interference to non-emergency services or communications, such as garage door openers or home appliances, as well as amateur or Citizens Band (CB) radio issues.

FCC Form 2000 will allow the Commission to collect detailed information from complainants concerning possible violations of the Act and the Commission's rules, which will enable the Commission to investigate such allegations more efficiently and to initiate enforcement actions against violators as appropriate. By collecting complaint information in a single, comprehensive template, the form will provide a standardized way for complainants to provide their information, thus reducing the need for further documentation or questions from FCC investigators to determine whether violations have occurred. This approach will ensure that complainants present their information in a way that maximizes the FCC's ability to take enforcement action against violators and protects complainants from violations that are unjust, unreasonable, and potentially hazardous to life and property. Additionally, FCC Form 2000's format reduces the need for complainants to compose narratives with all the information necessary for the Commission to begin an investigation, principally by including fields for and examples of the information most commonly needed for investigations of the most common types of violations. The form will allow the Commission to gather and review this information more efficiently. The information collected by FCC Form 2000 may ultimately become the foundation for enforcement actions and/or rulemaking proceedings, as appropriate.

FCC Form 475–B, Obscene, Profane, and Indecent Complaint Form. This form is used by consumers to lay out precisely their complaint(s) and issue(s) concerning the practices of the communications entities, which consumers believe may have aired obscene, profane, and/or indecent programming. FCC Form 475–B will remain unchanged.

Note: In this document, the Commission corrects inaccuracies published in 71 FR 53686, September 12, 2006, regarding OMB Control No. 3060–0874.

Federal Communications Commission.

William F. Caton,

Deputy Secretary.

[FR Doc. E7–10575 Filed 5–31–07; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Announcement of First Meeting of the Physical Activity Guidelines Advisory Committee

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science.

ACTION: Notice of meeting.

Authority: 42 U.S.C. 217a, section 222 of the Public Health Service Act, as amended. The Committee is governed by the provision of Public Law 92–463, as amended (5 U.S.C. Appendix 2), which sets forth standards for the formation and use of advisory committees.

SUMMARY: The U.S. Department of Health and Human Services (HHS) announces the first in a series of three federal advisory committee meetings on the Physical Activity Guidelines for Americans, to be held in Washington, DC. These meeting will be open to the public. The Physical Activity Guidelines Advisory Committee will review existing scientific literature to identify where there is sufficient evidence to develop a comprehensive set of specific physical activity recommendations. The Committee will prepare a report to the Secretary of HHS that documents the scientific background and rationale for the issuance of Physical Activity Guidelines for Americans. The report will also identify areas where further scientific research is needed. HHS will use the Final Report of the Committee to develop Physical Activity Guidelines. The intent is to issue physical activity recommendations for all Americans that will be tailored as necessary for specific subgroups of the population.

DATES: The Committee will meet on June 28–29, 2007 for a day and a half meeting.

ADDRESSES: The meeting will be held at the U.S. Department of Health and Human Services, Hubert H. Humphrey Building, located at 200 Independence Avenue, SW., Room 800, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: CAPT Richard Troiano, Ph.D., Executive Secretary, Physical Activity Guidelines Advisory Committee, Department of Health and Human Services, Office of Public Health and Science, Office of Disease Prevention and Health Promotion, Room LL–100, 1101 Wootton Parkway, Rockville, MD 20852, 240/453–8280 (telephone), 240/453–8281 (fax). Additional information is available on the Internet at *http://www.health.gov/PAguidelines.*

SUPPLEMENTARY INFORMATION: The Physical Activity Guidelines Advisory Committee: The thirteen-member Committee is chaired by William Haskell, Ph.D., Professor of Medicine, Stanford University School of Medicine. The Vice-Chair is Miriam Nelson, Ph.D., Director, John Hancock Center for Physical Activity and Nutrition, Friedman School of Nutrition Science and Policy, Tufts University. Other members of the Committee include Rod K. Dishman, Ph.D., Professor of Exercise Science and Director, Exercise Psychology Laboratory, Department of Kinesiology, University of Georgia; Edward Howley, Ph.D., Professor Emeritus, Department of Exercise, Sport, and Leisure Studies, University of Tennessee; Wendy Kohrt, Ph.D., Professor of Medicine, Division of Geriatric Medicine, University of Colorado at Denver and Health Sciences Center; William Kraus, M.D., Professor, Division of Cardiovascular Medicine, Duke University School of Medicine; I-Min Lee, M.D., Sc.D., Associate Professor of Medicine, Harvard Medical School and Associate Professor of Epidemiology, Harvard School of Public Health; Anne McTiernan, M.D., Ph.D., Director, Prevention Center, Fred Hutchinson Cancer Research Center; Russell Pate, Ph.D., Associate Vice President for Health Sciences, Office of Research and Health Sciences and Professor, Department of Exercise Science, University of South Carolina; Kenneth Powell, M.D., M.P.H., Public Health and Epidemiologic Consultant; Judith Regensteiner, Ph.D., Professor Department of Medicine and Director, Center for Women's Health Research, University of Colorado at Denver and Health Sciences Center: James Rimmer, Ph.D., Professor and Director, National Center on Physical Activity and Disability, Department of Disability and Human Development, University of Illinois at Chicago; and Antronette Yancey, M.D., M.P.H., Professor, Department of Health Services, University of California at Los Angeles School of Public Health.

