

Dated: June 20, 2003
Laura M. Nagel,
Deputy Assistance Administrator, Office of
Diversion Control, Drug Enforcement
Administration.
[FR Doc. 03-17124 Filed 7-7-03; 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 May 8, 2003, Cody
Laboratories, Inc., made application by
letter to the Drug Enforcement
Administration (DEA) for registration as
a bulk manufacturer of the basic classes
of controlled substances listed below:
Diphenoxylate (9170)—Schedule II
Meperidine (9230)—Schedule II
Oxymorphone (9652)—Schedule II
Sufentanil (9740)—Schedule II

The firm plans to manufacture bulk
material 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.

Any such comments or objections
may be addressed, in quintuplicate, to
the Deputy Assistant Administrator,
Office of Diversion Control, Drug
Enforcement Administration, United
States Department of Justice,
Washington, DC 20537, Attention: DEA
Federal Register Representative (CCD)
and must be filed no later than
September 8, 2003.

Laura M. Nagel,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.
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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 April 21, 2003,
Lilly Del Caribe, Inc., Chemical Plant,
Kilometer 146.7, State Road 2,
Mayaguez, Puerto Rico 00680, made
application by renewal to the Drug
Enforcement Administration (DEA) for
registration as a bulk manufacturer of

Dextropropoxyphene (9273), a basic
class of controlled substance listed in
Schedule II.

The firm plans to bulk manufacture
product 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.

Any such comments or objections
may be addressed, in quintuplicate, to
the Deputy Assistant Administrator,
Office of Diversion Control, Drug
Enforcement Administration, United
States Department of Justice,
Washington, DC 20537, Attention:
Federal Register Representative, Office
of Chief Counsel (CCD) and must be
filed no later than September 8, 2003.

Dated: June 20, 2003.
Laura M. Nagel,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.
[FR Doc. 03-17125 Filed 7-7-03; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled
Substances; Notice of Registration

By Notice dated January 27, 2003, and
published in the Federal Register on
February 6, 2003, (68 FR 6184),
Noramco, Inc., 1440 Olympic Drive,
Athens, GA 30601, made application by
renewal to the Drug Enforcement
Administration to be registered as a bulk
manufacturer of the basic classes of
controlled substances listed below:

Table with 2 columns: Drug, Schedule. Rows include Amphetamine, Oxycodone (9143), Hydrocodone (9193), Morphine (9300), Thebaine (9333), Sufentanil (9740), and Fentanyl (9801).

The firm plans to support its other
manufacturing facility with
manufacturing and analytical testing.

No comments or objections have been
received. DEA has considered the
factors in Title 21, United States Code,
section 823(a) and determined that the
registration of Noramco, Inc., to
manufacture the listed controlled
substances is consistent with the public
interest at this time. DEA has
investigated Noramco, Inc., to ensure
that the company's registration is
consistent with the public interest. This

investigation has included inspection
and testing of the company's physical
security systems, verification of the
company's compliance with state and
local laws, and a review of the
company's background and history.
Therefore, pursuant to 21 U.S.C. 823
and 28 CFR 0.100 and 0.104, the Deputy
Assistant Administrator, Office of
Diversion Control, hereby orders that
the application submitted by the above
firm for registration as a bulk
manufacturer of the basic classes of
controlled substances listed above is
granted.

Dated: June 20, 2003.
Laura M. Nagel,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.
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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 March 14, 2003,
Pressure Chemical Company, 3419
Smallman Street, Pittsburgh,
Pennsylvania 15201, made application
by renewal to the Drug Enforcement
Administration (DEA) for registration as
a bulk manufacturer of 2, 5-
Dimethoxyamphetamine (7396), a basic
class of controlled substance listed in
Schedule I.

The firm plans to manufacturer the
substance for distribution to its
customers.

Any other such applicant and any
person who is presently registered with
DEA to manufacturer such substances
may file comments or objections to the
issuance of the proposed registration.

Any such comments or objections
may be addressed, in quintuplicate, to
the Deputy Assistant Administrator,
Office of Diversion Control, Drug
enforcement Administration, United
States Department of Justice,
Washington, DC 20537, Attention:
Federal Register Representative, Office
of Chief Counsel (CCD) and must be
filed no later than September 8, 2003.

Dated: June 20, 2003.
Laura M. Nagel,
Deputy Assistant Administrator, Office of
Diversion Control, Drug Enforcement
Administration.
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