

Chester, OH 45069. Officers: John J. Plnes, CEO, (Qualifying Individual).
Guaranteed International Freight and Trade, Inc., dba International Freight and Trading, 239–241 Kingston Ave., Brooklyn, NY 11213. Officers: Lawrence Medas, Sr., President, (Qualifying Individual), Cornelius Medas, CEO.

Florida Freight Forwarders, LLC, 2041 NW 12th Ave., Miami, FL 33127.

Officer: Jose Maria Rivas, Vice President, (Qualifying Individual).

Sunset Transportation, Inc., 11406

Gravois Road, St. Louis, MO 63126.

Officers: Deborah L. Kopeny, Director, (Qualifying Individual), James A. Williams.

D.M.C. Logistics Incorporated, 207

Meadow Road, Edison, NJ 08817.

Officers: Julia Ertler, Vice President, (Qualifying Individual), Francis S. Molfetta, President.

Cargo Shipping Expedition International

Inc., 6 Sandow Court, Fair Lawn, NJ

07410. Officers: Gerry Lysogorsky,

Vice President, (Qualifying

Individual), Alexander Zilberman, President.

Ocean Freight Forwarder—Ocean Transportation Intermediary Applicants

Hal-Mari International Logistics, Inc., 935 Knotty Elmwood Trail, Houston, TX 77062. Officer: Ilkka Halmari,

President, (Qualifying Individual).

Matt Global Freight Co. LLC, 3517

Langrehr Road, Suite 102, Baltimore, MD 21244. Officers: Mathew T.

Chacko, President, (Qualifying

Individual), Ann T. Mathews, Vice

President.

Allfreight Worldwide Cargo, Inc., dba

Allfreight, 4810 Beauregard Street,

Suite 100, Alexandria, VA 22312.

Officers: Demeke Meri, CEO/

President, (Qualifying Individual),

Abel Meri, Director.

Dated: March 3, 2006.

Karen V. Gregory,

Assistant Secretary.

[FR Doc. E6–3315 Filed 3–8–06; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Sunshine Act; Notice

TIME AND DATE: 9 a.m. (est); March 20, 2006.

PLACE: 4th Floor Conference Room, 1250 H Street, NW., Washington, DC.

STATUS: Open.

MATTERS TO BE CONSIDERED:

1. Approval of the minutes of the February 21, 2006, Board Member meeting.

2. Thrift Savings Plan activity report by the Executive Director.

CONTACT PERSON FOR MORE INFORMATION:

Thomas J. Trabucco, Director, Office of External Affairs, (202) 942–1640.

Dated: March 6, 2006.

Thomas K. Emswiler,

Acting General Counsel, Federal Retirement Thrift Investment Board.

[FR Doc. 06–2325 Filed 3–7–06; 1:01 pm]

BILLING CODE 6760–01–P

GOVERNMENT ACCOUNTABILITY OFFICE

Publication of Volume II of GAO's Principles of Federal Appropriations

AGENCY: Government Accountability Office.

ACTION: Notice of publication.

SUMMARY: The third edition of Volume II of GAO's *Principles of Federal Appropriations Law* is being prepared for publication by the Government Printing Office (GPO). Government departments, agencies, and other federal organizations that normally require more than one copy have been given an opportunity to request them through their agencies' account representatives at pre-publication rate. This notice is intended for other parties who might be interested in purchasing the book.

SUPPLEMENTARY INFORMATION: The Government Accountability Office (GAO) will shortly publish Volume II of *Principles of Federal Appropriations Law*, third edition—also known as “The Red Book” This publication is part of a multi-volume set intended to present a basic reference covering those areas of law in which the Comptroller General renders decisions. Our approach is to lay a foundation with text discussion, using specific legal authorities to illustrate the principles discussed, their application, and exceptions. These authorities include GAO decisions and opinions, judicial decisions, statutory provisions, and other relevant sources.

GAO will provide copies of this volume to the heads of Federal agencies, and agencies have already been given an opportunity to place advance (rider) orders for additional copies of this volume with their account representatives at the Government Printing Office (GPO).

This notice is intended to tell the general public that they will be able to place pre-issue orders for this publication through GPO's new online

bookstore, at http://bookstore.gpo.gov/collections/gao_appropriation.jsp.

Otherwise, we expect this publication will be available for purchase from the Superintendent of Documents, United States Government Printing Office, P.O. Box 371954, Pittsburgh, PA 15250–7954, by April 2006. The price is \$69.

