DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committee Meeting; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee. This meeting was announced in the **Federal Register** of June 19, 1997. The amendment is being made to reflect a change in the procedure for the meeting to add a second day for oral presentations from the public. An extension has been designated for notification of written and oral presentations. There are no other changes. This amendment will be announced at the beginning of the open portion of the meeting.

FOR FURTHER INFORMATION CONTACT: Elisa D. Harvey, Center for Devices and Radiological Health (HFZ–470), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–1180.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of June 19, 1997 (62 FR 33426), FDA announced that a meeting of the Obstetrics and Gynecology Devices Panel of the Medical Devices Advisory Committee would be held on July 14 and 15, 1997. On page 33426, in the third column, the ''*Procedure*'' portion is amended to read as follows:

Procedure: On July 14, 1997, from 9:30 a.m. to 5 p.m., and on July 15, 1997, from 8:30 a.m. to 5 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 July 8, 1997. Oral presentations from the public will be scheduled between approximately 10 a.m. and 11 a.m. on July 14, 1997, and between approximately 8:30 a.m. and 9:30 a.m. on July 15, 1997. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 8, 1997, 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.

Dated: July 1, 1997.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 97–17675 Filed 7–2–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

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: Vaccines and Related Biological Products Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA regulatory issues.

Date and Time: The meeting will be held on August 7, 1997, 12:30 p.m. to 3:30 p.m.

Location: Food and Drug Administration, Bldg. 29, conference room 121, 8800 Rockville Pike, Bethesda, MD. This meeting will be held by a telephone conference call. A speaker phone will be provided in the conference room to allow public participation in the meeting.

Contact Person: Nancy T. Cherry or Denise H. Royster, Center for Biologics Evaluation and Research (HFM–21), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852, 301–827–0314, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12388. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss the intramural scientific programs of the Laboratory of Method Development.

Procedure: On August 7, 1997, from 12:30 p.m. to 1:15 p.m., and from 2:15 p.m. to 3:15 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 July 31, 1997. Oral presentations from the public will be scheduled between approximately 2:15 p.m. and 3:15 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 31, 1997, 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 August 7, 1997, from 1:15 p.m. to 2:15 p.m., the meeting will be closed to permit discussion where disclosure would constitute a clearly unwarranted invasion of personal privacy (5 U.S.C. 552b(c)(6)). The meeting will be closed to discuss personal information concerning individuals associated with the research program.

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

Dated: June 26, 1997.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 97–17516 Filed 7-3-97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

Statement of Organization, Functions, and Delegations of Authority

Part F of the Statement of Organization, Functions, and Delegations of Authority for the Department of Health and Human Services, Health Care Financing Administration (HCFA), **Federal Register**, Vol. 62, No. 85, pp. 24120–24126, dated Friday, May 2, 1997) is amended to reflect a change to the organization structure that includes the Chief of Operations and changes to the administrative codes for the Offices of Internal Customer Support, Information Services, and Financial Management.

The specific amendments to Part F are described below:

- Section F.10.A.5. (Organization) is amended to read as follows:

1. Press Office (FAE)
2. Office of Legislation (FAF)
3. Office of Equal Opportunity and Civil Rights (FAJ)
4. Office of Strategic Planning (FAK)
5. Office of Communications and Operations Support (FAL)
6. Office of Clinical Standards and Quality (FAM)
7. Center for Beneficiary Services (FAQ)
8. Center for Health Plans and Providers (FAR)
9. Center for Medicaid and State Operations (FAS)
10. Consortium # 1 (FAU)
11. Consortium # 2 (FAV)

12. Consortium # 3 (FAW)
13. Consortium # 4 (FAX)
14. Chief of Operations (FB)
15. Office of Internal Customer Support (FBA)
16. Office of Information Services (FBB)
17. Office of Financial Management (FBC)

Dated: June 25, 1997.

Bruce C. Vladeck,

Administrator, Health Care Financing Administration.

[FR Doc. 97–17578 Filed 7–3–97; 8:45 am]

BILLING CODE 4120–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget, in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301)–443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: A Study To Review Activities Conducted To Assess Client Satisfaction With HIV/AIDS-related Clinical and Support Services—

New—A mail survey will be conducted of two groups: grantees that are currently funded under the Ryan White CARE Act (RWCA); and a purposive sample of 50 organizations that provide services to people with HIV/AIDS but are not currently funded under the RWCA. This second group of participants will be selected from the National Association of People With AIDS database.

The survey will collect information about the evaluation/tracking activities that were implemented from 1991 to 1996 to assess consumer/client satisfaction with services. The purpose of this study is to find out what types of evaluation/tracking activities have been implemented, and to identify gaps within these activities. The study will also identify ''model'' evaluation/tracking activities that have assessed consumer/client satisfaction and implemented findings to improve HIV/AIDS-related services, and consequently, have improved consumer/client satisfaction.

The study's final report will include a description of evaluation/tracking activities among organizations that provide services to people with HIV/AIDS and in-depth case studies of three model evaluations/tracking activities that can be easily replicated and used by other projects. The report will be disseminated to program-level and project-level officers as a guide on how to develop and implement effective evaluation/tracking activities on consumer/client satisfaction.

Estimates of respondent burden for the survey are as follows:

Type of respondent	Number of respondents	Responses per respondent	Average burden per response (in hours)	Total burden hours
Respondents who *have* assessed client satisfaction	463	1	1 hour	463
Respondents who *have not* assessed client satisfaction	87	1	.17	15
Total	550	1	.87	478

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: Virginia Huth, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, D.C. 20503.

Dated: June 27, 1997.

James J. Corrigan,

Acting Associate Administrator for Management and Program Support.

[FR Doc. 97–17513 Filed 7–3–97; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB review; Comment Request; Pretesting of Office of Cancer Communications Messages

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval. This proposed information collection was previously published in **Federal Register** on April 4, 1997, page 16168 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection

Title: Pretesting of Office of Cancer Communications Messages,

Type of Information Collection Request: EXTENSION (OMB# 0925–0046, expires 8/31/97).

Need and Use of Information Collection: In order to carry out NCI's legislative mandate to educate and disseminate information about cancer prevention, detection diagnosis, and treatment to a wide variety of audiences and organizations (e.g., cancer patients, their families, the general public, health providers, the media, voluntary groups, scientific and medical organizations), the Office of Cancer Communications (OCC) needs to pretest its communications strategies, concepts, and messages while they are under development. The primary purpose of this pretesting, or formative evaluation, is to ensure that the messages, communication materials, and information services created by OCC have the greatest capacity of being received, understood, and accepted by their target audiences. By utilizing appropriate qualitative and quantitative methodologies, OCC is able to (1) understand characteristics of the