

information from public review, we cannot guarantee that we will be able to do so.

Abstract: The regulations at 30 CFR part 250, subpart O, concern well control and production safety training and are the subject of this collection. This request also covers any related Notices to Lessees and Operators (NTLs) that BSEE issues to clarify, supplement, or provide additional guidance on some aspects of our regulations.

The BSEE will use the information collected under subpart O regulations to ensure that workers in the OCS are properly trained with the necessary skills to perform their jobs in a safe and pollution-free manner.

In some instances, we may conduct oral interviews of offshore employees to evaluate the effectiveness of a company's training program. The oral interviews are used to gauge how effectively the companies are implementing their own training program.

Title of Collection: 30 CFR 250, Subpart O, *Well Control and Production Safety Training*.

OMB Control Number: 1014-0008.

Form Number: None.

Type of Review: Extension of a currently approved collection.

Respondents/Affected Public: Potential respondents comprise Federal OCS oil, gas, and sulfur lessees/operators and holders of pipeline rights-of-way.

Total Estimated Number of Annual Respondents: Varies, not all of the potential respondents will submit information in any given year and some may submit multiple times.

Total Estimated Number of Annual Responses: 6.

Estimated Completion Time per Response: Varies from 1 hour to 105 hours, depending on activity.

Total Estimated Number of Annual Burden Hours: 202.

Respondent's Obligation: Most responses are mandatory, while others are required to obtain or retain benefits.

Frequency of Collection: On occasion.

Total Estimated Annual Nonhour Burden Cost: We have not identified any non-hour cost burdens associated with this collection of information.

An agency may not conduct or sponsor and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

The authority for this action is the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*)

Dated: March 5, 2018.

Doug Morris,

Chief, Office of Offshore Regulatory Programs.

[FR Doc. 2018-07976 Filed 4-16-18; 8:45 am]

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INTERNATIONAL TRADE COMMISSION

[Investigation No. U.S.-Korea FTA-103-031]

U.S.-Korea FTA: Advice on Modifications to Duty Rates for Certain Motor Vehicles

AGENCY: United States International Trade Commission.

ACTION: Institution of investigation and notice of opportunity to provide written comments.

SUMMARY: Following receipt on April 6, 2018, of a request from the U.S. Trade Representative (USTR), the Commission instituted investigation No. U.S.-Korea FTA-103-031, *U.S.-Korea FTA: Advice on Modifications to Duty Rates for Certain Motor Vehicles*, for the purpose of providing advice on the probable economic effect of modifications to the United States-Korea Free Trade Agreement regarding the staging of duty treatment for certain motor vehicles.

DATES: May 1, 2018: Deadline for filing written submissions.

June 1, 2018: Transmittal of Commission report to USTR.

ADDRESSES: All Commission offices, including the Commission's hearing rooms, are located in the United States International Trade Commission Building, 500 E Street SW, Washington, DC. All written submissions should be addressed to the Secretary, United States International Trade Commission, 500 E Street SW, Washington, DC 20436. The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at <http://www.usitc.gov/secretary/edis.htm>.

FOR FURTHER INFORMATION CONTACT: Project Leader Jeff Horowitz (202-205-2750 or jeffrey.horowitz@usitc.gov) or Deputy Project Leader Mitch Semanik (202-205-2034 or mitchell.semanik@usitc.gov) for information specific to this investigation. For information on the legal aspects of this investigation, contact William Gearhart of the Commission's Office of the General Counsel (202-205-3091 or william.gearhart@usitc.gov). The media should contact Margaret O'Laughlin, Office of External Relations (202-205-1819 or margaret.olaughlin@usitc.gov). Hearing-impaired individuals may

obtain information on this matter by contacting the Commission's TDD terminal at 202-205-1810. General information concerning the Commission may also be obtained by accessing its internet server (<http://www.usitc.gov>). Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202-205-2000.

