(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.

Type of Information Collection Request: Extension of a currently approved collection; *Title of Information Collection:* End Stage Renal Disease Death Notification; *Form No.:* HCFA–2746 (OMB# 0938–0448); *Use:* This form is completed by all Medicare approved ESRD facilities upon death of an ESRD patient. The forms primary purpose is to collect fact and cause of death. Reports of deaths are used to show cause of death and demographic characteristics of these patients; *Frequency:* On occasion; *Affected Public:* Business or other for-profit; Federal Gov't., Not-for-profit institutions; *Number of Respondents:* 4,000; *Total Annual Responses:* 56,258; *Total Annual Hours:* 9,564.

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: 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. 01–14947 Filed 6–13–01; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Request for Clearance To Evaluate the National Institutes of Health (NIH) Small Business Innovation Research (SBIR) Program

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 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.

Proposed Collection

Title: The National Survey to Evaluate the NIH SBIR Program. Type of Information Collection Request: NEW. Need and Use of the Information Collection: The NIH, Office of Extramural Research, Office of Extramural Programs seeks to obtain OMB's approval to conduct a survey to evaluate the Small Business Innovation Research (SBIR) Program. The SBIR Program, established by Congress in 1982 (Public Law 97–219) and recently reauthorized through September 30, 2008 (P.L. 106–554), provides research support to small businesses for innovative technology. Primary objectives are to assess the extent to which SBIR program goals are being met, particularly those dealing with the commercialization of research products, processes or services and the uncovering of new knowledge that will lead to better health for everyone. With survey information, NIH is enabled to accurately assess the results of its large financial investment in funding innovative research conducted by small business concerns. Findings will help to (1) understand if innovative projects supported through the NIH SBIR Program are being commercialized, and if so, to classify the types of products, processes or services that are derived through SBIR funding; (2) determine if other measures of success defined within the NIH mission are being achieved; and (3) enhance NIH's administration of the SBIR Program and the support that it provides to small business concerns. Overall, the NIH will use the survey results to assess the outcomes from NIH-supported SBIR awards. OD will collect information from SBIR awardees using an Internet survey. The online survey will be implemented using Secure Socket Layer (SSL) encryption technology and password access. OD will use first-class mail and email messages to advise awardees that they have been selected to participate in the survey. Frequency of Reponse: One time survey. Affected Public: Small business concerns supported by NIH through the SBIR Program. Type of Respondents: For-profit small business concerns that have received NIH SBIR awards. The annual reporting burden is as follows: Estimated Number of Respondents: 1,000; Estimated Number of Responses Per Respondent: 1; Averaged Burden Hours Per Response: .5; and Estimated Total Annual Burden Hours Requested: 500. The annualized cost to the public is estimated at $37,500. There are no Capital Costs, Operating Costs and/or Maintenance Costs to report.

Requests for Comments

Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (2) The accuracy of the agency's estimate of the burden (including hours and cost) of the proposed information collection; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on respondents, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Ms. Jo Anne Goodnight, NIH SBIR/STTR Program Coordinator, Rockledge II Bldg., Room 6186, 6701 Rockledge Drive, Bethesda, MD 20892–7910, or call non-toll-free number (301) 435–2688 or email your request, including your address, to: *jg128w@nih.gov.*

COMMENTS DUE DATE: Comments regarding this information collection are best assured of having their full effect if received on or before August 13, 2001.

Dated: June 7, 2001.

Jo Anne Goodnight,

Coordinator, Small Business Innovation Research/Small Business Technology Transfer Program, Office of Extramural Programs, Office of Extramural Research, National Institutes of Health.

[FR Doc. 01–14972 Filed 6–13–01; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; The Jackson Heart Study: Annual Follow-Up With Third Party Respondents

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Heart, Lung, and Blood Institute (NHLBI), the National Institute of Health