

20170, 703-421-5826, FAX 703-444-1737.

**Registration:** Preregistration is recommended on or before May 29, 2002. Onsite registration will be done on a space-available basis on both days of the workshop, beginning at 7:30 a.m. You may obtain registration forms and information about registration fees from HelmsBriscoe Resource One (see the *Contact* section of this document) or from Joseph Wilczek, Project Manager, at wilczek@cber.fda.gov. Mail or fax your registration information and registration fee to HelmsBriscoe Resource One by May 29, 2002.

If you need special accommodations due to a disability, please contact HelmsBriscoe Resource One at least 7 days in advance.

**Transcripts:** Transcripts of the public workshop may be requested in writing from the Freedom of Information Office (HFI-35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A-16, Rockville, MD 20857, approximately 15 working days after the meeting at a cost of 10 cents per page. The public workshop transcript will also be available on the Internet at <http://www.fda.gov/cber/minutes/workshop-min.htm>.

**SUPPLEMENTARY INFORMATION:** FDA and PPTA are jointly cosponsoring a public workshop on comparability studies for human plasma-derived therapeutics. The workshop will discuss critical issues and approaches for establishing the comparability of human plasma derivatives for supporting changes in manufacturing processes, equipment, or facilities. On May 30, 2002, the workshop will address the three levels of comparability studies—physical/chemical characterization, preclinical studies, and clinical evaluations as they are related to manufacturing changes for a human plasma derivative, as well as information on reporting manufacturing changes, comparability protocols, and several case studies.

On May 31, 2002, the workshop will focus on issues related to comparing fractionation intermediates, a topic specific to the plasma derivative industry. FDA will present historical perspectives and current guidance on cooperative manufacturing arrangements. Industry will discuss the current status of the necessity for fractionation intermediates from sources outside of the company and the criteria for acceptance. The complexities involved in characterizing the source material, intermediates, and the drug products will be discussed. The public workshop agenda will be posted on the

Internet at <http://www.fda.gov/cber/whatsnew.htm>.

Dated: April 29, 2002.

**Margaret M. Dotzel,**

*Associate Commissioner for Policy.*

[FR Doc. 02-11208 Filed 5-6-02; 8:45 am]

**BILLING CODE 4160-01-S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### Joint Meeting of the Nonprescription Drugs Advisory Committee and the Gastrointestinal Drugs Advisory Committee

**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 Committees:*

Nonprescription Drugs Advisory Committee and the Gastrointestinal Drugs 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 June 21, 2002, from 8 a.m. to 4:30 p.m.

*Location:* Holiday Inn, Versailles Ballroom, 8120 Wisconsin Ave., Bethesda, MD.

*Contact Person:* Sandra Titus, 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: [Tituss@cder.fda.gov](mailto:Tituss@cder.fda.gov), or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area) code 12541. Please call the Information Line for up-to-date information on this meeting.

*Agenda:* The committees will consider the safety and efficacy of new drug application (NDA) 21-229, proposing over-the-counter (OTC) use of PRILOSEC1 (omeprazole magnesium), AstraZeneca LP/Procter and Gamble, for the prevention of the symptoms of frequent heartburn. The sponsor proposes a 20 milligram dose to be taken for 14 days. The background material for this meeting will be posted one working day before the meeting under the Nonprescription Drugs Advisory Committee (NDAC) on the Dockets Management Branch Web site at

<http://www.fda.gov/ohrms/dockets/ac/acmenu.htm>. (Click on the year 2002 and scroll down to NDAC.)

*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 June 12, 2002. Oral presentations from the public will be scheduled on June 21, 2002, between approximately 8:15 a.m. and 9:15 a.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before June 12, 2002, 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.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Sandra Titus at least 7 days in advance of the meeting.

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

Dated: April 29, 2002.

**Linda A. Suydam,**

*Senior Associate Commissioner for Communications and Constituent Relations.*

[FR Doc. 02-11205 Filed 5-6-02; 8:45 am]

**BILLING CODE 4160-01-S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Resources and Services Administration

#### Fiscal Year 2002 Competitive Application Cycle for the Radiation Exposure Screening and Education Program 93.257

**AGENCY:** Health Resources and Services Administration, HHS.

**ACTION:** Correction.

**SUMMARY:** In notice document FR Doc. 02-10634, in the issue of Tuesday, April 30, 2002, make the following correction:

On page 21257 in the third column, under section "Funding Preferences," replace the first bullet (which reads

“Applicants that propose a Statewide service area.”) with the following bullet:

- Applicants that propose a Statewide service area and provide or arrange for services at multiple locations to serve a widely dispersed population so that all eligible individuals throughout the State have access to services.

Dated: April 30, 2002.

