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


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

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, 2017, 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.

FEDERAL MARITIME COMMISSION

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

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,
the NAMCS became an independent 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

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