Background: In his request letter (received April 6, 2018), the USTR stated that U.S. negotiators have recently reached an agreement in principle with representatives of the government of Korea on modifications to the FTA regarding the staging of duty treatment for certain motor vehicles. The USTR noted that section 201(b)(2) of the United States-Korea Free Trade Agreement Implementation Act (the Act) authorizes the President, subject to the consultation and layover requirements of section 104 of the Act, to proclaim such tariff modifications as the President determines to be necessary or appropriate to maintain the general level of reciprocal and mutually advantageous concessions with respect to Korea provided for by the FTA. He noted that one of the requirements set out in section 104 of the Act is that the President obtain advice regarding the proposed action from the U.S. International Trade Commission.

In the request letter, the USTR asked that the Commission provide advice on the probable economic effect of the modifications on U.S. trade under the FTA and on domestic producers of the affected articles. He asked that the Commission provide its advice at the earliest possible date but no later than eight weeks from receipt of the request. He also asked that the Commission issue, as soon as possible thereafter, a public version of its report with any confidential business information deleted.

The products identified in the proposal are motor vehicles for the transport of goods provided for in subheadings 8704.21.00, 8704.22.50, 8704.23.00, 8704.31.00, 8704.32.00, and 8704.90.00 of the U.S. Harmonized Tariff Schedule. The request letter and the proposed modification are available on the Commission's website at http://www.usitc.gov/research_and_analysis/what_we_are_working_on.htm. As requested, the Commission will provide its advice to USTR by June 1, 2018.

Written Submissions: No public hearing is planned. However, interested parties are invited to file written submissions. All written submissions should be addressed to the Secretary, and should be received no later than

5:15 p.m., May 1, 2018. All written submissions must conform with the provisions of section 201.8 of the Commission's *Rules of Practice and Procedure* (19 CFR 201.8). Section 201.8 and the Commission's Handbook on Filing Procedures require that interested parties file documents electronically on or before the filing deadline and submit eight (8) true paper copies by 12:00 p.m. eastern time on the next business day. In the event that confidential treatment of a document is requested, interested parties must file, at the same time as the eight paper copies, at least four (4) additional true paper copies in which the confidential information must be deleted (see the following paragraph for further information regarding confidential business information). Persons with questions regarding electronic filing should contact the Office of the Secretary, Docket Services Division (202-205-1802).

Confidential Business Information: Any submissions that contain confidential business information must also conform with the requirements of section 201.6 of the Commission's *Rules of Practice and Procedure* (19 CFR 201.6). Section 201.6 of the rules requires that the cover of the document and the individual pages be clearly marked as to whether they are the "confidential" or "non-confidential" version, and that the confidential business information is clearly identified by means of brackets. All written submissions, except for those containing CBI, will be made available for inspection by interested parties.

The Commission may include some or all of the confidential business information submitted in the course of this investigation in the report it sends to the USTR. Additionally, all information, including CBI, submitted in this investigation may be disclosed to and used: (i) By the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the

Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel for cybersecurity purposes. The Commission will not otherwise disclose any confidential business information in a manner that would reveal the operations of the firm supplying the information.

Summaries Of Written Submissions: The Commission intends to publish summaries of the positions of interested persons in an appendix to its report. Persons wishing to have a summary of their position included in the appendix should include a summary with their written submission. The summary may not exceed 500 words, should be in MSWord format or a format that can be easily converted to MSWord, and should not include any CBI. The summary will be included in the report as provided if it meets these requirements and is germane to the subject matter of the investigation. In the appendix, the Commission will identify the name of the organization furnishing the summary and will include a link to the Commission's Electronic Document Information System (EDIS) where the full written submission can be found.

By order of the Commission.

Issued: April 12, 2018.

Lisa Barton,

Secretary to the Commission.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances

Application: Clinical Supplies Management Holdings, Inc.

ACTION: Notice of application.

DATES: Registered bulk importers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the

proposed registration on or before May 17, 2018. Such persons may also file a written request for a hearing on the application on or before May 17, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All request for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417, (January 25, 2007)

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division ("Assistant Administrator") pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on March 14, 2018, Clinical Supplies Management Holdings, Inc., 342 42nd Street South, Fargo, ND 58103 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I

The company plans to import analytical reference standards for distribution to its customers for research and analytical purposes. Placement of these drug codes onto the company's registration does not translate into

automatic approval of subsequent permit applications to import controlled substances. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized

under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.