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