will be available for listening in the Board's Freedom of Information Office, and copies may be ordered for $6 per cassette by calling 202–452–3684 or by writing to: Freedom of Information Office, Board of Governors of the Federal Reserve System, Washington, DC 20551.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board, 202–452–3204.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 for a recorded announcement of this meeting; or you may contact the Board's Web site at http://www.federalreserve.gov for an electronic announcement. (The Web site also includes procedural and other information about the open meeting.)

Dated: November 1, 2000.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 00–28346 Filed 11–1–00; 10:55 am]

BILLING CODE 6210–01–P

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: Approximately 10:30 a.m., Wednesday, November 8, 2000, following a recess at the conclusion of the open meeting.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, NW., Washington, DC 20551.

STATUS: Closed.

MATTERS TO BE CONSIDERED:

1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

2. Any matters carried forward from a previously announced meeting.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board, 202–452–3204.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board's Web site at http://www.federalreserve.gov for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: November 1, 2000.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 00–28347 Filed 11–1–00; 10:56 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Dermatologic and Ophthalmic Drugs Advisory Committee; Notice of Meeting; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of October 19, 2000 (65 FR 62722). The notice announced a meeting of the Dermatologic and Ophthalmic Drugs Advisory Committee, which was scheduled for November 16, 2000. The document was published with an error. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Anita Prout, Committee Management Office (HFA–306), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–5503.

SUPPLEMENTARY INFORMATION: In FR Doc. 00–26787, appearing on page 62722 in the **Federal Register** of Thursday, October 19, 2000, the following correction is made:

1. On page 62722, in the first column, under the ''Location'' caption, ''Holiday Inn, Grand Ballroom, Two Montgomery Village Ave., Gaithersburg, MD.'' is corrected to read ''CDER Advisory Committee, conference room 1066, 5630 Fishers Lane, Rockville, MD.''

Dated: October 31, 2000.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 00–28349 Filed 11–01–00; 2:45 pm]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Clinical Chemistry and Clinical Toxicology 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). At least one portion of the meeting will be closed to the public.

Name of Committee: Clinical Chemistry and Clinical Toxicology 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 November 13, 2000, 10 a.m. to 4:30 p.m., and November 14, 2000, 8:30 a.m. to 4 p.m.

Location: Hilton, Salons C, D, and E, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: Veronica J. Calvin, Center for Devices and Radiological Health (HFZ–440), Food and Drug Administration, 2098 Gaither Rd., Rockville, MD 20850, 301–594–1243, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12514. Please call the Information Line for up-to-date information on this meeting.

Agenda: On November 13, 2000, the committee will discuss two draft guidances: ''Guidance for Prescription Use Drugs of Abuse Assays Premarket Notifications'' and ''Over the Counter (OTC) Screening Tests for Drugs of Abuse: Guidance for Premarket Notifications.'' The prescription use guidance will be available to the public on the Internet at http://www.fda.gov/cdrh/ode/odecl052.html and supersedes the document entitled ''Review Criteria for Assessment of In Vitro Diagnostic Devices for Drugs of Abuse Assays Using Various Methodologies.'' The OTC use guidance will be available to the public on the Internet at http://www.fda.gov/cdrh/ode/91.html and supersedes the document entitled ''Guidance for Premarket Submissions for Kits for Screening Drugs of Abuse To Be Used by the Consumer.'' Draft questions for the committee regarding these guidances will be available to the public on the Internet at http://www.fda.gov/cdrh/panelmtg.html. On November 14, 2000, the committee will discuss and make recommendations on a premarket notification (510(k)) for a first-of-a-kind prescription use screening device for heroin in human hair.

Procedure: On November 13, 2000, from 10 a.m. to 4:30 p.m., and on November 14, 2000, from 9 a.m. to 4 p.m., the meeting is open to the public. 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 November 6, 2000. On November 13, 2000, oral presentations from the public regarding the prescription use guidance will be scheduled between approximately 10:45 a.m. and 11:15 a.m., and oral presentations from the public regarding the OTC use guidance will be scheduled between approximately 1:15 p.m. and 2 p.m. On November 14, 2000, oral presentations from the public will be scheduled between approximately 9:15 a.m. and 9:45 a.m. and between approximately 3 p.m. and 3:30 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before November 6, 2000, 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.

Closed Committee Deliberations: On November 14, 2000, from 8:30 a.m. to 9 a.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential commercial information (5 U.S.C. 552b(c)(4)) relating to present and future agency issues.

FDA regrets that it was unable to publish this notice 15 days prior to the November 13 and 14, 2000, Clinical Chemistry and Clinical Toxicology Devices Panel of the Medical Devices Advisory Committee meeting. Because the agency believes there is some urgency to bring these issues to public discussion and qualified members of the Clinical Chemistry and Clinical Toxicology Devices Panel of the Medical Devices Advisory Committee were available at this time, the Commissioner of Food and Drugs concluded that it was in the public interest to hold this meeting even if there was not sufficient time for the customary 15-day public notice.

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

Dated: October 27, 2000.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 00–28350 Filed 11–1–00; 2:45 pm]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–10018]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration, DHHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), 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.

Type of Information Collection Request: New Collection;

Title of Information Collection: Survey of Medicaid Home and Community-Based Services Waiver and Personal Care Option Recipients for the Multi-Site Study of Medicaid Home and Community-Based Services;

Form No.: HCFA–10018 (OMB# 0938–NEW);

Use: Information collected will pertain to a description of the person, information regarding service use, unmet need for HCBS, quality of life, satisfaction with services, general health and functional status, care management and consumer direction. These data will be combined with secondary data on utilization of health care services to analyze the coordination of care; utilization; outcomes; and cost of providing services.

Frequency: One Time;

Affected Public: Individuals or Households;

Number of Respondents: 4,800; *Total Annual Responses:* 4,800;

Total Annual Hours: 3,200.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access HCFA's Web Site address at http://www.hcfa.gov/regs/prdact95.htm, or E-mail your request, including your address, phone number, OMB number, and HCFA document identifier, to Paperwork@hcfa.gov, or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Information Services, Security and Standards Group, Division of HCFA Enterprise Standards, Attention: Melissa Musotto, Room N2–14–26, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: October 24, 2000.

John P. Burke III,

HCFA Reports Clearance Officer, HCFA Office of Information Services, Security and Standards Group, Division of HCFA Enterprise Standards.

[FR Doc. 00–28206 Filed 11–2–00; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–565]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration, DHHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), 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.

Type of Information Collection Request: Extension of a previously approved collection for which approval has been expired; *Title of Information Collection:* Medicare Qualification