Dated: February 15, 2005.

James A. Esget,

Program Manager.

[FR Doc. 05–3751 Filed 2–25–05; 8:45 am]

BILLING CODE 4310–MN–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to 21 CFR 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 13, 2005, Boehringer Ingelheim Chemical Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Tetrahydrocannabinols (7370), a basic class of controlled substance listed in Schedule I.

The company plans to manufacture the listed controlled substance in bulk for use in analysis and drug test standards.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance 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 being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than [60 days from publication].

Dated: February 17, 2005.

William J. Walker,

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

[FR Doc. 05–3798 Filed 2–25–05; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to 21 CFR 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 28, 2004, Green Acres Farms, Inc., Rebecca Marie Yale, 5532 Frances Avenue, Tacoma, Washington 98422, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic class of controlled substances listed in Schedule I:

Drug	Schedule
Marijuana (7360)	I
Tetrahydrocannabinols (7370)	I

The applicant plans to manufacture (cultivate) Marijuana and Tetrahydrocannabinols in bulk for distribution. As documented in the applicant's response to the bulk manufacturer questionnaire submitted to the Drug Enforcement Administration (DEA), Green Acres Farms, Inc. stated its plans ''to support the medical marijuana market. It is our intention to manufacture, package and sell to the various authorized outlets within each state that has passed a law by its citizens to allow the medicinal use of marijuana.''

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 being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than (60 days from publication).

Dated: February 17, 2005.

William J. Walker,

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

[FR Doc. 05–3799 Filed 2–25–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(1), 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)(b) authorizing the importation of such substances, provide manufacturers holding registrations for the bulk manufacture of the substances an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on November 11, 2004, JFC Technologies, LLC, 100 West Main Street, PO Box 669, Bound Brook, New Jersey 08805, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of Meperidine-Intermediate B (9233), a basic class of controlled substance listed in Schedule II.

The company plans to import the listed controlled substance for the manufacture of controlled substances in bulk for distribution to its customers.

Any manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file written 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 written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than March 30, 2005.

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 43745–46), all applicants for registration to import 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 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: February 17, 2005.

William J. Walker,

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

[FR Doc. 05–3800 Filed 2–25–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

[Docket Nos. NRTL2–98, NRTL1–89]

NSF International, Expansion of Recognition; Application for Renewal of Recognition; Intertek Testing Services, NA, Interim Approval Subject to Review

AGENCY: Occupational Safety and Health Administration (OSHA), Labor.

ACTION: Notice.

SUMMARY: This notice announces the Agency's final decision on the application of NSF International (NSF) for expansion of its recognition as a Nationally Recognized Testing Laboratory under 29 CFR 1910.7. This notice also announces NSF's Application for renewal of its recognition and presents the Agency's preliminary finding on the renewal. This preliminary finding does not constitute an interim or temporary approval of the renewal application.

In an unrelated matter, we are adding one test standard, NFPA 72, Installation, Maintenance, and Use of Protective Signaling Systems, to the scope of recognition of Intertek Testing Services, NA (ITSNA), on an interim basis, subject to review.

DATES: *Recognition:* The expansion of recognition becomes effective on February 28, 2005 and, unless modified in accordance with 29 CFR 1910.7, continues in effect while NSF remains recognized by OSHA as an NRTL. *Renewal:* Comments on the renewal of recognition must be received no later than March 15, 2005. *Comments on Interim Approval:* Comments on the interim approval for test standard NFPA 72 must be received no later than March 15, 2005.

You may submit comments in response to the renewal application and the interim approval portions of this notice, or any request for extension of the time to comment, by (1) regular mail, (2) express or overnight delivery service, (3) hand delivery, (4) messenger service, or (5) FAX transmission (facsimile). Because of security-related problems there may be a significant delay in the receipt of comments by regular mail. Comments (or any request for extension of the time to comment) must be submitted by the following dates:

Regular mail and express delivery service: Your comments must be postmarked by March 15, 2005.

