

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](mailto: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](mailto: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

continues until FDA grants permission to market the drug product. Although only a portion of a regulatory review period may count toward the actual amount of extension that the Director of Patents and Trademarks may award (for example, half the testing phase must be subtracted as well as any time that may have occurred before the patent was issued), FDA's determination of the length of a regulatory review period for an animal drug product will include all of the testing phase and approval phase as specified in 35 U.S.C. 156(g)(4)(B).

FDA recently approved for marketing the animal drug product ZILMAX (zilpaterol hydrochloride). ZILMAX is indicated for increased rate of weight gain, improved feed efficiency, and increased carcass leanness in cattle fed in confinement for slaughter during the last 20 to 40 days on feed. Subsequent to this approval, the Patent and Trademark Office received a patent term restoration application for ZILMAX (U.S. Patent No. 4,900,735) from Intervet International B.V., and the Patent and Trademark Office requested FDA's assistance in determining this patent's eligibility for patent term restoration. In a letter dated March 5, 2007, FDA advised the Patent and Trademark Office that this animal drug product had undergone a regulatory review period and that the approval of ZILMAX represented the first permitted commercial marketing or use of the product. Shortly thereafter, the Patent and Trademark Office requested that FDA determine the product's regulatory review period.

FDA has determined that the applicable regulatory review period for ZILMAX is 4,720 days. Of this time, 4,674 days occurred during the testing phase of the regulatory review period, while 46 days occurred during the approval phase. These periods of time were derived from the following dates:

1. *The date an exemption under section 512(j) of the act involving this animal drug product became effective:* September 9, 1993. FDA has verified the applicant's claim that the date the investigational new animal drug application (INAD) became effective was on September 9, 1993.

2. *The date the application was initially submitted with respect to the animal drug product under section 512(b) of the act:* June 26, 2006. FDA has verified the applicant's claim that the new animal drug application (NADA) for ZILMAX (NADA 141-258) was initially submitted on June 26, 2006.

3. *The date the application was approved:* August 10, 2006. FDA has verified the applicant's claim that

NADA 141-258 was approved on August 10, 2006.

This determination of the regulatory review period establishes the maximum potential length of a patent extension. However, the U.S. Patent and Trademark Office applies several statutory limitations in its calculations of the actual period for patent extension. In its application for patent extension, this applicant seeks 5 years of patent term extension.

Anyone with knowledge that any of the dates as published is incorrect may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments and ask for a redetermination by July 31, 2007. Furthermore, any interested person may petition FDA for a determination regarding whether the applicant for extension acted with due diligence during the regulatory review period by November 28, 2007. To meet its burden, the petition must contain sufficient facts to merit an FDA investigation. (See H. Rept. 857, part 1, 98th Cong., 2d sess., pp. 41-42, 1984.) Petitions should be in the format specified in 21 CFR 10.30.

Comments and petitions should be submitted to the Division of Dockets Management. Three copies of any mailed information are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Comments and petitions may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: May 7, 2007.

**Jane A. Axelrad,**

*Associate Director for Policy, Center for Drug Evaluation and Research.*

[FR Doc. E7-10602 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-0011]

#### Determination of Regulatory Review Period for Purposes of Patent Extension; LANTUS

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) has determined the regulatory review period for LANTUS 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 human 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 human drug products, the testing phase begins when the exemption to permit the clinical investigations of the human drug product becomes effective and runs until the approval phase begins. The approval phase starts with the initial submission of an application to market the human drug product and continues until FDA grants permission to market the drug product. Although only a portion of a regulatory review period may count toward the actual amount of extension that the Director of Patents and Trademarks may award (for example, half the testing phase must be subtracted as well as any time that may have occurred before the patent was issued), FDA's determination of the length of a regulatory review period for a human drug product will include all of the testing phase and approval phase as specified in 35 U.S.C. 156(g)(1)(B).

FDA approved for marketing the human drug product LANTUS (insulin glargine [rDNA origin]). LANTUS is indicated for once-daily subcutaneous administration for the treatment of adult and pediatric patients with type 1 diabetes mellitus or adult patients with type 2 diabetes mellitus who require basal (long-acting) insulin for the