the Commission's TDD terminal on 202–205–1810. 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. General information concerning the Commission may also be obtained by accessing its internet server (http://www.usitc.gov).

SUPPLEMENTARY INFORMATION:

Background

The final phase of this investigation is being scheduled as a result of an affirmative preliminary determination by the Department of Commerce that imports of bulk acetylsalicylic acid (aspirin) from China are being sold in the United States at less than fair value within the meaning of section 733 of the Act (19 U.S.C. 1673b). The investigation was requested in a petition filed on May 28, 1999, by Rhodia, Inc., Cranbury, NJ.

Participation in the Investigation and Public Service List

Persons, including industrial users of the subject merchandise and, if the merchandise is sold at the retail level, representative consumer organizations, wishing to participate in the final phase of this investigation as parties must file an entry of appearance with the Secretary to the Commission, as provided in section 201.11 of the Commission's rules, no later than 21 days prior to the hearing date specified in this notice. A party that filed a notice of appearance during the preliminary phase of the investigation need not file an additional notice of appearance during this final phase. The Secretary will maintain a public service list containing the names and addresses of all persons, or their representatives, who are parties to the investigation.

Limited Disclosure of Business Proprietary Information (BPI) Under an Administrative Protective Order (APO) and BPI Service List

Pursuant to section 207.7(a) of the Commission's rules, the Secretary will make BPI gathered in the final phase of this investigation available to authorized applicants under the APO issued in the investigation, provided that the application is made no later than 21 days prior to the hearing date specified in this notice. Authorized applicants must represent interested parties, as defined by 19 U.S.C. § 1677(9), who are parties to the investigation. A party granted access to BPI in the preliminary phase of the investigation need not reapply for such access. A separate service list will be maintained by the Secretary for those parties authorized to receive BPI under the APO.

Staff Report

The prehearing staff report in the final phase of this investigation will be placed in the nonpublic record on May 5, 2000, and a public version will be issued thereafter, pursuant to section 207.22 of the Commission's rules.

Hearing

The Commission will hold a hearing in connection with the final phase of this investigation beginning at 9:30 a.m. on May 18, 2000, at the U.S. International Trade Commission Building. Requests to appear at the hearing should be filed in writing with the Secretary to the Commission on or before May 10, 2000. A nonparty who has testimony that may aid the Commission's deliberations may request permission to present a short statement at the hearing. All parties and nonparties desiring to appear at the hearing and make oral presentations should attend a prehearing conference to be held at 9:30 a.m. on May 15, 2000, at the U.S. International Trade Commission Building. Oral testimony and written materials to be submitted at the public hearing are governed by sections 201.6(b)(2), 201.13(f), and 207.24 of the Commission's rules. Parties must submit any request to present a portion of their hearing testimony *in camera* no later than 7 days prior to the date of the hearing.

Written Submissions

Each party who is an interested party shall submit a prehearing brief to the Commission. Prehearing briefs must conform with the provisions of section 207.23 of the Commission's rules; the deadline for filing is May 12, 2000. Parties may also file written testimony in connection with their presentation at the hearing, as provided in section 207.24 of the Commission's rules, and posthearing briefs, which must conform with the provisions of section 207.25 of the Commission's rules. The deadline for filing posthearing briefs is May 25, 2000; witness testimony must be filed no later than three days before the hearing. In addition, any person who has not entered an appearance as a party to the investigation may submit a written statement of information pertinent to the subject of the investigation on or before May 25, 2000. On June 15, 2000, the Commission will make available to parties all information on which they have not had an opportunity to comment. Parties may submit final comments on this information on or before June 19, 2000, but such final comments must not contain new factual information and must otherwise comply with section 207.30 of the Commission's rules. All written submissions must conform with the provisions of section 201.8 of the Commission's rules; any submissions that contain BPI must also conform with the requirements of sections 201.6, 207.3, and 207.7 of the Commission's rules. The Commission's rules do not authorize filing of submissions with the Secretary by facsimile or electronic means.

In accordance with sections 201.16(c) and 207.3 of the Commission's rules, each document filed by a party to the investigation must be served on all other parties to the investigation (as identified by either the public or BPI service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

Authority: This investigation is being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.21 of the Commission's rules.

Issued: February 1, 2000.

By order of the Commission.

Donna R. Koehnke,

Secretary.

[FR Doc. 00–2525 Filed 2–3–00; 8:45 am]

BILLING CODE 7020–02–U

INTERNATIONAL TRADE COMMISSION

[Investigation No. 731–TA–377 (Review)]

Internal Combustion Industrial Forklift Trucks From Japan

AGENCY: United States International Trade Commission.

ACTION: Cancellation of the hearing and revision of the schedule of a full five-year review concerning the antidumping duty order on internal combustion industrial forklift trucks from Japan.

EFFECTIVE DATE: January 28, 2000.

FOR FURTHER INFORMATION CONTACT: Christopher J. Cassise (202–708–5408), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired persons can obtain information on this matter by contacting the Commission's TDD terminal on 202–205–1810. 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. General information concerning the Commission may also be obtained by

accessing its internet server (*http://www.usitc.gov*).

