notice, and to authorize the administrative law judge and the Commission, without further notice to the respondent, to find the facts to be as alleged in the complaint and this notice and to enter a final determination containing such findings, and may result in the issuance of a limited exclusion order or cease and desist order or both directed against the respondent.

By order of the Commission.

Issued: January 11, 2005.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. 05–905 Filed 1–14–05; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigations Nos. 731–TA–340E and H (Second Review)]

Solid Urea From Russia and Ukraine

AGENCY: United States International Trade Commission.

ACTION: Notice of Commission determinations to conduct full five-year reviews concerning the antidumping duty orders on solid urea from Russia and Ukraine.

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 duty orders on solid urea from Russia and Ukraine 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).

EFFECTIVE DATE: January 4, 2005.

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 January 4, 2005, 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. The Commission found that both the domestic and Russian respondent interested party group responses to its notice of institution (69 FR 58957, October 1, 2004) were adequate but it found that the Ukrainian respondent interested party group response was inadequate. However, the Commission determined to conduct a full review concerning subject imports from Ukraine to promote administrative efficiency in light of its decision to conduct a full review with respect to solid urea from Russia. 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: January 12, 2005.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. 05–904 Filed 1–14–05; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[USITC SE–05–001]

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: International Trade Commission.

TIME AND DATE: January 26, 2005, at 2 p.m.

PLACE: Room 101, 500 E Street, SW., Washington, DC 20436, telephone: (202) 205–2000.

STATUS: Open to the public.

MATTERS TO BE CONSIDERED:

1. Agenda for future meetings: None.
2. Minutes.
3. Ratification List.
4. Inv. No. 731–TA–653 (Second Review)(Sebacic Acid from China)—briefing and vote. (The Commission is currently scheduled to transmit its determination and Commissioners' opinions to the Secretary of Commerce on or before February 8, 2005.)
5. Outstanding action jackets:

(1) Document No. GC–04–152: Concerning administrative matters.

(2) Document No. GC–04–173: Concerning Inv. No. 337–TA–406 (Certain Lens-Fitted Film Packages)(Enforcement Proceedings II).

(3) Document No. GC–04–178: Concerning administrative matters.

In accordance with Commission policy, subject matter listed above, not disposed of at the scheduled meeting, may be carried over to the agenda of the following meeting.

Issued: January 12, 2005.

By order of the Commission.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. 05–1019 Filed 1–13–05; 12:10 pm]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Bureau of Alcohol, Tobacco, Firearms and Explosives

Agency Information Collection Activities: Proposed Collection; Comments Requested

ACTION: 60-day notice of information collection under review: certification on agency letterhead authorizing purchase of firearm for official duties of law enforcement officer.

The Department of Justice (DOJ), Bureau of Alcohol, Tobacco, Firearms and Explosives (ATF), 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 proposed information collection is published to obtain comments from the public and affected agencies. Comments are encouraged and will be accepted for ''sixty days'' until March 21, 2005. This process is conducted in accordance with 5 CFR 1320.10.

If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact David Chipman, Chief, Firearms Enforcement Branch, Room 7400, 650 Massachusetts Avenue, NW., Washington, DC 20226.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your

comments should address one or more of the following four points:

—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;

—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;

—Enhance the quality, utility, and clarity of the information to be collected; and

—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.

Overview of This Information Collection:

(1) *Type of Information Collection: Extension of a currently approved collection.*

(2) *Title of the Form/Collection:* Certification on Agency Letterhead Authorizing Purchase of Firearm for Official Duties of Law Enforcement Officer.

(3) *Agency Form Number, if Any, and the Applicable Component of the Department of Justice Sponsoring the Collection:* Form Number: None. Bureau of Alcohol, Tobacco, Firearms and Explosives.

(4) *Affected Public Who Will be Asked or Required to Respond, as Well as a Brief Abstract:* Primary: State, Local or Tribal Government. Other: None. The letter is used by a law enforcement officer to purchase handguns to be used in his/her official duties from a licensed firearm dealer anywhere in the country. The letter shall state that the officer will use the firearm in official duties and that a records check reveals that the purchasing officer has no convictions for misdemeanor crimes of domestic violence.

