Commission's Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(d)–(e) and 210.13(a), such responses will be considered by the Commission if received not later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this 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 an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

The Süd-Chemie respondents may present to the presiding ALJ the matter raised in their June 6, 2011 confidential letter to the Commission.

By order of the Commission.

Issued: June 16, 2011.

James R. Holbein,

Secretary to the Commission.

[FR Doc. 2011–15534 Filed 6–21–11; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Notice of Lodging of Consent Decree Pursuant to the Clean Water Act

Notice is hereby given that on June 16, a proposed Consent Decree in *United States and the State of Nebraska* v. *Swift Beef Company,* Civil Action No. 8:11–cv–216 was lodged with the United States Court for the District of Nebraska. In this action, Plaintiffs the United States and State of Nebraska sought the penalties and injunctive relief for violations of the Clean Water Act (''CWA'') by Swift Beef Company (''Swift'') at a beef processing plant it owns and operates in Grand Island, Nebraska. Pursuant to the proposed Consent Decree, Defendants will pay to the United States and the State of Nebraska $1,300,000 in civil penalties and undertake injunctive measures designed to prevent future violations.

For 30 days after the date of this publication, the Department of Justice will receive comments relating to the proposed consent decree. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and either emailed to *pubcomment-ees.enrd@usdoj.gov* or mailed to P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044–7611, and should refer to *United States* v. *Swift Beef Company,* Civil Action No. 8:11–cv–216 (D. Neb.), DJ Reference No. 90–5–1–1–09466.

During the public comment period, the Consent Decree may also be examined on the following Department of Justice Web site: *http://www.usdoj.gov/enrd/Consent_Decrees.html.* A copy of the proposed consent decree may be obtained by mailing a request to the Consent Decree Library, P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044–7611. When requesting a copy by mail, please enclose a check payable to the U.S. Treasury in the amount of $12.00 (25 cents per page reproduction cost). A copy may also be obtained by faxing or e-mailing a request to Tonia Fleetwood, *tonia.fleetwood@usdoj.gov,* fax number (202) 514–0097, phone confirmation number (202) 514–1547, and sending a check to the Consent Decree Library at the stated address.

Robert E. Maher, Jr.,

Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.

[FR Doc. 2011–15465 Filed 6–21–11; 8:45 am]

BILLING CODE 4410–15–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 11, 2011, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Chattanooga, Tennessee 37409, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Opium tincture (9630)	II
Opium, powdered (9639)	II
Opium, granulated (9640)	II
Tapentadol (9780)	II

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers. Regarding (9640) the company plans to manufacture another controlled substance for sale to its customers.

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

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, **Federal Register** Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 22, 2011.

Dated: June 14, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011–15478 Filed 6–21–11; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 4, 2011, Boehringer Ingelheim Chemicals Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805–9372, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Amphetamine (1100)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Methadone (9250)	II
Methadone Intermediate (9254)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers for formulation into finished pharmaceuticals.

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

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive,