and reissue patents as appropriate. Under 37 CFR 1.510–1.570 and 37 CFR 1.902–1.997, the USPTO may grant requests for *ex parte* and *inter partes* reexamination proceedings. The public uses this collection to request corrections of errors in issued patents, to request reissue patents, to request reexamination proceedings, and to ensure that the necessary fees and documentation are submitted to the USPTO. The USPTO is adding two items to this information collection, an electronic version of the Issue Fee Transmittal (Form PTOL–85B) and a petition to request an extension of time in *ex parte* or *inter partes* reexamination proceedings. This petition is an existing requirement that was not previously covered in this collection.

Affected Public: Individuals or households, businesses or other for-profits, and not-for-profit institutions.

Frequency: On occasion.

Respondent's Obligation: Required to obtain or retain benefits.

OMB Desk Officer: David Rostker, (202) 395–3897.

Copies of the above information collection proposal can be obtained by any of the following methods:

• *E-mail: Susan.Fawcett@uspto.gov.* Include ''0651–0033 copy request'' in the subject line of the message.

• *Fax:* 571–273–0112, marked to the attention of Susan Fawcett.

• *Mail:* Susan K. Fawcett, Records Officer, Office of the Chief Information Officer, Customer Information Services Group, Public Information Services Division, U.S. Patent and Trademark Office, P.O. Box 1450, Alexandria, VA 22313–1450.

Written comments and recommendations for the proposed information collection should be sent on or before May 10, 2007 to David Rostker, OMB Desk Officer, Room 10202, New Executive Office Building, 725 17th Street, NW., Washington, DC 20503.

Dated: April 3, 2007.

Susan K. Fawcett,

Records Officer, USPTO, Office of the Chief Information Officer, Customer Information Services Group, Public Information Services Division.

[FR Doc. E7–6735 Filed 4–9–07; 8:45 am]

BILLING CODE 3510–16–P

DEPARTMENT OF COMMERCE

Patent and Trademark Office

[Docket No. PTO–P–2007–0014]

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 third 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.*

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, 2007, 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 third 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 an additional one year as required by 35 U.S.C. 156(d)(5)(B). Because it is apparent that the regulatory review period will continue beyond the extended expiration date of the patent (April 25, 2007), a third interim extension of the patent term under 35 U.S.C. 156(d)(5) is appropriate.

A third interim extension under 35 U.S.C. 156(d)(5) of the term of U.S. Patent No. 4,650,787 is granted for a period of one year from the extended expiration date of the patent, i.e., until April 25, 2008.

Dated: April 3, 2007.

Jon W. Dudas,

Under Secretary of Commerce for Intellectual Property and Director of the United States Patent and Trademark Office.

[FR Doc. E7–6698 Filed 4–9–07; 8:45 am]

BILLING CODE 3510–16–P

COMMISSION OF FINE ARTS

Notice of Meeting

The next meeting of the U.S. Commission of Fine Arts is scheduled for 19 April 2007, at 10 a.m. in the Commission's offices at the National Building Museum, Suite 312, Judiciary Square, 401 F Street, NW., Washington, DC 20001–2728. Items of discussion affecting the appearance of Washington, DC, may include buildings, parks and memorials.

Draft agendas and additional information regarding the Commission are available on our Web site: *http://www.cfa.gov.* Inquiries regarding the agenda and requests to submit written or oral statements should be addressed to Thomas Luebke, Secretary, U.S. Commission of Fine Arts, at the above address or call 202–504–2200. Individuals requiring sign language interpretation for the hearing impaired should contact the Secretary at least 10 days before the meeting date.

Dated in Washington, DC, 4 April 2007.

Thomas Luebke,

Secretary

[FR Doc. 07–1772 Filed 4–9–07; 8:45 am]

BILLING CODE 6330–01–M

DEPARTMENT OF DEFENSE

Office of the Secretary

[No. DoD–2007–HA–0022]

Proposed Collection; Comment Request

AGENCY: Office of the Assistant Secretary of Defense for Health Affairs, DoD.

ACTION: Notice.

In accordance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Assistant Secretary of Defense for Health Affairs announces the proposed extension of a public information collection and seeks public comment on the provisions thereof. Comments are invited on: (a) Whether the proposed extension of collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the information collection; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology.

DATES: Consideration will be given to all comments received June 11, 2007.

