With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (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 automated collection techniques when appropriate, and other forms of information technology.

AoA plans to submit to the Office of Management and Budget for approval, an extension, with no revisions, of a semi-annual reporting form and instructions pursuant to requirements in Title IV of the Older Americans Act. AoA estimates the burden of this collection of information as follows: *Frequency:* Semi-annually. *Respondents:* States, public agencies, private nonprofit agencies, institutions of higher education, and organizations including tribal organizations. *Estimated Number of Responses:* 300. *Total Estimated Burden Hours:* 12,000.

Dated: April 3, 2004.

Josefina G. Carbonell,

Assistant Secretary for Aging.

[FR Doc. 04–9594 Filed 4–27–04; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration on Aging

Agency Information Collection Activities; Proposed Collection; Comment Request; Extension of Supplemental Form to the Financial Status Report for All AoA Title III Grantees

AGENCY: Administration on Aging, HHS.

ACTION: Notice.

SUMMARY: The Administration on Aging (AoA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to the Supplemental form to the Financial Status Report for all AoA Title III Grantees.

DATES: Submit written or electronic comments on the collection of information by June 28, 2004.

ADDRESSES: Submit electronic comments on the collection of information to: *Margaret.Tolson@aoa.gov*.

Submit written comments on the collection of information to Administration on Aging, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Margaret Tolson, Director of Grants Management, Administration on Aging, Washington, DC 20201.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, AoA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (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 automated collection techniques when appropriate, and other forms of information technology.

The Supplemental form to the Financial Status Report for all AoA Title III Grantees provides an understanding of how projects funded by the Older Americans Act are being administered by grantees, in conformance with legislative requirements, pertinent Federal regulations and other applicable instructions and guidelines issued by Administration on Aging (AoA). This information will be used for Federal oversight of Title III Projects. AoA estimates the burden of this collection of information as follows: 56 State Agencies on Aging respond semiannually which should be an average burden of 1 hour per State agency per submission.

Dated: April 23, 2004.

Josefina G. Carbonell,

Assistant Secretary for Aging.

[FR Doc. 04–9595 Filed 4–27–04; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Implementation of the National Violent Death Reporting System, Program Announcement Number 04061

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Implementation of the National Violent Death Reporting System, Program Announcement Number 04061.

Times and Dates: 8:30 a.m.–8:50 a.m., May 17, 2004 (Open), 8:50 a.m.–4:00 p.m., May 17, 2004 (Closed).

Place: Marriott Atlanta Century Center, 2000 Century Boulevard NE, Atlanta, GA 30345, Telephone 404.325.0000.

Status: Portions of the meeting will be closed to the public in accordance with provisions set forth in Section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters to be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement Number 04061.

FOR FURTHER INFORMATION CONTACT: James Belloni, Deputy Director, National Center for Injury Prevention and Control, Office of the Director, Office of Program Management, CDC, 4770 Buford Highway, NE, MS–K62, Atlanta, GA 30341, Telephone 770.488.4538.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register**

notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: April 21, 2004.

Bill Atkinson,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 04–9584 Filed 4–27–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004D–0146]

Draft Guidance for Industry: Validation of Analytical Procedures for Type C Medicated Feeds

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance for industry (#135) entitled ''Validation of Analytical Procedures for Type C Medicated Feeds.'' This draft guidance represents the agency's current thinking on the characteristics that should be considered during the validation of non-microbiological analytical procedures for the analysis of drugs in Type C medicated feeds included as part of original and supplemental new animal drug applications (NADAs) and abbreviated new animal drug applications (ANADAs) for Type A medicated articles submitted to the FDA. This draft guidance is the first in a series of three guidances that will discuss assay methods for Type C medicated feeds.

DATES: Submit written or electronic comments on the draft guidance by July 12, 2004, to ensure their adequate consideration in preparation of the final document. General comments on agency guidance documents are welcome at any time.

ADDRESSES: Submit written requests for single copies of the draft guidance to the Communications Staff (HFV–12), Center for Veterinary Medicine (CVM), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855.

Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments on the draft guidance via the Internet at *http://www.fda.gov/dockets/ecomments*. Comments should be identified with the full title of the draft guidance and the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Mary G. Leadbetter, Center for Veterinary Medicine (HFV–141), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6964, e-mail: *mleadbet@cvm.fda.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

This draft guidance document provides assistance and recommendations to industry on how to consider the various validation characteristics for each analytical procedure used in medicated feed assays submitted as part of original and supplemental NADAs and ANADAs.

II. Paperwork Reduction Act of 1995

According to the Paperwork Reduction Act of 1995, a collection of information must display a valid OMB control number. The existing valid OMB control numbers for this information collection are 0910–0032 and 0910–0154. This draft guidance contains no new collections of information.

III. Significance of Guidance

This draft Level 1 guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). This draft guidance, when finalized, will represent the agency's current thinking on the topic. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternate method may be used as long as it satisfies the requirements of applicable statutes and regulations.

IV. Comments

This draft guidance is being distributed for comment purposes only and is not intended for implementation at this time. Interested persons may submit written comments to the Division of Dockets Management (see **ADDRESSES**) regarding this draft guidance document. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments should be identified with the docket number found in brackets in the heading of this document. A copy of the document and received comments are available for public examination in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

V. Electronic Access

Copies of the draft guidance document entitled ''Validation of Analytical Procedures for Type C Medicated Feeds'' may be obtained from the CVM Home Page (*http://www.fda.gov/cvm*) and from the Division of Dockets Management Web site (*http://www.fda.gov/ohrms/dockets/default.htm*).

Dated: April 19, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–9566 Filed 4–27–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of an Exclusive License: Interleukin-2 Stimulated T Lymphocyte Cell Death for the Treatment of Autoimmune Diseases, Allergic Responses, and Graft Rejection

AGENCY: National Institutes of Health, Public Health Service, DHHS.

ACTION: Notice.

SUMMARY: This notice, in accordance with 35 U.S.C. 209(c)(1) and 37 CFR 404.7(a)(1)(i), announces that the National Institutes of Health, Department of Health and Human Services, is contemplating the grant of an exclusive license to practice the inventions embodied in U.S. Patent No. 6,083,503, entitled ''Interleukin-2 stimulated T lymphocyte cell death for the treatment of autoimmune diseases, allergic responses, and graft rejection;'' U.S. Patent No. 5,989,546, entitled ''Interleukin-2 stimulated T lymphocyte cell death for the treatment of allergic responses;'' and U.S. Patent No. 5,935,575, entitled ''Interleukin-4 stimulated T lymphocyte cell death for the treatment of allergic disorders'' to Kasha Corporation, having a place of business in Rockville, Maryland. The patent rights in these inventions have been assigned to the United States of America.

The prospective exclusive license territory will be worldwide and the field of use may be limited to therapeutics for the treatment of autoimmune diseases.

DATES: Only written comments and/or application for a license which are received by the NIH Office of Technology Transfer on or before June 28, 2004 will be considered.