(ii) Identify any public health, safety, or welfare concerns in the United States relating to the potential orders;

(iii) Indicate the extent to which like or directly competitive articles are produced in the United States or are otherwise available in the United States, with respect to the articles potentially subject to the orders; and

(iv) Indicate whether Complainant, Complainant’s licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to an exclusion order and a cease and desist order within a commercially reasonable time.

Written submissions must be filed no later than by close of business, five business days after the date of publication of this notice in the Federal Register. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation.

Persons filing written submissions must file the original document and 12 true copies thereof on or before the deadlines stated above with the Office of the Secretary. Submissions should refer to the docket number (“Docket No. 2863”) in a prominent place on the cover page and/or the first page. The Commission’s rules authorize filing submissions with the Secretary by facsimile or electronic means only to the extent permitted by section 201.8 of the rules (see Handbook for Electronic Filing Procedures, http://www.usitc.gov/secretary/fed_reg_notices/rules/documents/handbook_onElectronic_filing.pdf). Persons with questions regarding electronic filing should contact the Secretary (202) 205–2000.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary of Commerce on December 5, 2011.

The views of the Commission are properly sought will be treated accordingly. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary.

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of sections 201.10 and 210.50(a)(4) of the Commission’s Rules of Practice and Procedure (19 CFR 201.10, 210.50(a)(4)).

By order of the Commission.

James R. Holbein,
Secretary to the Commission.

Issued: December 6, 2011.

DEPARTMENT OF JUSTICE

[AA/A Order No. 001/2011]

Privacy Act of 1974; Computer Matching Agreement

AGENCY: Department of Justice.

ACTION: Notice—computer matching between the Department of Justice and the Internal Revenue Service, Department of Treasury.

SUMMARY: In accordance with the Privacy Act of 1974 (5 U.S.C. 552a), as amended by the Computer Matching and Privacy Protection Act of 1988 (Pub. L. 100–503), Office of Management and Budget (OMB) Guidelines on the Conduct of Matching Programs 54 FR 25818 (June 19, 1989), OMB Bulletin 89–22, “Instructions on Reporting Computer Matching Programs to the Office of Management and Budget (OMB), Congress and the Public,” and OMB Circular No. A–130, Revised November 28, 2000, “Management of Federal Information Resources”, the Department of Justice is issuing a public notice of its intent to conduct a computer matching program with the Internal Revenue Service, Department of the Treasury. Under this matching program, entitled Taxpayer Address Request, the IRS will provide information relating to taxpayers’ mailing addresses to the DOJ for purposes of enabling DOJ to locate debtors to initiate litigation and/or enforce the collection of debts owed by the taxpayers to the United States.

DATES: Effective Date: The matching program will become effective 40 days after a copy of the agreement, as approved by the Data Integrity Board of each agency, is sent to Congress and the Office of Management and Budget, or 30 days after publication of this notice in the Federal Register, whichever is later. The matching program will continue for 18 months after the effective date and may be extended for an additional 12 months, if the conditions specified in 5 U.S.C. 552a(o)(2)(D) have been met.


By order of the Commission.

James R. Holbein,
Secretary to the Commission.

Issued: December 6, 2011.

INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 731 TA 340–E and 340–H (Third Review)]

Solid Urea From Russia and Ukraine

Determination

On the basis of the record 1 developed in the subject five-year reviews, the United States International Trade Commission (Commission) determines, pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)), that 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 to an industry in the United States within a reasonably foreseeable time.2

Background

The Commission instituted these reviews on December 1, 2010 (75 FR 74746) and determined on March 7, 2011 that it would conduct full reviews (76 FR 15339, March 21, 2011). Notice of the scheduling of the Commission’s reviews and of a public hearing to be held in connection therewith was given by posting copies of the notice in the Office of the Secretary, U.S. International Trade Commission, Washington, DC, and by publishing the notice in the Federal Register on April 28, 2011 (76 FR 23835). The hearing was held in Washington, DC, on October 4, 2011, and all persons who requested the opportunity were permitted to appear in person or by counsel.


By order of the Commission.

Issued: December 5, 2011.

James R. Holbein,
Secretary to the Commission.

[FR Doc. 2011–31596 Filed 12–8–11; 8:45 am]

BILLING CODE 7020–02–P

1The record is defined in sec. 207.2(f) of the Commission’s Rules of Practice and Procedure (19 CFR 207.2)[f].

2Chairman Deanna Tanner Okun and Commissioner Daniel R. Pearson dissenting.
DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–343F]

Controlled Substances: Final Adjusted Aggregate Production Quotas for 2011

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

ACTION: Notice.

SUMMARY: This notice establishes final adjusted 2011 aggregate production quotas for controlled substances in Schedules I and II of the Controlled Substances Act (CSA).

FOR FURTHER INFORMATION CONTACT: John W. Partridge, Chief, Liaison and Policy Section, Drug Enforcement Administration, 8701 Morrissette Drive, Springfield, VA 22152, Telephone: (202) 307–4564.

SUPPLEMENTARY INFORMATION:

Background

Section 306 of the Controlled Substances Act (CSA) (21 U.S.C. 826) requires that the Attorney General establish aggregate production quotas for each basic class of controlled substance listed in Schedules I and II. This responsibility has been delegated to the Administrator of the DEA by 28 CFR 0.100. In accordance with 21 U.S.C. 826 and 21 CFR 1303.11, DEA published in the Federal Register on December 20, 2010, notice of the established 2011 aggregate production quotas for controlled substances in Schedules I and II (75 FR 79404). That notice stated that the Administrator would adjust, as needed, the established aggregate production quotas in 2011 as provided for in 21 CFR 1303.13. The 2011 proposed adjusted aggregate production quotas were subsequently published in the Federal Register on September 14, 2011, (76 FR 56810) in consideration of the outlined criteria. All interested persons were invited to comment on or object to the proposed adjusted aggregate production quotas on or before October 14, 2011.

The September 14, 2011, proposed adjusted aggregate production quotas also included proposed aggregate production quotas for five newly scheduled substances. On March 1, 2011, the DEA Administrator published a final order (76 FR 11075) which temporarily placed five synthetic cannabinoids in Schedule I: 1-[4-Morpholinyl](ethyl)-3-(1-naphthoyl)indole (JWH-018); 1-Butyl-3-(1-naphthoyl)indole (JWH-073); 1-Pentyl-3-(1-naphthoyl)indole (JWH-018); 6-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxy-cyclohexyl]-phenol; and 5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxy-cyclohexyl]-phenol. That final order stated that quotas for the five substances would be “established based on registrations granted and quota applications received pursuant to part 1303 of Title 21 of the Code of Federal Regulations.” 76 FR 11077. Aggregate production quotas for these newly scheduled substances had not been previously established and were initially proposed in the above referenced notice published in the Federal Register on September 14, 2011 (76 FR 56810). All interested persons were invited to comment on or object to the proposed aggregate production quotas on or before October 14, 2011.

Analysis for Final Adjusted 2011 Aggregate Production Quotas

Consideration has been given to the criteria outlined in the September 14, 2011, notice of proposed adjusted aggregate production quotas in accordance with 21 CFR 1303.13. In addition, six companies, four DEA registered manufacturers and two non-registrants, submitted timely comments regarding a total of 22 Schedule I and II controlled substances. Comments received proposed that the aggregate production quotas for 4-anilino-N,N-dipropyl-1-piperidinyl-4-piperidine (ANPP), alfentanil, amphetamine (for sale), diphenoxylate, fentanyl, gamma hydroxybutyric acid, hydrocodone, meperidine, methadone, methadone...