Dated: October 8, 2008.

Lynn Bryant,

Department Clearance Officer, PRA, United States Department of Justice.

[FR Doc. E8–24284 Filed 10–10–08; 8:45 am]

BILLING CODE 4410–FY–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this section to a bulk manufacturer of a controlled substance in schedule I or II, and prior to issuing a regulation under 21 U.S.C. 952(a)(2) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on August 26, 2008, Hospira Inc., 1776 North Centennial Drive, McPherson, Kansas 67460–1247, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of Remifentanil (9739), a basic class of controlled substance listed in schedule II.

The company plans to import Remifentanil for use in dosage form manufacturing.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

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 November 13, 2008.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the **Federal Register** on September 23, 1975, (40 FR 43745), all applicants for registration to import a basic class of any controlled substance in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 USC 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: October 6, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8–24308 Filed 10–10–08; 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) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 13, 2008, Halo Pharmaceutical Inc., 30 North Jefferson Road, Whippany, New Jersey 07981, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
Dihydromorphine (9145)	I
Hydromorphone (9150)	II

Dihydromorphine is an intermediate in the manufacture of Hydromorphone and is not for commercial distribution. The company plans to manufacture Hydromorphone HCL for sale to other manufacturers and for the manufacture of other controlled substance dosage units for distribution to its customers.

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 written 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 December 15, 2008.

Dated: October 7, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8–24310 Filed 10–10–08; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket Nos. 05–13 and 05–45]

Sunny Wholesale, Inc. Revocation of Registration and Denial of Application; Correction

On October 3, 2008, the Drug Enforcement Administration (DEA) published an order in the **Federal Register** (73 FR 57655) that, among other things, revoked the registration of Sunny Wholesale, Inc. In the order taking this action, the DEA Certificate of Registration was incorrectly cited. The correct Certificate of Registration for Sunny Wholesale, Inc., 120 Forest Parkway, Forest Park, Georgia, is 004550SLY.

Therefore, the Certificate of Registration referenced at 73 FR 57668, first column, sixth line down, is corrected to read ''004550SLY.''

Dated: October 6, 2008.

Michele M. Leonhart,

Deputy Administrator.

[FR Doc. E8–24305 Filed 10–10–08; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Office of Justice Programs

[OMB Number 1121–NEW]

Agency Information Collection Activities: New Information Collection, Comments Requested

ACTION: 30-Day Notice of Information Collection Under Review: NICS Act State Record Estimates Data Collection.

The Department of Justice (DOJ), Office of Justice Programs, Bureau of Justice Statistics (BJS), will be submitting 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. This proposed information collection was previously published in the **Federal Register** Volume 73, Number 150, page 45245 on August 4, 2008, allowing for a 60 day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until November 13, 2008. This process is conducted in accordance with 5 CFR 1320.10.

The proposed information collection is available online at *http://www.ojp.usdoj.gov/bjs/niaa.htm.*