individuals are exposed to insecticides, recommending individuals to be observed up to 96 hours after treatment, and revising the statistical analyses recommendations.

Authority: 7 U.S.C. 136 *et seq.;* 15 U.S.C. 2601 *et seq.;* 21 U.S.C. 301 *et seq.*

Dated: May 16, 2017.

Wendy C. Hamnett,

Acting Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

[FR Doc. 2017–12347 Filed 6–13–17; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–9963–68–ORD]

Human Studies Review Board; Notification of Public Meetings

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA), Office of the Science Advisor announces two separate public meetings of the Human Studies Review Board (HSRB) to advise the Agency on the ethical and scientific review of research involving human subjects.

DATES: A virtual public meeting will be held on Wednesday, July 26, 2017, from 1:00 p.m. to approximately 5:00 p.m. Eastern Time. A separate, subsequent teleconference meeting is planned for Friday, September 15, 2017, from 2:00 p.m. to approximately 3:30 p.m. Eastern Time for the HSRB to finalize its Final Report of the July 26, 2017 meeting and review other possible topics.

ADDRESSES: Both of these meetings will be conducted entirely by telephone and on the Internet using Adobe Connect. For detailed access information visit the HSRB Web site: *https://www2.epa.gov/osa/human-studies-review-board*

FOR FURTHER INFORMATION CONTACT: Any member of the public who wishes to receive further information should contact the HSRB Designated Federal Official (DFO), Jim Downing on telephone number (202) 564–2468; fax number: (202) 564–2070; email address: *downing.jim@epa.gov;* or mailing address: Environmental Protection Agency, Office of the Science Advisor, Mail code 8105R, 1200 Pennsylvania Avenue NW., Washington, DC 20460.

SUPPLEMENTARY INFORMATION:

Meeting access: These meetings are open to the public. The full Agenda and meeting materials are available at the HSRB Web site: *https://www2.epa.gov/osa/human-studies-review-board* . For questions on document availability, or if you do not have access to the Internet, consult with the DFO, Jim Downing listed under **FOR FURTHER INFORMATION CONTACT**.

Special accommodations. For information on access or services for individuals with disabilities, or to request accommodation of a disability, please contact the DFO listed under **FOR FURTHER INFORMATION CONTACT** at least 10 days prior to the meeting to give EPA as much time as possible to process your request.

How may I participate in this meeting?

The HSRB encourages the public's input. You may participate in these meetings by following the instructions in this section.

1. Oral comments. Requests to present oral comments during either meeting will be accepted up to Noon Eastern Time on Wednesday, July 19, 2017, for the July 26, 2017 meeting and up to Noon Eastern Time on Friday, September 8, 2017 for the September 15, 2017 teleconference. To the extent that time permits, interested persons who have not pre-registered may be permitted by the HSRB Chair to present oral comments during either meeting at the designated time on the agenda. Oral comments before the HSRB are generally limited to five minutes per individual or organization. If additional time is available, further public comments may be possible.

2. Written comments. Submit your written comments prior to the meetings. For the Board to have the best opportunity to review and consider your comments as it deliberates, you should submit your comments by Noon Eastern Time on Wednesday, July 19 2016, for the July 26, 2017 meeting, and by noon Eastern Time on Friday, September 8, 2017 for the September 15, 2017 teleconference. If you submit comments after these dates, those comments will be provided to the HSRB members, but you should recognize that the HSRB members may not have adequate time to consider your comments prior to their discussion. You should submit your comments to the DFO, Jim Downing listed under **FOR FURTHER INFORMATION CONTACT**. There is no limit on the length of written comments for consideration by the HSRB.

Background

The HSRB is a Federal advisory committee operating in accordance with the Federal Advisory Committee Act 5 U.S.C. App.2 section 9. The HSRB provides advice, information, and recommendations on issues related to scientific and ethical aspects of third-party human subjects research that are submitted to the Office of Pesticide Programs (OPP) to be used for regulatory purposes.

Topic for discussion. On Wednesday, July 26, 2017, EPA's Human Studies Review Board will consider one topic: Field evaluation of three topically applied insect repellent products containing IR3535 against mosquitoes in Florida.

