

WYOMING**Albany County**

Mountain View Hotel, 2747 WY 130,
Centennial, 07000541

[FR Doc. E7-9171 Filed 5-11-07; 8:45 am]

BILLING CODE 4312-51-P

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701-TA-409 and 731-TA-909 (Review)]

Low Enriched Uranium From France

AGENCY: United States International Trade Commission.

ACTION: Notice of Commission determinations to conduct full five-year reviews concerning the antidumping and countervailing duty orders on low enriched uranium ("LEU") from France.

SUMMARY: The Commission hereby gives notice that it will proceed with full reviews pursuant to section 751(c)(5) of the Tariff Act of 1930 (19 U.S.C. 1675(c)(5)) to determine whether revocation of the antidumping and countervailing duty orders on LEU from France would be likely to lead to continuation or recurrence of material injury within a reasonably foreseeable time. A schedule for the reviews will be established and announced at a later date. For further information concerning the conduct of these reviews and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

DATES: *Effective Date:* April 9, 2007.

FOR FURTHER INFORMATION CONTACT:

Mary Messer (202-205-3193), 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>). The public record for these reviews may be viewed on the Commission's electronic docket (EDIS) at <http://edis.usitc.gov>.

SUPPLEMENTARY INFORMATION: On April 9, 2007, the Commission determined that it should proceed to full reviews in

the subject five-year reviews pursuant to section 751(c)(5) of the Act.¹ The Commission found that both the domestic and respondent interested party group responses to its notice of institution (72 F.R. 144, January 3, 2007) were adequate.² A record of the Commissioners' votes, the Commission's statement on adequacy, and any individual Commissioner's statements will be available from the Office of the Secretary and at the Commission's Web site.

Authority: These reviews are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.62 of the Commission's rules.

By order of the Commission.

Issued: May 8, 2007.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E7-9148 Filed 5-11-07; 8:45 am]

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DEPARTMENT OF JUSTICE**Drug Enforcement Administration****Manufacturer of Controlled Substances; Notice of Application**

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 5, 2006, Noramco Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedule I and II:

Drug	Schedule
Codeine-N-Oxide (9053)	I
Morphine-N-Oxide (9307)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Thebaine (9333)	II
Opium, raw (9600)	II
Opium poppy (9650)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Carfentanil (9743)	II
Fentanyl (9801)	II

¹ Commissioner Okun did not participate.

² Commissioner Williamson dissented with respect to the adequacy of the respondent interested party group response, finding that the respondent interested party group response was inadequate. Commissioner Williamson also found that other circumstances warranted conducting full reviews.

The company plans to manufacture small quantities of the Schedule I controlled substances for internal testing; the Schedule II controlled substances will be manufactured in bulk for distribution to its customers.

Any other such applicant, and any person who is presently registered with the DEA to bulk manufacture such a substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than July 13, 2007.

Dated: May 7, 2007.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E7-9200 Filed 5-11-07; 8:45 am]

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DEPARTMENT OF JUSTICE**Parole Commission**

[(Public Law 94-409) (5 U.S.C. 552b)]

Record of Vote of Meeting Closure

I, Edward F. Reilly, Jr., Chairman of the United States Parole Commission, was present at a meeting of said Commission, which started at approximately 1:30 p.m., on Thursday, April 26, 2007, at the U.S. Parole Commission, 5550 Friendship Boulevard, 4th Floor, Chevy Chase, Maryland 20815. The purpose of the meeting was to decide four petitions for reconsideration pursuant to 28 CFR 2.27. Five Commissioners were present, constituting a quorum when the vote to close the meeting was submitted.

Public announcement further describing the subject matter of the meeting and certifications of General Counsel that this meeting may be closed by vote of the Commissioners present were submitted to the Commissioners prior to the conduct of any other business. Upon motion duly made, seconded, and carried, the following Commissioners voted that the meeting be closed: Edward F. Reilly, Jr., Cranston J. Mitchell, Deborah A.