**Jane M. Harrison,**

*Director, Division of Policy Review and Coordination.*

[FR Doc. 02-11209 Filed 5-6-02; 8:45 am]

BILLING CODE 4165-15-P

## DEPARTMENT OF THE INTERIOR

### Fish and Wildlife Service

#### Information Collection To Be Submitted to the Office of Management and Budget (OMB) for Approval Under the Paperwork Reduction Act for the Woodcock Singing Ground Survey

**AGENCY:** Fish and Wildlife Service, Interior.

**ACTION:** Notice; request for comments.

**SUMMARY:** The U.S. Fish and Wildlife Service will submit the collection of information listed below to OMB for approval under the provisions of the Paperwork Reduction Act. A copy of the information collection requirement is included in this notice. If you wish to obtain copies of the proposed information collection requirement, related forms, and explanatory material, contact the Service Information Collection Clearance Officer at the address listed below.

**DATES:** We will accept comments until July 8, 2002.

**ADDRESSES:** Send your comments and suggestions on the requirement to the Information Collection Clearance Officer, U.S. Fish and Wildlife Service, ms 222-ARLSQ, 1849 C Street, NW., Washington, DC 20204.

**FOR FURTHER INFORMATION CONTACT:** To request a copy of the information collection request, explanatory information and related forms, contact Rebecca A. Mullin at (703) 358-2287, or electronically to [rmullin@fws.gov](mailto:rmullin@fws.gov).

**SUPPLEMENTARY INFORMATION:** The Office of Management and Budget (OMB) regulations at 5 CFR 1320, which implement provisions of the Paperwork Reduction Act of 1995 (Pub. L. 104-13), require that interested members of the public and affected agencies have an opportunity to comment on information collection and recordkeeping activities (see 5 CFR 1320.8(d)). The U.S. Fish and Wildlife Service (We) plan to submit a

request to OMB to renew its approval of the collection of information for the North American Woodcock Singing Ground Survey. We are requesting a 3-year term of approval for this information collection activity.

Federal agencies may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this collection of information is 1018-0019.

The Migratory Bird Treaty Act (16 U.S.C. 703-711) and Fish and Wildlife Act of 1956 (16 U.S.C. 742d) designate the Department of the Interior as the key agency responsible for the wise [Page 7661] management of migratory bird populations frequenting the United States and for the setting of hunting regulations that allow appropriate harvests that are within the guidelines that will allow for those populations' well being. These responsibilities dictate the gathering of accurate data on various characteristics of migratory bird populations. The North American Woodcock Singing Ground Survey is an essential part of the migratory bird management program. This survey is conducted annually by State and Federal conservation agencies to provide the necessary data to determine the population status of the woodcock. The information is used primarily by us to develop recommendations for hunting regulations. It is also used by us, State conservation agencies, University associates and other interested parties for various research and management projects. Without information on the population's status, we might promulgate hunting regulations that were too liberal thus causing harm to the woodcock population, or too conservative, thus unduly restricting recreational opportunities afforded by woodcock hunting.

*Title:* North American Woodcock Singing Ground Survey.

*Approval Number:* 1018-0019.

*Service Form Number:* 3-156.

*Frequency of Collection:* Annually.

*Description of Respondents:* State, local, tribal, provincial, or Federal employees.

*Total Annual Burden Hours:* The reporting burden is estimated to average 0.67 hours per respondent. With an estimated 40% entering data electronically, the reporting burden is estimated to average 0.75 hours per respondent. The Total Annual Burden hours is 527 hours.

*Total Annual Responses:* About 750 individuals are expected to participate in the survey.

We invite comments concerning this renewal on: (1) Whether the collection of information is necessary for the proper performance of our migratory bird management functions, including whether the information will have practical utility; (2) the accuracy of our estimate of the burden of the collection of information; (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. The information collections in this program are part of a system of record covered by the Privacy Act (5 U.S.C. 552(a)).

Dated: April 23, 2002.

**Rebecca A. Mullin,**

*Information Collection Officer, Fish and Wildlife Service.*

[FR Doc. 02-11317 Filed 5-6-02; 8:45 am]

BILLING CODE 4310-55-M

## DEPARTMENT OF THE INTERIOR

### Fish and Wildlife Service

#### Information Collection To Be Submitted to the Office of Management and Budget (OMB) for Approval Under the Paperwork Reduction Act Mourning Dove Call Count Survey

**AGENCY:** Fish and Wildlife Service, Interior.

**ACTION:** Notice; request for comments.

**SUMMARY:** The U.S. Fish and Wildlife Service will submit the collection of information listed below to OMB for approval under the provisions of the Paperwork Reduction Act. A copy of the information collection requirement is included in this notice. If you wish to obtain copies of the proposed information collection requirement, related forms, and explanatory material, contact the Service Information Collection Clearance Officer at the address listed below.

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**SUPPLEMENTARY INFORMATION:** The Office of Management and Budget (OMB)