The Agenda and meeting materials for this topic will be available in advance of the meeting at *https://www2.epa.gov/osa/human-studies-review-board.*

On September 15, 2017, the Human Studies Review Board will review and finalize their draft Final Report from the July 26, 2017 meeting, in addition to other topics that may come before the Board. The HSRB may also discuss planning for future HSRB meetings. The agenda and the draft report will be available prior to the teleconference at *https://www2.epa.gov/osa/human-studies-review-board.*

Meeting minutes and final reports. Minutes of these meetings, summarizing the matters discussed and recommendations made by the HSRB, will be released within 90 calendar days of the meeting. These minutes will be available at *https://www2.epa.gov/osa/human-studies-review-board.* In addition, information regarding the HSRB's Final Report, will be found at *https://www2.epa.gov/osa/human-studies-review-board* or from Jim Downing listed under **FOR FURTHER INFORMATION CONTACT**.

Robert J. Kavlock,

EPA Science Advisor.

[FR Doc. 2017–12345 Filed 6–13–17; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL MARITIME COMMISSION

[Docket No. 17–06]

Notice of Filing of Complaint and Assignment: Tarik Afif Chaouch v. Demetrios Air Freight Co., Demetrios International Shipping Co., Inc., and Troy Container Line Ltd

Notice is given that a complaint has been filed with the Federal Maritime Commission (Commission) by Tarik Afif Chaouch, hereinafter ''Complainant,'' against Demetrios Air Freight Co., Demetrios International Shipping Co., Inc., and Troy Container Line LTD, hereinafter ''Respondents.'' Complainant states it hired the Respondents to ship two cars to Algiers, Algeria.

Complainant alleges that due to an error the Respondents made on the bill

of lading, the shipment was ''impounded in Algiers, Algeria for approximately four months.'' Complainant alleges that this error resulted in costs for which complainant would not have otherwise been responsible. Complainant alleges that it is ''subject to injury as a result of the violations by respondent of sections 46 U.S.C. code § 41104 and more specifically paragraphs 4 and 5.''

Complainant seeks reparations in the amount of $21,086.70, and other relief. The full text of the complaint can be found in the Commission's Electronic Reading Room at *www.fmc.gov/17-06/.*

This proceeding has been assigned to the Office of Administrative Law Judges. The initial decision of the presiding officer in this proceeding shall be issued by June 8, 2018, and the final decision of the Commission shall be issued by December 21, 2018.

Rachel E. Dickon,

Assistant Secretary.

[FR Doc. 2017–12296 Filed 6–13–17; 8:45 am]

BILLING CODE 6731–AA–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–17–1015]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) 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; (b) 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; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) 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; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

The National Electronic Health Records Survey (NEHRS) (OMB Control No. 0920–1015, Expires 04/30/2017)—Reinstatement with Change—National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Section 306 of the Public Health Service (PHS) Act (42 U.S.C. 242k), as amended, authorizes that the Secretary of Health and Human Services (DHHS), acting through NCHS, shall collect statistics on ''utilization of health care'' in the United States. NEHRS was originally designed as a mail supplement to the National Ambulatory Medical Care Survey (NAMCS). Questions in NEHRS have been asked in NAMCS starting in 2001.

The purpose of NEHRS is to measure progress toward goals for electronic health records (EHRs) adoption. NEHRS target universe consists of all non-Federal office-based physicians (excluding those in the specialties of anesthesiology, radiology, and pathology) who are engaged in direct patient care.

NEHRS is the principal source of data on national and state-level EHR adoption in the United States. In 2008 and 2009, the sample size was 2,000 physicians annually. Starting in 2010, the annual sample size was increased five-fold, from 2,000 physicians to 10,302 physicians. The increased sample size allows for more reliable national estimates as well as state-level estimates on EHR adoption without having to be combined with NAMCS. For these reasons, in 2012 NEHRS became an independent survey, not as a supplement under NAMCS.

NEHRS collects information on characteristics of physician practices, the capabilities of EHRs in those practices, and intent to apply for meaningful use incentive payments. These data, together with trend data, may be used to monitor the adoption of EHR as well as accessing factors associated with EHR adoption. In 2017, a set of follow-up questionnaires will be incorporated into the survey that focuses on content related to physician attitudes on using EHRs.

Users of NEHRS data include, but are not limited to, Congressional offices, Federal agencies, state and local governments, schools of public health, colleges and universities, private industry, nonprofit foundations, professional associations, clinicians, researchers, administrators, and health planners. There is no cost to the respondents other than their time. The total estimated annualized burden hours are 6,295.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (hours)
Office-based physicians	NEHRS	10,302	1	30/60
Office-based physicians	Follow-up NEHRS Elec Resp	858	1	20/60
Office-based physicians	Follow-up NEHRS Non-Elec Resp	859	1	20/60
Office-based physicians	Follow-up NEHRS Nonresp	1,717	1	20/60