

C. The Nature and Extent of Voluntary Steps To Mitigate the Impact on the Public

Next, I consider whether Liu has raised specific facts showing that there is a genuine and substantial issue of fact that requires a hearing concerning the nature and extent of voluntary steps to mitigate the impact of his offense on the public, including full cooperation with any investigations (including the extent of disclosure to appropriate authorities of all wrongdoing) and any other actions taken to substantially limit potential or actual adverse effects on the public health.

In the Notice of Opportunity for a Hearing, ORA stated, "You took no steps to mitigate the impact on the public of your actions." Liu has not challenged this statement. As such, I agree with ORA that the nature and extent of Liu's voluntary steps to mitigate the impact to the public weighs in favor of debarment.

D. The Impact of Changes in Ownership, Management, or Operations

In the Notice of Opportunity for Hearing, ORA determined that this factor was not applicable for consideration. Liu has not challenged that determination.

E. Prior Convictions Under the FD&C Act or Related Acts

In the Notice of Opportunity for Hearing, ORA acknowledged that the Agency was unaware of any prior convictions involving matters within the jurisdiction of FDA. The lack of previous violations of the FD&C Act or related statutes by Liu weighs against debarment.

III. Findings and Order

The Director of the Office of Scientific Integrity, under section 306(b)(3)(A) of the FD&C Act and under authority delegated to him, finds that Liu has been convicted of a felony for conduct relating to the importation of food into the United States. Accordingly, FDA may debar Liu from importing articles of food or offering such articles for import into the United States for a period of not more than 5 years.

I have considered the arguments raised by Liu regarding the relevant factors listed in section 306(c)(3) of the FD&C Act and have determined that Liu has raised no genuine and substantial issues of fact that require resolution at an evidentiary hearing. I have considered the factors in section 306(c)(3) of the FD&C Act. The nature and seriousness of Liu's offense, Liu's management participation in the offense, and the lack of any voluntary

steps to mitigate the impact of the offense weigh in favor of debarment. Although Liu appears to have no prior criminal convictions involving matters within the jurisdiction of FDA, that consideration does not counterbalance to a sufficient degree the remaining considerations to warrant decreasing the period of debarment. Of particular note is the nature and seriousness of the offense, in light of the volume of honey that was imported, the amount of duties that were avoided, and the fact that false statements were made with regard to two shipments of honey. I agree with ORA's proposed period of debarment and find that a debarment of 5 years is appropriate.

As a result of the foregoing findings, Liu is debarred for a period of 5 years from importing articles of food or offering such articles for import into the United States, effective (see DATES). Under section 301(cc) of the FD&C Act, the importing or offering for import into the United States of an article of food by, with the assistance of, or at the direction of Liu is a prohibited act.

Any application by Liu for termination of debarment under section 306(d) of the FD&C Act should be identified with Docket No. FDA-2011-N-0169 and sent to the Division of Dockets Management Branch (see ADDRESSES). All such submissions are to be filed in four copies. The public availability of information in these submissions is governed by 21 CFR 10.20(j). Publicly available submissions may be seen in the Division of Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday. Persons with access to the Internet may obtain documents in the Docket at <http://www.regulations.gov>.

Dated: June 25, 2015.

Nathan Doty,

Director, Office of Scientific Integrity.

[FR Doc. 2015-16561 Filed 7-6-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Advisory Council on Nurse Education and Practice; Notice of Meeting

SUMMARY: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), notice is hereby given of the following meeting of the National Advisory Council on Nurse Education and Practice (NACNEP).

DATES: July 28 and 29, 2015, 8:30 a.m.–5 p.m. EST.

ADDRESSES: This meeting will be via Webinar Format. U.S. Department of Health and Human Services, Health Resources and Services Administration, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: For additional information regarding NACNEP, please contact Jeanne Brown, Staff Assistant, National Advisory Council on Nurse Education and Practice, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. The telephone number is: (301) 443-5688. The email is jbrown@hrsa.gov.

SUPPLEMENTARY INFORMATION:

Status: This advisory council meeting will be open to the public.

Purpose: The purpose of the 131st National Advisory Council on Nurse Education and Practice (NACNEP) meeting is to provide advice and recommendations on policy and program development related to the role of nursing in Interprofessional Education (IPE) and Practice. The purpose is to discuss existing IPE models in an academic setting and the intersect between education and practice as it relates to Health Care delivery reform. The goal of the meeting is to solicit recommendations for IPE in an academic setting and the intersect important to IPE and practice. Strengths, challenges, achievable solutions, and replicable models required and/or available to move from discussion to action will be identified. Additionally, the meeting will discuss topics for future work of the council. This meeting will conclude with a formulation of recommendations and form the basis for NACNEP's mandated Thirteenth Annual Report to the Secretary of the U.S. Department of Health and Human Services and Congress.

Agenda: A final agenda will be posted on the *NACNEP Web site* 3 days prior to the meeting. Agenda items are subject to change as priorities dictate.

Further information regarding NACNEP including the roster of members, reports to Congress, and minutes from previous meetings is available at the NACNEP Web site. Members of the public and interested parties may request to participate in the meeting by contacting Staff Assistant, Jeanne Brown. Access to the meeting will be granted on a first come, first-served basis and space is limited. Public participants may submit written statements in advance of the scheduled meeting. If you would like to provide oral public comment during the meeting

you will need to register with Erin Fowler, Designated Federal Official (DFO). Public comment will be limited to 3 minutes per speaker and is tentatively scheduled for after lunch on the first day of the meeting. Statements and comments can be addressed to Erin Fowler. Please send by email to: nacnep@hrsa.gov.

Please be advised that committee members are given copies of all written statements submitted by the public prior to the meeting. Any further public participation will be at the discretion of the Chair, with approval of the DFO in attendance. Registration through the designated contact for the public comment session is required. Any member of the public who wishes to have printed materials distributed to the Advisory Group should submit materials to the point of contact no later than 12:00 p.m. EST on July 13, 2015.

Jackie Painter,

Director, Division of the Executive Secretariat.

[FR Doc. 2015-16135 Filed 7-6-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Council on Blood Stem Cell Transplantation; Notice of Meeting

SUMMARY: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), notice is hereby given of the following meeting of the Advisory Council on Blood Stem Cell Transplantation (ACBSCT).

DATES: September 11, 2015 from 8:00 a.m. to 4:30 p.m. Eastern Time.

ADDRESSES: Health Resources and Services Administration, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Patricia Stroup, MBA, MPA, Executive Secretary, Healthcare Systems Bureau, Health Resources and Services Administration, 5600 Fishers Lane, Room 17W65, Rockville, Maryland 20857; telephone (301) 443-1127.

SUPPLEMENTARY INFORMATION:

Status: The meeting will be open to the public.

Purpose: Pursuant to Public Law 109-129, 42 U.S.C. 274k (section 379 of the Public Health Service Act, as amended), the ACBSCT advises the Secretary of the Department of Health and Human Services and the Administrator, Health Resources and Services Administration, on matters related to the activities of the C.W. Bill Young Cell Transplantation

Program (Program) and the National Cord Blood Inventory Program.

Agenda: The Council will hear a report from the ACBSCT Work Group on Advancing Hematopoietic Stem Cell Transplantation for Hemoglobinopathies. The Council also will hear presentations and discussions on topics including: past recommendations made to the Secretary, cord blood bank economics, status of Food and Drug Administration licensure of cord blood banks, and potential impact of haploidentical transplants. Agenda items are subject to changes as priorities indicate.

After Council discussions, members of the public will have an opportunity to provide comment. Because of the Council's full agenda and timeframe in which to cover the agenda topics, public comment may be limited. All public comments will be included in the record of the ACBSCT meeting. Meeting summary notes will be posted on HRSA's Program Web site at http://bloodcell.transplant.hrsa.gov/ABOUT/Advisory_Council/index.html.

The draft meeting agenda will be posted on <https://www.blsmmeetings.net/acbsct>. Those participating at this meeting should pre-register by visiting <https://www.blsmmeetings.net/acbsct>. The deadline to pre-register for this meeting is Thursday, September 10, 2015. Registration will be confirmed on site. For all logistical questions and concerns, please contact Anita Allen, Seamon Corporation, at (301) 658-3442 or send an email to aallen@seamoncorporation.com.

Participants can also join this meeting via teleconference by:

1. (Audio Portion) Calling the Conference Phone Number (800-988-0202) and providing the Participant Passcode (9115853); and
2. (Visual Portion) Connecting to the ACBSCT Adobe Connect Pro Meeting using the following URL and entering as GUEST: https://hrsa.connectsolutions.com/acbsct_webinar/ (copy and paste the link into your browser if it does not work directly, and enter as a guest).

Participants should call and connect 15 minutes prior to the meeting for logistics to be set up. If you have never attended an Adobe Connect meeting, please test your connection using the following URL: https://hrsa.connectsolutions.com/common/help/en/support/meeting_test.htm. In order to obtain a quick overview, go to the following URL: http://www.adobe.com/go/connectpro_overview. Call (301) 443-0437 or send an email to ptonge@hrsa.gov if you are

having trouble connecting to the meeting site.

Public Comment: It is preferred that persons interested in providing an oral presentation email a written request, along with a copy of their presentation, to Patricia Stroup, MBA, MPA, Executive Secretary, Healthcare Systems Bureau, Health Resources and Services Administration, at pstroup@hrsa.gov. Requests should contain the name, address, telephone number, email address, and any business or professional affiliation of the person desiring to make an oral presentation. Groups having similar interests are requested to combine their comments and present them through a single representative.

The allocation of time may be adjusted to accommodate the level of expressed interest. Persons who do not file an advance request for a presentation, but desire to make an oral statement, may request it during the public comment period. Public participation and ability to comment will be limited as time permits.

Jackie Painter,

Director, Division of the Executive Secretariat.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Vaccine Injury Compensation Program; List of Petitions Received

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: The Health Resources and Services Administration (HRSA) is publishing this notice of petitions received under the National Vaccine Injury Compensation Program (the Program), as required by Section 2112(b)(2) of the Public Health Service (PHS) Act, as amended. While the Secretary of Health and Human Services is named as the respondent in all proceedings brought by the filing of petitions for compensation under the Program, the United States Court of Federal Claims is charged by statute with responsibility for considering and acting upon the petitions.

FOR FURTHER INFORMATION CONTACT: For information about requirements for filing petitions, and the Program in general, contact the Clerk, United States Court of Federal Claims, 717 Madison Place NW., Washington, DC 20005,