Purpose of Meeting: Over the past 40 years, many organizations, including the Federal Government, have issued physical activity recommendations. While the various recommendations illustrate scientific consensus on the health benefits of physical activity, they differ from each other in the particular recommendations and highlighted benefits. The Physical Activity Guidelines Advisory Committee will review existing scientific literature to identify where there is sufficient evidence to develop a comprehensive

set of specific physical activity recommendations. The Committee will prepare a report to the Secretary of HHS that documents the scientific background and rationale for the issuance of Physical Activity Guidelines for Americans. The report will also identify areas where further scientific research is needed. HHS will use the Final Report of the Committee to develop Physical Activity Guidelines. The intent is to develop physical activity recommendations for all Americans that will be tailored as necessary for specific subgroups of the population.

Public Participation at Meeting: Members of the public are invited to observe the Advisory Committee meeting. Please note it is anticipated that there will be no oral public comments during the initial Physical Activity Guidelines Advisory Committee meeting, however, written comments are welcome throughout the Guidelines development process and may be e-mailed to *PA.guidelines@hhs.gov.* A summary of the Advisory Committee meetings will be made available shortly after each meeting.

To observe the Committee meeting, individuals must pre-register at the Physical Activity Guidelines Web site at *http://www.health.gov/PAguidelines.* Registrations must be completed by June 22, 2007. Space for the meeting is limited. Registrations will be accepted until maximum room capacity is reached. A waiting list will be maintained should registrations exceed room capacity. Individuals on the waiting list will be contacted as additional space for the meeting becomes available.

Registrants for the Physical Activity Advisory Committee meeting must present valid government-issued photo identification (i.e., driver's license) and should arrive 45 minutes prior to the start of the meeting to pass through security.

Registration questions may be directed to Experient at *PAguidelines@experient-inc.com* (e-mail), (703) 525–8333 x3349 (phone) or (703) 525–8557 (fax).

Dated: May 22, 2007.

Penelope Slade Royall,

RADM, USPHS, Deputy Assistant Secretary for Health, Office of Disease Prevention and Health Promotion.

[FR Doc. E7–10440 Filed 5–31–07; 8:45 am]

BILLING CODE 4150–32–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0197]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Registration of Food Facilities Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration is announcing that a collection of information entitled ''Registration of Food Facilities Under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of December 15, 2006 (71 FR 75555), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency 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. OMB has now approved the information collection and has assigned

OMB control number 0910–0502. The approval expires on May 31, 2010. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets.*

Dated: May 29, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–10617 Filed 5–31–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007E–0046]

Determination of Regulatory Review Period for Purposes of Patent Extension; ZILMAX

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for ZILMAX and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of Patents and Trademarks, Department of Commerce, for the extension of a patent which claims that animal drug product.

ADDRESSES: Submit written comments and petitions to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments.*

FOR FURTHER INFORMATION CONTACT: Beverly Friedman, Office of Regulatory Policy (HFD–007), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION: The Drug Price Competition and Patent Term Restoration Act of 1984 (Public Law 98–417) and the Generic Animal Drug and Patent Term Restoration Act (Public Law 100–670) generally provide that a patent may be extended for a period of up to 5 years so long as the patented item (human drug product, animal drug product, medical device, food additive, or color additive) was subject to regulatory review by FDA before the item was marketed. Under these acts, a product's regulatory review period forms the basis for determining the amount of extension an applicant may receive.

A regulatory review period consists of two periods of time: A testing phase and an approval phase. For animal drug products, the testing phase begins on the earlier date when either a major environmental effects test was initiated for the drug or when an exemption under section 512(j) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360b(j)) became effective and runs until the approval phase begins. The approval phase starts with the initial submission of an application to market the animal drug product and