Orders for Volume II should specify GPO Stock No. 020–000–00287–5 or the ISBN 0–16–0075602–2. Through periodic training courses on federal appropriations law, GAO believes that this publication might be useful in particular to law offices, to accounting firms, to the financial, budget, or accounting officers of government contractors, to university and state law libraries, to corporate chief financial officers, and to people who follow Federal financial management, contracts, grants, and loans.

As with the second edition of *Principles*, we are publishing the third edition in loose-leaf format but will include a CD-ROM as well. Volume II covers chapters 6 through 11: *availability of appropriations; amount; obligation of appropriations; continuing resolutions; liability and relief of accountable officers; Federal assistance, grants and cooperative agreements; and Federal assistance, guaranteed and insured loans*. We plan three volumes with annual updates. The updates will only be published electronically. Users should retain copies of the remaining volumes of the second edition until each volume is revised. Volume III of the second edition addresses provisions that the GAO Act of 1996 transferred to the Executive Branch and will not be updated. The first annual update of Volume I is currently available online at <http://www.gao.gov/legal.htm>.

Authority: 31 U.S.C. 712, 717, 719, 3511, 3526–29.

Susan Poling,

Managing Associate General Counsel, Government Accountability Office.

[FR Doc. 06–2235 Filed 3–8–06; 8:45 am]

BILLING CODE 1610–02–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Toxicology Program (NTP); Liaison and Scientific Review Office; Meeting of the NTP Board of Scientific Counselors Technical Reports Review Subcommittee

AGENCY: National Institute of Environmental Health Sciences (NIEHS), National Institutes of Health (NIH).

ACTION: Meeting announcement and request for comments.

SUMMARY: Pursuant to Public Law 92-463, notice is hereby given of a meeting of the NTP Board of Scientific Counselors Technical Reports Review Subcommittee (TRR Subcommittee). The primary agenda topic is the peer review of the findings and conclusions presented in four draft NTP Technical Reports of rodent toxicology and carcinogenicity studies conducted by the NTP (see Preliminary Agenda below). The TRR Subcommittee meeting is open to the public with time scheduled for oral public comment. The NTP also invites written comments on any draft technical report discussed at the meeting. The TRR Subcommittee deliberations on the draft technical reports will be reported to the NTP Board of Scientific Counselors (NTP Board) at a future date.

DATES: The TRR Subcommittee meeting will be held on June 12, 2006. All individuals who plan to attend are encouraged to register online by May 30, 2006, at the NTP Web site ([http://ntp.niehs.nih.gov/select "Advisory Boards & Committees"](http://ntp.niehs.nih.gov/select/AdvisoryBoards&Committees)). In order to facilitate planning for this meeting, persons wishing to make an oral presentation are asked to notify Dr. Barbara Shane via online registration, phone, or e-mail (see **ADDRESSES** below) by May 30, 2006, and if possible, to send a copy of the statement or talking points at that time. Written comments on the draft reports are also welcome and should also be received by May 30, 2006, to enable review by the TRR Subcommittee and NIEHS/NTP staff prior to the meeting. Persons needing special assistance, such as sign language interpretation or other reasonable accommodation in order to attend, should contact 919-541-2475 (voice), 919-541-4644 TTY (text telephone), through the Federal TTY Relay System at 800-877-8339, or by e-mail to niehsoeeo@niehs.nih.gov. Requests should be made at least 7 days in advance of the event.

ADDRESSES: The TRR Subcommittee meeting will be held in the Rodbell Auditorium, Rall Building at the NIEHS, 111 T. W. Alexander Drive, Research Triangle Park, NC 27709. A copy of the preliminary agenda, committee roster, and any additional information, when available, will be posted on the NTP Web site ([http://ntp.niehs.nih.gov/select "Advisory Boards and Committees"](http://ntp.niehs.nih.gov/select/AdvisoryBoards&Committees)) and provided upon request. Public comments and any other correspondence should be submitted to Dr. Barbara Shane, Executive Secretary

for the NTP Board (NTP Liaison and Scientific Review Office, NIEHS, P.O. Box 12233, MD A3-01, Research Triangle Park, NC 27709; telephone: 919-541-4253, fax: 919-541-0295; or e-mail: shane@niehs.nih.gov).