ADDRESSES: You may submit comments, identified by docket number and title, by any of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Federal Docket Management System Office, 1160 Defense Pentagon, Washington, DC 20301–1160.

Instructions: All submissions received must include the agency name, docket number and title for this **Federal Register** document. The general policy for comments and other submissions from members of the public is to make these submissions available for public viewing on the Internet at *http://www.regulations.gov* as they are received without change, including any personal identifiers or contact information.

FOR FURTHER INFORMATION CONTACT: To request more information on this proposed information collection, please write TRICARE Management Activity, Office of General Counsel, 16401 E. Centretech Parkway, *Attn:* Helen Hilton, Aurora, CO 80011, or call TRICARE Management Activity, Office of General Counsel, at (303) 676–3542.

Title Associated With Form, and OMB Number: Statement of Personal Injury—Possible Third Party Liability, TRICARE Management Activity; DD Form 2527; OMB Number 0720–0003.

Needs and Uses: This information collection is completed by CHAMPUS beneficiaries suffering from personal injuries and receiving medical care at Government expense. The information is necessary in the assertion of the Government's right to recovery under the Federal Medical Care Recovery Act. The data is used in the evaluation and processing of these claims.

Affected Public: Individuals or households; Federal government.

Annual Burden Hours: 33,250.

Number of Respondents: 133,000.

Responses per Respondent: 1.

Average Burden per Response: 20 minutes.

Frequency: On occasion, only when a beneficiary is injured under circumstances creating possible liability in a third party.

SUPPLEMENTARY INFORMATION:

Summary of Information Collection

The Federal Medical Recovery Act, 42 U.S.C. 2651–2653 as implemented by Executive Order No. 11060 and 28 CFR part 43 provides for recovery of the reasonable value of medical care provided by the United States to a person who is injured or suffers a disease under circumstances creating tort liability in some third person. DD Form 2527 is required for investigating and asserting claims in favor of the United States arising out of such incidents.

When a claim for CHAMPUS benefits is identified as involving possible third party liability and the information is not submitted with the claim, the TRICARE/CHAMPUS contractor requests that the injured party (or a designee) complete DD Form 2527. To protect the interests of the Government, the contractor suspends claims processing until the requested third party liability information is received. The contractor conducts a preliminary evaluation based upon the collection of information and refers the case to a designated appropriate legal officer of the Uniformed Services. The responsible Uniformed Services legal officer uses the information as a basis for asserting and settling the Government's claim. When appropriate, the information is forwarded to the Department of Justice as the basis for litigation.

Section 1 of the Form is used to collect general information, such as name, address and telephone numbers about the military sponsor and the injured beneficiary and the date, time and location where the injured occurred.

Section 2 of the Form is used to collect information about accidental injuries. Most of the investigations for third party liability involve motor vehicle accidents. Information about insurance coverage for the parties involved in the accident is collected. Section 2 of the Form is also used to collect information about accidents that do not involve motor vehicles. Information such as the type of accident, the place where the injury occurred, the name of the property owner where the injury occurred and cause of the injury is collected. The name and address of the employer is collected when the injury was work related.

Section 3 of the Form is used for miscellaneous information such as possible medical treatment at a Government hospital, the name and address of the beneficiary's attorney, and information regarding any possible releases or settlements with another party to the accident. It also contains the certification, date and signature of the beneficiary (or a designee).

Dated: April 3, 2007.

Patricia L. Toppings,

Alternate OSD Federal Register Liaison Officer, Department of Defense.

[FR Doc. 07–1757 Filed 4–9–07; 8:45 am]

BILLING CODE 5001–06–M

DEPARTMENT OF DEFENSE

Office of the Secretary

[No. DoD–2007–HA–0029]

Proposed Collection; Comment Request

AGENCY: Office of the Assistant Secretary of Defense for Health Affairs, DoD.

ACTION: Notice.

In accordance with Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Assistant Secretary of Defense for Health Affairs announces the proposed extension of a public information collection and seeks public comment on the provisions thereof. Comments are invited on: (a) Whether the proposed extension of collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the information collection; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the information collection on respondents, including through the use of automated collection techniques or other forms of information technology.

DATES: Consideration will be given to all comments received June 11, 2007.

ADDRESSES: You may submit comments, identified by docket number and title, by any of the following methods: