

National Institutes of Health/NIAID, 6700B Rockledge Drive, MSC 7616, Bethesda, MD 20892-7616. 301-496-2550. [qvos@niaid.nih.gov](mailto:qvos@niaid.nih.gov).

(Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: November 17, 2010.

**Jennifer S. Spaeth,**

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010-29525 Filed 11-22-10; 8:45 am]

**BILLING CODE 4140-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Heart, Lung, and Blood Institute; Notice of Closed Meetings**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* National Heart, Lung, and Blood Institute Special Emphasis Panel Institutional National Research Service Awards.

*Date:* December 15, 2010.

*Time:* 1 p.m. to 4:30 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20817. (Virtual Meeting).

*Contact Person:* Roy L White, PhD, Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, 6701 Rockledge Drive, Room 7176, Bethesda, MD 20892-7924. 301-435-0310. [whiterl@nhlbi.nih.gov](mailto:whiterl@nhlbi.nih.gov).

*Name of Committee:* National Heart, Lung, and Blood Institute Special Emphasis Panel Conference Grants.

*Date:* December 16-17, 2010.

*Time:* 8 a.m. to 5 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892. (Virtual Meeting).

*Contact Person:* William J Johnson, PhD, Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and

Blood Institute, 6701 Rockledge Drive, Room 7178, Bethesda, MD 20892-7924. 301-435-0725. [johnsonwj@nhlbi.nih.gov](mailto:johnsonwj@nhlbi.nih.gov).

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: November 17, 2010.

**Jennifer S. Spaeth,**

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010-29524 Filed 11-22-10; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2010-N-0001]

**Oncologic Drugs Advisory Committee; Amendment of Notice**

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an amendment to the notice of a meeting of the Oncologic Drugs Advisory Committee. This meeting was announced in the **Federal Register** of October 19, 2010 (75 FR 64314). The amendment is being made to reflect changes in the *Date and Time*, *Agenda*, and *Procedure* portions of the document. We also are postponing a session regarding biologics license application (BLA) 125377, with the proposed trade name Yervoy (ipilimumab), manufactured by Bristol-Myers Squibb Co. The proposed indication (use) for this product is for the treatment of advanced melanoma in patients who have received prior therapy. This portion of the meeting has been postponed due to the need to complete the review of additional data submitted by the applicant. Future meeting dates may be announced in the **Federal Register**.

**FOR FURTHER INFORMATION CONTACT:**

Nicole Vesely, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993-0002, 301-796-9001, FAX: 301-847-8533, e-mail: [Nicole.vesely@fda.hhs.gov](mailto:Nicole.vesely@fda.hhs.gov), or FDA Advisory Committee Information Line, 1-800-741-8138, 301-443-0572 in the Washington, DC area, code 301-451-2542. Please call the Information Line

for up-to-date information on this meeting.

**SUPPLEMENTARY INFORMATION:** In the **Federal Register** of October 19, 2010 (75 FR 64314), FDA announced that a meeting of the Oncologic Drugs Advisory Committee would be held on December 2, 2010. On page 64314, in the first column, the *Date and Time* portion of the document is changed to read as follows:

*Date and Time:* The meeting will be held on December 2, 2010, from 8 a.m. to 12:30 p.m.

On page 64314, in the second column, the *Agenda* portion of the document is changed to read as follows:

*Agenda:* On December 2, 2010, the committee will discuss new drug application (NDA) 022-405, with the proposed trade name Zictifa (vandetanib) Tablets, manufactured by iPR Pharmaceuticals, Inc., represented by AstraZeneca Pharmaceuticals LP (authorized U.S. agent). The proposed indication (use) for this product is for the treatment of patients with unresectable (non-operable) locally advanced or metastatic medullary thyroid cancer.

On page 64314, in the second column, the third sentence in the *Procedure* portion of the document is changed to read as follows:

*Procedure:* Oral presentations from the public will be scheduled between approximately 10:30 a.m. and 11:30 a.m.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to the advisory committees.

Dated: November 17, 2010.

**Joanne Less,**

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2010-29522 Filed 11-22-10; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Heart, Lung, and Blood Institute; Notice of Meeting**

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of a meeting of the Sleep Disorders Research Advisory Board.

The meeting will be open to the public. Individuals who plan to attend and need special assistance should notify the Contact Person listed below in advance of the meeting.

*Name of Committee:* Sleep Disorders Research Advisory Board.

Date: January 4, 2011.

Time: 1 p.m. to 5 p.m.

