

The scope of the 2007 draft document is solely fisheries research to support the Magnuson-Stevens Act. It does not include the regulatory and enforcement components of NMFS' mission. NMFS currently conducts a comprehensive program of fisheries research and involves industry and others interested in planning and implementing its fisheries objectives.

NMFS intends that the final version of the Strategic Plan for Fisheries Research will take advantage of information and recommendations from all interested parties. Therefore, comments and suggestions on this draft NMFS Strategic Plan for Fisheries Research are hereby solicited from the public, other concerned government agencies, the scientific community, industry, and any other interested parties.

Dated: January 18, 2007.

**Steven A. Murawski,**

*Director of Scientific Programs and Chief Science Advisor, National Marine Fisheries Service.*

[FR Doc. E7-1017 Filed 1-23-07; 8:45 am]

**BILLING CODE 3510-22-S**

## DEPARTMENT OF COMMERCE

### National Oceanic and Atmospheric Administration

[I.D. 011107G]

#### Endangered Species; File No. 1596

**AGENCY:** National Marine Fisheries Service (NMFS), National Oceanic and Atmospheric Administration (NOAA), Commerce.

**ACTION:** Notice; issuance of permit.

**SUMMARY:** Notice is hereby given that NMFS Southwest Fisheries Science Center, 8604 La Jolla Shores Drive, La Jolla, CA 92037-1508 has been issued a permit to take leatherback (*Dermochelys coriacea*) sea turtles for purposes of scientific research.

**ADDRESSES:** The permit and related documents are available for review upon written request or by appointment in the following office(s):

Permits, Conservation and Education Division, Office of Protected Resources, NMFS, 1315 East-West Highway, Room 13705, Silver Spring, MD 20910; phone (301) 713-2289; fax (301) 427-2521;

Southwest Region, NMFS, 501 West Ocean Blvd., Suite 4200, Long Beach, CA 90802-4213; phone (562) 980-4001; fax (562) 980-4018.

**FOR FURTHER INFORMATION CONTACT:** Patrick Opay or Amy Hapeman, (301) 713-2289.

**SUPPLEMENTARY INFORMATION:** On October 20, 2006, notice was published

in the **Federal Register** (71 FR 61960) that a request for a scientific research permit to take leatherback sea turtles had been submitted by the above-named organization. The requested permit has been issued under the authority of the Endangered Species Act of 1973, as amended (ESA; 16 U.S.C. 1531 *et seq.*) and the regulations governing the taking, importing, and exporting of endangered and threatened species (50 CFR parts 222-226).

The researchers will continue long-term monitoring of the status of leatherback sea turtles off the coasts of California, Oregon, and Washington to determine their abundance, distribution, size ranges, sex ratio, health status, diving behavior, local movements, habitat use, and migration routes. Up to 38 animals will be captured using a breakaway hoop net and be measured, weighed, blood and tissue sampled, photographed, and flipper and passive integrated transponder (PIT) tagged. A subset of animals are to have biotelemetry devices (e.g., transmitters) attached to them. An additional 40 animals will be approached (but not captured) and have a VHF/TDR/sonic tag unit attached to them by suction cup using a long pole or these animals would be tissue sampled with a biopsy pole. The primary goal is to address priorities outlined in the U.S. Pacific leatherback Recovery Plan and identify critical forage habitats, genetic stock structure, migratory corridors, and potential fishery impacts on this species in the Pacific. This information is necessary to make informed management decisions concerning these turtles and their habitat. The permit is issued for 5 years.

Issuance of this permit, as required by the ESA, was based on a finding that such permit (1) was applied for in good faith, (2) will not operate to the disadvantage of any endangered or threatened species, and (3) is consistent with the purposes and policies set forth in section 2 of the ESA.

Dated: January 18, 2007.

**P. Michael Payne,**

*Chief, Permits, Conservation and Education Division, Office of Protected Resources, National Marine Fisheries Service.*

[FR Doc. E7-1014 Filed 1-23-07; 8:45 am]

**BILLING CODE 3510-22-S**

## DEPARTMENT OF COMMERCE

### Patent and Trademark Office

[Docket No. PTO-P-2006-0050]

#### Grant of Interim Extension of the Term of U.S. Patent No. 4,650,787; Sanvar®

**AGENCY:** United States Patent and Trademark Office, Commerce.

**ACTION:** Notice of Interim Patent Term Extension.

**SUMMARY:** The United States Patent and Trademark Office has issued a certificate under 35 U.S.C. 156(d)(5) for a second one-year interim extension of the term of U.S. Patent No. 4,650,787.

#### FOR FURTHER INFORMATION CONTACT:

Mary C. Till by telephone at (571) 272-7755; by mail marked to her attention and addressed to the Commissioner for Patents, Mail Stop Hatch-Waxman PTE., P.O. Box 1450, Alexandria, VA 22313-1450; by fax marked to her attention at (571) 273-7755, or by e-mail to [Mary.Till@uspto.gov](mailto:Mary.Till@uspto.gov).

**SUPPLEMENTARY INFORMATION:** Section 156 of Title 35, United States Code, generally provides that the term of a patent may be extended for a period of up to five years if the patent claims a product, or a method of making or using a product, that has been subject to certain defined regulatory review, and that the patent may be extended for interim periods of up to a year if the regulatory review is anticipated to extend beyond the expiration date of the patent.

On March 23, 2006, Debiovision Inc., the exclusive agent of Debiopharm S.A. and Debio Recherche Pharmaceutique S.A., who is the exclusive licensee of the Administrators of the Tulane Educational Fund of New Orleans, Louisiana, the patent owner, timely filed an application under 35 U.S.C. 156(d)(5) for a second interim extension of the term of U.S. Patent No. 4,650,787. The patent claims the human drug product Sanvar® (vapreotide acetate). The application indicates that a New Drug Application for the human drug product Sanvar® (vapreotide acetate) has been filed and is currently undergoing regulatory review before the Food and Drug Administration for permission to market or use the product commercially.

Review of the application indicates that except for permission to market or use the product commercially, the subject patent would be eligible for an extension of the patent term under 35 U.S.C. 156, and that the patent should be extended for one year as required by 35 U.S.C. 156(d)(5)(B). Because it is