SUPPLEMENTARY INFORMATION:

Background

The primary agenda topic is the peer review of the findings and conclusions of four draft NTP Technical Reports of rodent toxicology and carcinogenicity studies conducted by the NTP (see Preliminary Agenda below) on studies with conventional strains of rats and mice.

Attendance and Registration

The meeting is scheduled for June 12, 2006, from 8:30 a.m. to adjournment and is open to the public with attendance limited only by the space available. Individuals who plan to attend are encouraged to register online at the NTP Web site by May 30, 2006, at [http://ntp.niehs.nih.gov/select "Advisory Boards and Committees"](http://ntp.niehs.nih.gov/select/AdvisoryBoards&Committees) to facilitate access to the NIEHS campus. Please note that a photo ID is required to access the NIEHS campus. The NTP is making plans to videocast the meeting through the Internet at <http://www.niehs.nih.gov/external/video.htm>.

Availability of Meeting Materials

A copy of the preliminary agenda, committee roster, and any additional information, when available, will be posted on the NTP Web site ([http://ntp.niehs.nih.gov/select "Advisory Boards and Committees"](http://ntp.niehs.nih.gov/select/AdvisoryBoards&Committees)) or may be requested in hardcopy from the NTP (see **ADDRESSES** above). Following the meeting, summary minutes will be prepared and made available on the NTP Web site.

Request for Comments

Public input at this meeting is invited and time is set aside for the presentation of public comments on any draft technical report. Each organization is allowed one time slot per agenda topic. At least 7 minutes will be allotted to each speaker, and if time permits, may be extended to 10 minutes. Registration for oral comments will also be available on-site, although time allowed for presentation by on-site registrants may be less than that for pre-registered speakers and will be determined by the number of persons who register at the meeting.

Persons registering to make oral comments are asked, if possible, to send a copy of their statement to Dr. Shane (see **ADDRESSES** above) by May 30, 2006, to enable review by the TRR

Subcommittee and NIEHS/NTP staff prior to the meeting. Written statements can supplement and may expand the oral presentation. If registering on-site and reading from written text, please bring 40 copies of the statement for distribution to the TRR Subcommittee and NIEHS/NTP staff and to supplement the record. Written comments received in response to this notice will be posted on the NTP Web site. Persons submitting written comments should include their name, affiliation, mailing address, phone, fax, e-mail, and sponsoring organization (if any) with the document.

Background Information on the NTP Board of Scientific Counselors

The NTP Board is a technical advisory body comprised of scientists from the public and private sectors who provide primary scientific oversight to the overall program and its centers. Specifically, the NTP Board advises the NTP on matters of scientific program content, both present and future, and conducts periodic review of the program for the purposes of determining and advising on the scientific merit of its activities and their overall scientific quality. The TRR Subcommittee is a standing subcommittee of the NTP Board. NTP Board members are selected from recognized authorities knowledgeable in fields, such as toxicology, pharmacology, pathology, biochemistry, epidemiology, risk assessment, carcinogenesis, mutagenesis, molecular biology, behavioral toxicology and neurotoxicology, immunotoxicology, reproductive toxicology or teratology, and biostatistics. Its members are invited to serve overlapping terms of up to four years. NTP Board meetings are held annually or biannually.

Dated: February 27, 2006.

Samuel H. Wilson,

Deputy Director, National Institute of Environmental Health Sciences and the National Toxicology Program.

Preliminary Agenda

National Toxicology Program (NTP) Board of Scientific Counselors Technical Reports Review Subcommittee Meeting

June 12, 2006

Rodbell Auditorium, Rall Building, National Institute of Environmental Health Sciences, 111 TW Alexander Drive, Research Triangle Park, NC

NTP Technical Reports (TR) Scheduled for Review

- TR 539 Genistein (CASRN 446-72-0)—Multigenerational Study.

- Naturally occurring phytoestrogen, found in soy products.
- TR 545: Genistein (CASNR 446-72-0)—2-year Bioassay. Naturally occurring phytoestrogen, found in soy products.
 - TR 543: α -Methylstyrene (CASNR 98-83-9).
 - Used in the production of resins and polymers.
 - TR 540: Methylene blue trihydrate (CAS No. 7220-79-3).
 - Dye used to stain tissues and bacteriological samples for microscopy and an antidote for methemoglobinemia; previously used as a hair colorant.