Agenda: To discuss sleep research plan development. Public meeting observers should call 1-888-791-5525 to access the teleconference and the observer passcode is 3737665. Public meeting observers should send comments or questions for this meeting to the National Center on Sleep Disorders Research by e-mail [mt2d@nih.gov](mailto:mt2d@nih.gov) or fax 301-480-3557.

Place: National Institutes of Health, 6701 Rockledge Drive, Suite 10170, Bethesda, MD 20892. (Telephone Conference Call)

Contact Person: Michael J. Twery, PhD, Director, National Center on Sleep Disorders Research, Division of Lung Diseases, National Heart, Lung, and Blood Institute, National Institutes of Health, 6701 Rockledge Drive, Suite 10038, Bethesda, MD 20892-7952. 301-435-0199. [twerym@nhlbi.nih.gov](mailto:twerym@nhlbi.nih.gov).

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

Information is also available on the Institute's/Center's home page: [www.nhlbi.nih.gov/meetings/index.htm](http://www.nhlbi.nih.gov/meetings/index.htm), where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: November 17, 2010.

**Jennifer S. Spaeth,**

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2010-29523 Filed 11-22-10; 8:45 am]

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## DEPARTMENT OF HOMELAND SECURITY

### U.S. Citizenship and Immigration Services

#### Agency Information Collection Activities: Form I-130, Extension of a Currently Approved Information Collection; Comment Request

**ACTION:** 30-Day Notice of Information Collection Under Review: Form I-130, Petition for Alien Relative; OMB Control No. 1615-0012.

The Department of Homeland Security, U.S. Citizenship and Immigration Services (USCIS) will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of

1995. The information collection was previously published in the **Federal Register** on August 26, 2010, at 75 FR 52540, allowing for a 60-day public comment period. USCIS did not receive any comments for this information collection.

The purpose of this notice is to allow an additional 30 days for public comments. Comments are encouraged and will be accepted until December 23, 2010. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Department of Homeland Security (DHS), and to the Office of Management and Budget (OMB) USCIS Desk Officer. Comments may be submitted to: USCIS, Chief, Regulatory Products Division, 20 Massachusetts Avenue, Washington, DC 20529-2020. Comments may also be submitted to DHS via facsimile to 202-272-8352 or via e-mail at [rfs.regs@dhs.gov](mailto:rfs.regs@dhs.gov), and to the OMB USCIS Desk Officer via facsimile at 202-395-5806 or via e-mail at [oir\\_submission@omb.eop.gov](mailto:oir_submission@omb.eop.gov). When submitting comments by e-mail please make sure to add OMB Control Number 1615-0012 in the subject box. Written comments and suggestions from the public and affected agencies should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

#### Overview of This Information Collection

(1) *Type of Information Collection:* Extension of a currently approved information collection.

(2) *Title of the Form/Collection:* Petition for Alien Relative.

(3) *Agency form number, if any, and the applicable component of the Department of Homeland Security sponsoring the collection:* Form I-130; U.S. Citizenship and Immigration Services (USCIS).

(4) Affected public who will be asked or required to respond, as well as a brief abstract: *Primary: Individuals or Households.* This form allows citizens or lawful permanent residents of the United States to petition on behalf of certain alien relatives who wish to immigrate to the United States.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* 690,520 responses at 1.5 hours per response.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 1,035,780 annual burden hours.

If you need a copy of the information collection instrument, please visit the Web site at: <http://www.regulations.gov>.

We may also be contacted at: USCIS, Regulatory Products Division, 20 Massachusetts Avenue, NW., Washington, DC 20529-2020; Telephone 202-272-8377.

Dated: November 18, 2010.

**Stephen Tarragon,**

Deputy Chief, Regulatory Products Division, U.S. Citizenship and Immigration Services, Department of Homeland Security.

[FR Doc. 2010-29521 Filed 11-22-10; 8:45 am]

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## DEPARTMENT OF HOMELAND SECURITY

### U.S. Citizenship and Immigration Services

#### Agency Information Collection Activities: Form N-470, Extension of a Currently Approved Information Collection; Comment Request

**ACTION:** 30-Day Notice of Information Collection Under Review: Form N-470, Application to Preserve Residence for Naturalization; OMB Control No. 1615-0056.

The Department of Homeland Security, U.S. Citizenship and Immigration Services (USCIS) will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995. The information collection was previously published in the **Federal Register** on August 18, 2010, at 75 FR 51096, allowing for a 60-day public