Parties: ZIM Integrated Shipping Services, Ltd. and Turkon Container Transportation & Shipping Inc.

Filing Party: Mark E. Newcomb; ZIM American Integrated Shipping Services Co., LLC; 5801 Lake Wright Dr.; Norfolk, VA 23508.

Synopsis: The agreement authorizes Turkon to charter space to ZIM in the trade between Greece and the U.S. East Coast. The parties have requested expedited review.

By Order of the Federal Maritime Commission.

Dated: March 21, 2014.

Karen V. Gregory,

Secretary.

[FR Doc. 2014–06711 Filed 3–26–14; 8:45 am]

BILLING CODE 6730–01–P

FEDERAL MINE SAFETY AND HEALTH REVIEW COMMISSION

Sunshine Act Notice

March 24, 2014.

TIME AND DATE: 10:00 a.m., Thursday, April 10, 2014.

PLACE: The Richard V. Backley Hearing Room, Room 511N, 1331 Pennsylvania Avenue NW., Washington, DC 20004 (entry from F Street entrance).

STATUS: Open.

MATTERS TO BE CONSIDERED: The Commission will consider and act upon the following in open session: *Secretary of Labor* v. *Wolf Run Mining Co.,* Docket Nos. WEVA 2006–853, *et al.* (Issues include whether the Administrative Law Judge erred in concluding that a violation of a lightning arrester standard was not ''significant and substantial.'')

Any person attending this meeting who requires special accessibility features and/or auxiliary aids, such as sign language interpreters, must inform the Commission in advance of those needs. Subject to 29 CFR 2706.150(a)(3) and 2706.160(d).

CONTACT PERSON FOR MORE INFO: Jean Ellen (202) 434–9950/(202) 708–9300 for TDD Relay/1–800–877–8339 for toll free.

Emogene Johnson,

Administrative Assistant.

[FR Doc. 2014–06904 Filed 3–25–14; 11:15 am]

BILLING CODE 6735–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: 201403–0990–004–30D]

Agency Information Collection Activities; Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, has submitted an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for a new generic clearance for information collection. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Comments on the ICR must be received on or before April 28, 2014.

ADDRESSES: Submit your comments to *OIRA_submission@omb.eop.gov* or via facsimile to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, *Information.CollectionClearance@hhs.gov* or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the Information Collection Request Title and document identifier 201403–0990–004–30D for reference.

Information Collection Request Title: ASPE Generic Clearance for the Collection of Qualitative Research and Assessment.

Abstract: The Office of the Assistant Secretary for Planning and Evaluation at the Department of Health and Human Services (HHS) is requesting a generic clearance for purposes of conducting qualitative research.

Need and Proposed Use of the Information: The information collected will be used to gain a better understanding of emerging health policy issues, develop future intramural and extramural research projects, and to ensure HHS leadership, agencies and offices have recent data and information to inform program and policy decision-making.

Likely Respondents: Policy experts, national, state, and local health representatives, healthcare providers, and representatives of other health organizations.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Health Policy Stakeholder	747	1	1	747

Keith A. Tucker,

Information Collection Clearance Officer.

[FR Doc. 2014–06765 Filed 3–26–14; 8:45 am]

BILLING CODE 4150–05–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0238]

Neurological Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Neurological Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on April 24, 2014, from 8 a.m. to 6 p.m.

Location: Holiday Inn, Main Ballroom, 2 Montgomery Village Ave., Gaithersburg, MD 20879. The hotel's telephone number is 301–948–8900.

Contact Person: Avena Russell, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1535, Silver Spring, MD 20993–0002, *Avena.Russell@fda.hhs.gov,* 301–796–3805, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site at *http://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

Agenda: On April 24, 2014, the committee will discuss the current knowledge about the safety and effectiveness of aversive conditioning devices that are intended to deliver a noxious electrical stimulus to a patient to modify undesirable behavioral characteristics. FDA is convening this committee to seek clinical and scientific expert opinion on the risks and benefits of certain aversive conditioning devices based on available scientific data and information. The Agency is considering whether to ban aversive conditioning devices that are intended to administer a noxious electrical stimulus to a patient to modify undesirable behavioral characteristics. The meeting will concern only devices classified under 21 CFR 882.5235 (aversive conditioning device, class II) that are not self-administered. Devices which deliver a noxious electrical stimulus automatically are not considered to be self-administered devices. Section 516 of the FD&C Act (21 U.S.C. 360f) sets forth the standard for banning devices. Under that provision, in order to ban a device, FDA must make a finding that a device ''presents substantial deception or an unreasonable and substantial risk of illness or injury'' based on all available data and information. FDA regulations provide additional details about the procedures and standards for banning a device (21 CFR part 895).

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

Procedure: FDA will work with affected industry, professional organizations, and societies that have an interest in aversive conditioning devices and who wish to make a presentation separate from the general open public hearing; time slots on April 24, 2014, between approximately 11 a.m. and 12 p.m. Representatives from industry, professional organizations and societies interested in making formal presentations to the committee should notify the contact person on or before March 28, 2014.

Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before April 14, 2014. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before April 4, 2014. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by April 7, 2014.

FDA is opening a docket for public comment on this document. The docket number is FDA–2014–N–0238. The docket will close on June 24, 2014. Interested persons are encouraged to use the docket to submit electronic or written comments regarding this meeting. Comments received on or before April 14, 2014, will be provided to the committee for their consideration. Comments received after May 27, 2014 will be taken into consideration by the Agency.

Interested persons may submit either electronic comments regarding this document to *http://www.regulations.gov* or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at *http://www.regulations.gov.*

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams at *Annmarie.Williams@fda.hhs.gov,* or 301–796–5966 at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.