[FR Doc. E6-3317 Filed 3-8-06; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel: National Institute for Occupational Safety and Health (NIOSH) Support for Conferences and Scientific Meetings, Request for Applications PAR 06-014

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces the following meeting:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): NIOSH Support for Conferences and Scientific Meetings, Request for Applications PAR 06-014.

Time and Date: 1 p.m.–4 p.m., March 29, 2006 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92-463.

Matters to Be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to NIOSH Support for Conferences and Scientific Meetings, Request for Applications PAR 06-014.

For More Information Contact: George Bockosh, MS, Scientific Review Administrator, National Institute for Occupational Safety and Health, CDC, 626 Cochran Mill Road, MS PO-5, Pittsburgh, PA 15236, Telephone 412-386-6465. The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the CDC and the Agency for Toxic Substances and Disease Registry.

Dated: March 3, 2006.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. E6-3354 Filed 3-8-06; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Injury Prevention and Control Initial Review Group

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announce the following meeting:

Name: National Center for Injury Prevention and Control (NCIPC) Initial Review Group (IRG).

Times and Dates: 6:30 p.m.–10 p.m., April 10, 2006. 8:30 a.m.–6 p.m., April 11, 2006. 8 a.m.–5:30 p.m., April 12, 2006.

Place: Hilton Atlanta Airport and Towers, 1031 Virginia Avenue, Atlanta, Georgia 30354.

Status: Open: 6:30 p.m.–7:15 p.m., April 10, 2006.

Closed: 7:15 p.m. to 10 p.m., April 10, 2006. 8:30 a.m. to 6 p.m., April 11, 2006. 8 a.m. to 5:30 p.m., April 12, 2006.

Purpose: This group is charged with providing advice and guidance to the Secretary, Department of Health and Human Services (HHS) and the Director, CDC, concerning the scientific and technical merit of grant and cooperative agreement applications received from academic institutions and other public and private profit and nonprofit organizations, including state and local government agencies, to conduct specific injury research that focuses on injury prevention and control.

Matters to be Discussed: Agenda items include an overview of the injury program, discussion of the review process and panelists responsibilities, and the review of, and vote on, applications. Beginning at 7:15 p.m., April 10, through 5:30 p.m., April 12, the Group will review individual research grant and cooperative agreement applications submitted in response to six Fiscal Year 2006 Requests for Applications (RFAs) related to the following individual research announcements: #06001, Research Grants to Prevent Unintentional Injuries; #06002, Dissertation Grant Awards for Violence-Related Injury Prevention Research in Minority Communities; #06003, Research Grants to Describe Traumatic Brain Injury Consequences; #06004, Grants for Violence-Related Injury Prevention Research: Youth Violence, Suicidal Behavior, Child Maltreatment, Intimate Partner Violence, and Sexual Violence; #06005, Research Grants for the Care of the Acutely Injured; #06007, Evaluation of Community-based Approaches to Increasing Seat Belt Use Among Adolescent Drivers and Their Passengers.

This portion of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c)(4) and (6), Title 5, U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to section 10(d) of Public Law 92-463.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Gwendolyn H. Cattledge, Ph.D., M.S.E.H., Executive Secretary, NCIPC IRG, CDC, 4770 Buford Highway, NE, M/S K02, Atlanta, Georgia 30341-3724, telephone 770/488-4655.

The Director, Management Analysis and Services Office has been delegated the authority to sign **Federal Register** Notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: March 3, 2006.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. E6-3346 Filed 3-8-06; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Institute for Occupational Safety and Health Advisory Board on Radiation and Worker Health

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces the following committee meeting:

Name: Advisory Board on Radiation and Worker Health (ABRWH), National Institute for Occupational Safety and Health (NIOSH).

Time and Date: 10 a.m.–4 p.m., EST, Tuesday, March 14, 2006.

Place: Audio Conference Call via FTS Conferencing. The USA toll free dial in number is 1-866-643-6504, pass code of 9448550.

Status: Open to the public, but without a public comment period.

Background: The ABRWH was established under the Energy Employees Occupational Illness Compensation Program Act (EEOICPA) of 2000 to advise the President, delegated to the Secretary of Health and Human Services (HHS), on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Board include providing advice on the development of probability of causation guidelines which have been promulgated by HHS as a final rule, advice on methods of dose reconstruction which have also been promulgated by HHS as a final rule, advice on the scientific validity and quality of dose