Hand delivery and messenger service: Your comments must be received in the OSHA Docket Office by March 15, 2005. OSHA Docket Office and Department of Labor hours of operation are 8:15 a.m. to 4:45 p.m.

Facsimile and electronic transmission: Your comments must be sent by March 15, 2005.

ADDRESSES: Regular mail, express delivery, hand-delivery, and messenger service: You must submit three copies of your comments and attachments to the OSHA Docket Office, Docket No. NRTL2–98 or Docket No. NRTL1–89 (as applicable), Room N–2625, U.S. Department of Labor, Occupational Safety and Health Administration, 200 Constitution Avenue, NW., Washington, DC 20210. Please contact the OSHA Docket Office at (202) 693–2350 for information about security procedures concerning the delivery of materials by express delivery, hand delivery and messenger service.

Facsimile: If your comments, including any attachments, are 10 pages or fewer, you may fax them to the OSHA Docket Office at (202) 693–1648. You must include the docket number of this notice, Docket No. NRTL2–98 or Docket No. NRTL1–89 (as applicable), in your comments.

Internet access to comments and submissions: OSHA will place comments and submissions in response to this notice on the OSHA Web page *http://www.osha.gov*. Accordingly, OSHA cautions you about submitting information of a personal nature (*e.g.*, social security number, date of birth). There may be a lag time between when comments and submissions are received and when they are placed on the Webpage. Please contact the OSHA Docket Office at (202) 693–2350 for information about materials not available through the OSHA Webpage and for assistance in using the Webpage to locate docket submissions. Comments and submissions will also be available for inspection and copying at the OSHA Docket Office at the address above.

Extension of Comment Period: Submit requests for extensions concerning this notice to: Office of Technical Programs and Coordination Activities, NRTL Program, Occupational Safety and Health Administration, U.S. Department of Labor, Room N3653, 200 Constitution Avenue, NW., Washington, DC 20210. Or fax to (202) 693–1644.

FOR FURTHER INFORMATION CONTACT: Bernard Pasquet, Office of Technical Programs and Coordination Activities, NRTL Program, Occupational Safety and Health Administration, U.S. Department of Labor, 200 Constitution Avenue, NW., Room N3653, Washington, DC 20210, or phone (202) 693–2110.

SUPPLEMENTARY INFORMATION:

Notice of Final Decision on the Expansion of Recognition

The Occupational Safety and Health Administration (OSHA) hereby gives notice of the expansion of recognition of NSF International (NSF) as a Nationally Recognized Testing Laboratory (NRTL). NSF's expansion covers the use of an additional test standard and two supplemental programs. OSHA's current scope of recognition for NSF may be found in the following informational Web page: *http://www.osha.gov/dts/otpca/nrtl/nsf.html*.

OSHA recognition of an NRTL signifies that the organization has met the legal requirements in Section 1910.7 of Title 29, Code of Federal Regulations (29 CFR 1910.7). Recognition is an acknowledgment that the organization can perform independent safety testing and certification of the specific products covered within its scope of recognition and is not a delegation or grant of government authority. As a result of recognition, employers may use products ''properly certified'' by the NRTL to meet OSHA standards that require testing and certification.

The Agency processes applications by an NRTL for initial recognition or for expansion or renewal of this recognition following requirements in Appendix A to 29 CFR 1910.7. This appendix requires that the Agency publish two notices in the **Federal Register** in processing an application. In the first notice, OSHA announces the application and provides its preliminary finding and, in the second notice, the Agency provides its final decision on an application. These notices set forth the NRTL's scope of recognition or modifications of this scope.

NSF submitted an application, dated October 8, 2003 (*see* Exhibit 14), to expand its recognition to include one additional test standard. Prior to submitting this application, NSF submitted another application, dated July 31, 2003 (*see* exhibit 15) to include several additional programs within its current scope of recognition. The NRTL Program staff performed an on-site review (assessment) of NSF's NRTL facilities and in the on-site review