(5) *An Estimate of the Total Number of Respondents and the Amount of Time Estimated for an Average Respondent to Respond:* It is estimated that 50,000 respondents will take 5 seconds to file the letter.

(6) *An Estimate of the Total Public Burden (in Hours) Associated With the Collection:* There are an estimated 69 annual total burden hours associated with this collection.

If additional information is required contact: Brenda E. Dyer, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, Department of Justice, Patrick Henry Building, Suite 1600, 601 D Street, NW., Washington, DC 20530.

Dated: January 12, 2005.

Brenda E. Dyer,

Department Clearance Officer, Department of Justice.

[FR Doc. 05–901 Filed 1–14–05; 8:45 am]

BILLING CODE 4410–FY–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–261N]

Solicitation of Comments on Dispensing of Controlled Substances for the Treatment of Pain

AGENCY: Drug Enforcement Administration (DEA), Department of Justice.

ACTION: Notice; solicitation of comments.

SUMMARY: On November 16, 2004, DEA published in the **Federal Register** an Interim Policy Statement on the dispensing of controlled substances for the treatment of pain. The Interim Policy Statement stated that DEA would address the subject in greater detail in a future **Federal Register** document, taking into consideration the views of the medical community. DEA is hereby seeking comments from physicians and other interested members of the public as to what areas of the law relating to the dispensing of controlled substances for the treatment of pain they would like DEA to address in the upcoming **Federal Register** document.

DATES: Written comments must be postmarked, and electronic comments must be sent, on or before March 21, 2005.

ADDRESSES: To ensure proper handling of comments, please reference ''Docket No. DEA–261'' on all written and electronic correspondence. Written comments being sent via regular mail should be sent to the Deputy Administrator, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative/CCD. Written comments sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/CCD, 2401 Jefferson-Davis Highway, Alexandria, VA 22301.

Comments may be directly sent to DEA electronically by sending an electronic message to *dea.diversion.policy@usdoj.gov*. Comments may also be sent electronically through *http://www.regulations.gov* using the electronic comment form provided on that site. An electronic copy of this document is also available at the *http://www.regulations.gov* Web site. DEA will accept attachments to electronic comments in Microsoft Word, WordPerfect, Adobe PDF, or Excel file formats only. DEA will not accept any file format other than those specifically listed here.

FOR FURTHER INFORMATION CONTACT: Daniel Dormont, Senior Attorney, Drug Enforcement Administration, Washington, DC 20537; telephone: (202) 307–8010.

SUPPLEMENTARY INFORMATION:

On November 16, 2004, DEA published in the **Federal Register** an Interim Policy Statement on the dispensing of controlled substances for the treatment of pain. 69 FR 67170. The Interim Policy Statement explained why an earlier document, which appeared on the DEA Office of Diversion Control Web site in August 2004, contained misstatements and was not approved as an official statement of the agency. The Interim Policy Statement corrected some of the misstatements in the August 2004 document and announced that DEA would address, in greater detail, the subject of dispensing controlled substances for the treatment of pain in a future **Federal Register** document, taking into consideration the views of the medical community. This upcoming document will stay within the scope of DEA's authority by addressing the law the agency administers, the Controlled Substances Act (CSA), and the DEA regulations promulgated thereunder, as well as the pertinent court decisions. As indicated in the Interim Policy Statement, the document will contain a recitation of the relevant provisions of the CSA and DEA regulations relating to the dispensing of controlled substances for the treatment of pain. The purpose of this recitation will be to provide guidance and reassurance to the overwhelming majority of physicians who engage in legitimate pain treatment while deterring unlawful prescribing and dispensing of pharmaceutical controlled substances.

As was the case with the Interim Policy Statement, none of the principles addressed in the upcoming **Federal Register** document will be new. Rather, the document will reiterate legal concepts that have been incorporated in the federal laws and regulations for many years and are reflected in federal court decisions and DEA final administrative orders. DEA recognizes