compression) can be used for primary interpretation or storage;

(3) Whether images obtained from digitized film-screen mammograms can be used for primary interpretation or storage; and

(4) Revisions to Mammography Quality Standards Act (MQSA) compliance guidance.

The committee will also receive updates on recently approved alternative standards, full field digital mammography accreditation and certification, the inspection demonstration program, the status of MQSA reauthorization, and the new post inspection enforcement strategy.

The MQSA compliance guidance documents, which are in a question and answer format, are available to the public on the Internet at *http://www.fda.gov/cdrh/mammography.* This guidance is updated continually in response to questions that FDA receives from the public.

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 April 5, 2004. Oral presentations from the public will be scheduled between approximately 9:30 a.m. and 10:30 a.m. on April 19, 2004. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before April 5, 2004, 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.

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 Shirley Meeks, Conference Management Staff, at 301–594–1283, ext. 105, at least 7 days in advance of the meeting.

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

Dated: February 25, 2004.

Peter J. Pitts,

Associate Commissioner for External Relations.

[FR Doc. 04–4786 Filed 3–3–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Food and Drug Administration and Food and Drug Administration Medical Device Industry Coalition Quality Systems Educational Forum: Production and Process Controls; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA), Office of Regulatory Affairs (ORA), Southwest Region (SWR), Dallas District Office (DALDO), in collaboration with the FDA Medical Device Industry Coalition (FMDIC) is announcing a public workshop entitled ''Quality Systems Educational Forum: Production and Process Controls.'' This public workshop is intended to provide information about FDA's Medical Device Quality Systems Regulation (QSR) to the regulated industry, particularly small businesses.

Date and Time: The public workshop will be held on April 23, 2004, from 8 a.m. to 5 p.m.

Location: The public workshop will be held at the Crowne Plaza Dallas Market Center Hotel, 7050 I–35 Stemmons Freeway, Dallas, TX 75247. Directions to the facility are available at the FMDIC Web site at *http://www.fmdic.org.*[1]

Contact Person: David Arvelo or Sue Thomason, Food and Drug Administration, 4040 North Central Expressway, suite 900, Dallas, TX 75204, 214–253–4952 or 214–253–4951, FAX: 214–253–4970, e-mail *oraswrsbr@ora.fda.gov*.

Registration: FMDIC has a $150 early registration fee. Early registration begins on February 1 and ends March 26, 2004. Registration is $175 from March 27 to April 9, 2004. To register online, please visit *http://www.fmdic.org/*. As an alternative, you may send registration information including name, title, firm name, address, telephone and fax numbers, and e-mail along with a check or money order for the appropriate amount payable to the FMDIC to Dr. William Hyman, Texas A&M University, Department of Biomedical Engineering, 3120 Tamu, College Station, TX 75843–3120. Course space will be filled in order of receipt of registration *with appropriate fees*. Seats are limited, please submit registration form as soon as possible. Those accepted into the course will receive confirmation. Registration will close after the course is filled. Registration at the site will be done on a space available basis on the day of the public workshop beginning at 8 a.m. The cost of registration at the site is $175 payable to the FMDIC. The registration fee will be used to offset expenses of hosting the conference, including meals, refreshments, meeting rooms, and materials.

[1]FDA has verified the Web site address, but FDA is not responsible for subsequent changes to the Web site after this document publishes in the **Federal Register**.

If you need special accommodations due to a disability, please contact David Arvelo or Sue Thomason at least 7 days in advance.

Transcripts: Transcripts of the public workshop will *not* be available due to the format of this workshop. Course handouts may be requested in writing from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A–16, Rockville, MD 20857, approximately 15 working days after the public workshop at a cost of 10 cents per page.

SUPPLEMENTARY INFORMATION: The workshop is being held in response to the interest in the topics discussed from small medical device manufacturers in the Dallas District area. FMDIC and FDA present this workshop to help achieve objectives set forth in section 406 of the Food and Drug Administration Modernization Act of 1997 (21 U.S.C. 393), which include working closely with stakeholders and maximizing the availability and clarity of information to stakeholders and the public. This is also consistent with the purposes of FDA's Regional Small Business Program, which are in part to respond to industry inquiries, develop educational materials, sponsor workshops and conferences to provide firms, particularly small businesses, with firsthand working knowledge of FDA's requirements and compliance policies. This workshop is also consistent with the Small Business Regulatory Enforcement Fairness Act of 1996 (Public Law 104–121), as outreach activities by Government agencies to small businesses.

The goal of the workshop is to present information that will enable manufacturers and regulated industry to better comply with the Medical Device QSR. The following topics will be discussed at the workshop: (1) The production and process control subsystem of the QSR, (2) FDA 483 trends and applicable regulations, (3) the business friendly approach, (4) software validation, (5) process validation, (6) product acceptance including techniques and purchasing controls, and (7) device history records.

Dated: February 26, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–4785 Filed 3–3–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Drug Abuse; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of a meeting of the Board of Scientific Counselors, NIDA.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in section 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the National Institute on Drug Abuse, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, NIDA.

Date: May 12, 2004.

Time: 8:30 a.m. to 6 p.m.

Agenda: To review and evaluate personal qualifications and performance, and competence of individual investigators.

Place: Intramural Research Program, National Institute on Drug Abuse, NIH, Johns Hopkins Bayview Campus, Bldg. C, 2nd Floor Auditorium, Baltimore, MD 21224.

Contact Person: Stephen J. Heishman, PhD, Research Psychologist, Clinical Pharmacology Branch, Intramural Research Program, National Institute on Drug Abuse, National Institutes of Health, DHHS, 5500 Nathan Shock Drive, Baltimore, MD 21224, (410) 550–1547.

(Catalogue of Federal Domestic Assistance Program Nos. 93.277, Drug Abuse Scientist Development Award for Clinicians, Scientist Development Awards, and Research Scientist Awards; 93.278, Drug Abuse National Research Service Awards for Research Training; 93.279, Drug Abuse Research Programs, National Institutes of Health, HHS)

Dated: February 26, 2004.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 04–4795 Filed 3–3–04; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Drug Abuse; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as