SUPPLEMENTARY INFORMATION:

Background

On August 27, 1999 (64 FR 46952), the Commission published a notice in the **Federal Register** scheduling a full five-year review concerning the antidumping duty order on internal combustion industrial forklift trucks from Japan. The schedule provided for a public hearing on January 25, 2000. Requests to appear at the hearing were filed with the Commission on behalf of NACCO Materials Handling Group and on behalf of Clark Material Handling Co. However, the Federal Government was closed on January 25, 2000, because of snow and so the Commission hearing was not held as scheduled. Subsequently, each of the parties requesting to appear at the hearing withdrew their request. Since there are no current requests by interested parties to appear at a public hearing, the Commission determined to cancel, instead of reschedule, the public hearing on internal combustion industrial forklift trucks from Japan and provide those parties scheduled to appear an opportunity to present written testimony. The Commission unanimously determined that no earlier announcement of this cancellation was possible.

The Commission's new schedule for the review is as follows: the deadline for filing posthearing briefs is February 15, 2000; the Commission will make its final release of information on March 9, 2000; and final party comments are due on March 13, 2000.

For further information concerning the review, see the Commission's notice cited above and the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A and F (19 CFR part 207).

Authority: This review is being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to sections 201.35 and 207.62 of the Commission's rules.

Issued: January 31, 2000.

By order of the Commission.

Donna R. Koehnke,

Secretary.

[FR Doc. 00–2524 Filed 2–3–00; 8:45 am]

BILLING CODE 7020–02–U

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

James Garvey Cavanagh, M.D.; Revocation of Registration

On August 5, 1999, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA), issued an Order to Show Cause to James Garvey Cavanagh, M.D., of Hawthorne, Nevada, notifying him of an opportunity to show cause as to why DEA should not revoke his DEA Certificate of Registration AC9084485 pursuant to 21 U.S.C. 824(a)(3), and deny any pending applications for renewal of such registration pursuant to 21 U.S.C. 823(f), for reason that he is not currently authorized to handle controlled substances in the State of Nevada. The order also notified Dr. Cavanagh that should no request for a hearing be filed within 30 days, his hearing right would be deemed waived.

DEA received a signed receipt indicating that the Order to Show Cause was received on August 21, 1999. No request for a hearing or any other reply was received by the DEA from Dr. Cavanagh or anyone purporting to represent him in this matter. Therefore the Deputy Administrator, finding that (1) 30 days have passed since the receipt of the Order to Show Cause, and (2) no request for a hearing having been received, concludes that Dr. Cavanagh is deemed to have waived his hearing right. After considering material from the investigative file in this matter, the Deputy Administrator now enters his final order without a hearing pursuant to 21 C.F.R. 1301.43(d) and (e) and 1301.46. This final order replaces and supersedes the final order issued on December 22, 1999, and published at 64 FR 73,586 (December 30, 1999).

The Deputy Administrator finds that Dr. Cavanagh currently possesses DEA Certificate of Registration AC9084485 issued to him in Nevada. The Deputy Administrator further finds that on March 18, 1999, the Board of Medical Examiners of the State of Nevada issued its Findings of Fact, Conclusions of Law, and Order revoking Dr. Cavanagh's license to practice medicine in the State of Nevada.

The Deputy Administrator concludes that Dr. Cavanagh is not currently licensed to practice medicine in Nevada, and therefore, it is reasonable to infer that he is not currently authorized to handle controlled substances in that state. The DEA does not have the statutory authority under the Controlled Substances Act to issue or maintain a registration if the applicant or registrant is without state authority to handle controlled substances in the state in which he conducts his business. See 21 U.S.C. 802(21), 823(f) and 824(a)(3). This prerequisite has been consistently upheld. See Romeo J. Perez, M.D., 62 FR 16,193 (1997); Demetris A. Green, M.D., 61 FR 60,728 (1996); Dominick A. Ricci, M.D., 58 FR 51,104 (1993).

Here it is clear that Dr. Cavanagh is not currently authorized to handle controlled substances in the State of Nevada. As a result, Dr. Cavanagh is not entitled to a DEA registration in that state.

Accordingly, the Deputy Administrator of the Drug Enforcement Administration, pursuant to the authority vested in him by 21 USC 823 and 824 and 28 C.F.R. 0.100(b) and 0.104, hereby orders that DEA Certificate of Registration AC9084485, previously issued to James Garvey Cavanagh, M.D., be, and it hereby is, revoked. The Deputy Administrator further orders that any pending applications for the renewal of such registration, be, and they hereby are, denied. This order is effective March 6, 2000, and is considered the final agency action for appellate purposes pursuant to 21 U.S.C. 877.

Dated: January 18, 2000.

Donnie R. Marshall,

Deputy Administrator.

[FR Doc. 00–2526 Filed 2–3–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 99–9]

Michael G. Dolin, M.D., Denial of Request for Modification of Registration

On December 17, 1998, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA) issued an Order to Show Cause to Michael Glen Dolin, M.D. (Respondent) of Rockville Center, New York, notifying him of an opportunity to show cause as to why DEA should not revoke his DEA Certificate of Registration AD4476378 pursuant to 21 U.S.C. 824(a)(4), and deny any pending applications for modification or renewal of such registration pursuant to 21 U.S.C. 823(f), for reason that his registration would be inconsistent with the public interest.

On January 4, 1999, Respondent, through counsel, filed a request for a