SUMMARY: Notice is hereby given that a noncompetitive grant award is being made to the University of Hawaii to develop a virtual Data Center for Children and Families that contains the most comprehensive collection of data and information on Hawaii's children and families. As a Congressional setaside, this one-year project is being funded noncompetitively. The cost of this one-year project is $100,000.

FOR FURTHER INFORMATION CONTACT: K.A. Jagannathan, Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade, SW., Washington, DC 20447, Phone: 202–205–4829.

Dated: June 7, 2001.

Howard Rolston,

Director, Office of Planning, Research and Evaluation.

[FR Doc. 01–15032 Filed 6–13–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

A Conversation About Cancer Drug Development With Cancer Patient Advocates

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of meeting.

The Food and Drug Administration (FDA) is announcing the following meeting: A Conversation About Cancer Drug Development With Cancer Patient Advocates. The topics to be discussed are: Fast track, compassionate use, quality of life, the Patient Consultant Program, and other issues as they arise.

Date and Time: The meeting will be held on June 29, 2001, 2 p.m. to 4 p.m.

Location: The meeting will be held at the Westin Fairfax Hotel, Whitehall Room, 2100 Massachusetts Ave. NW., Washington, DC 20008, 202–293–2100.

Contacts: Patricia C. Delaney and JoAnn M. Minor, Office of Special Health Issues, Cancer Liaison Program, Office of International and Constituent Relations, Office of the Commissioner, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4460, FAX: 301–443–4555, or e-mail: pdelaney@oc.fda.gov and jminor@oc.fda.gov.

Registration: Send registration information (including name, title, firm name, address, telephone, fax number and/or e-mail address) to either contact person by June 22, 2001.

If you need special accommodations due to a disability, please contact Patricia C. Delaney or JoAnn M. Minor at least 7 days in advance.

Dated: June 8, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–14930 Filed 6–13–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committee for Pharmaceutical Science; 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: Advisory Committee for Pharmaceutical Science.

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 July 19, 2001, from 8:30 a.m. to 5:30 p.m. and July 20, 2001, from 8:30 a.m. to 3 p.m.

Location: Center for Drug Evaluation and Research Advisory Committee conference room 1066, 5630 Fishers Lane, Rockville, MD.

Contact: Nancy Chamberlin, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1076), Rockville, MD 20857, 301–827–7001, or e-mail: CHAMBERLINN@cder.fda.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12539. Please call the Information Line for up-to-date information on this meeting.

Agenda: On July 19, 2001, the committee will: (1) Discuss specific recommendations of the Orally Inhaled and Nasal Drug Products Subcommittee regarding dose response of locally acting nasal sprays and nasal aerosols, with particular application to bioequivalence studies; (2) hear reports and provide direction to the Nonclinical Studies Subcommittee; (3) provide comments and advice to the Risk-Based Chemistry, Manufacturing, and Controls Review Working Group for establishment of a list of low risk drugs; (4) discuss and provide direction on optimal applications of inline process controls in pharmaceutical production; and (5) discuss problems and provide comments to form a scientific basis for establishment of acceptance limits for microbiological tests that use newly developed technologies that do not rely on colony counts, and their application as process controls and product release criteria. On July 20, 2001, the committee will: (1) Provide comments and advice on methods to determine drug transfer into breast milk and interpretation of data; and (2) discuss and provide comments on the feasibility, scientific challenges, and approaches for establishment of pharmaceutical equivalence, bioavailability, and bioequivalence of liposome drug products.

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 July 12, 2001. Oral presentations from the public will be scheduled between approximately 1:15 p.m. to 2:15 p.m. on July 19, 2001, and between approximately 10:15 a.m. to 11:15 a.m. on July 20, 2001. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 12, 2001, 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.

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

Dated: June 6, 2001.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 01–14928 Filed 6–13–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Orally Inhaled and Nasal Drug Products Subcommittee of the Advisory Committee for Pharmaceutical Science; 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: Orally Inhaled and Nasal Drug Products Subcommittee of the Advisory Committee for Pharmaceutical Science.

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 July 17, 2001, from 8:30 a.m. to 5:30 p.m.

Location: Center for Drug Evaluation and Research Advisory Committee conference room 1066, 5630 Fishers Lane, Rockville, MD.

Contact: Nancy Chamberlin, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093) Rockville, MD 20857, 301–827–7001, or e-mail: CHAMBERLINN@cder.fda.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12539. Please call the Information Line for up-to-date information on this meeting.

Agenda: The subcommittee meeting will discuss the issue of dose-response of locally acting nasal sprays and nasal aerosols, with particular application to bioequivalence studies.

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 July 10, 2001. Oral presentations from the public will be scheduled between approximately 11:30 a.m. to 12:30 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 10, 2001, 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.

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

Dated: June 6, 2001.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 01–14929 Filed 6–13–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA–10043]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration, HHS.

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:* Evaluation of the BadgerCare Medicaid Demonstration; *Form No.:* HCFA–10043 (OMB #0938–NEW); *Use:* The subject surveys are components of the HCFA evaluation of the Wisconsin BadgerCare Section 1115 Medicaid demonstration and Title XXI (SCHIP) program. The goals of the evaluation are to assess the effectiveness of BadgerCare in reducing the number of Wisconsin residents who lack health insurance, increasing participation of eligible children in the SCHIP program, and supporting families making transitions from welfare to work. Other specific features of BadgerCare will be examined as well, including the State's outreach efforts and policy of charging premiums to selected families. Findings from the study will help to inform HCFA policy regarding Medicaid demonstrations and SCHIP, and will help states in designing similar health insurance programs.; *Frequency:* Other: One time; *Affected Public:* Individuals or Households; *Number of Respondents:* 5,680; *Total Annual Responses:* 5,680; *Total Annual Hours:* 1,914.

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: Dawn Willinghan, HCFA–10043, Room N2–14–26, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: June 5, 2001.

John P. Burke III,

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

[FR Doc. 01–14946 Filed 6–13–01; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA–2746]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration, HHS.

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.