Consent Decree. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044–7611, and should refer to *United States, et al* v. *Conoco, Inc.*, D.J. Ref. 90–5–2–1–07295/1.

During the public comment period the Consent Decree may be examined at the Office of the United States Attorney, Southern District of Texas, U.S. Courthouse, 515 Rusk, Houston, Texas 77002. The Amendment may also be examined on the following Department of Justice Web site, *http://www.usdoj.gov/enrd/open.html*. A copy of the Amendment may also be obtained by mail from the Consent Decree Library, P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044–7611 or by faxing or e-mailing a request to Tonia Fleetwood (*tonia.fleetwood@usdoj.gov*), fax no. (202) 514–0097, phone confirmation number (202) 514–1547. In requesting a copy from the Consent Decree Library, please enclose a check in the amount of $9.75 (25 cents per page reproduction cost) payable to the U.S. Treasury.

Robert D. Brook,

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

[FR Doc. 06–6488 Filed 7–25–06; 8:45 am]

BILLING CODE 4410–15–M

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Nanoparticle Benchmarking Research Project

Notice is hereby given that, on June 21, 2006, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), Nanoparticle Benchmarking Research Project (''Project'') has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) The identities of the parties to the venture and (2) the nature and objectives of the venture. The notifications were filed for the purpose of invoking the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances.

Pursuant to Section 6(b) of the Act, the identities of the parties to the venture are: E.I. du Pont de Nemours and Company, Wilmington, DE; Procter & Gamble Company, West Chester, OH; The Dow Chemical Company, Midland, MI; Cabot Corporation, Boston, MA; Air Products and Chemicals Incorporated, Allentown, PA; Rohm & Haas Company, Spring House, PA; PPG Industries Incorporated, Pittsburgh, PA; Intel Corporation, Santa Clara, CA; Degussa Corporation, Parsippany, NJ; and United Kingdom Health & Safety Executive, London, UNITED KINGDOM. The general area of Project's planned activity is to undertake research and development in the areas of health, safety, and environmental considerations raised by the exposure of workers to airborne nanoparticles in the production of goods. Specifically, Project's objectives include: (1) Design and development of portable workplace monitoring instrumentation; and (2) development and testing of protective clothing fabrics as a barrier to an aerosol of nanoparticles. This work is being jointly funded by DuPont, and the other entites names above, as sponsors who are interested in nanoparticle research.

Dorothy B. Fountain,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 06–6469 Filed 7–25–05; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 9, 2006, Lin Zhi International Inc., 687 North Pastoria Avenue, Sunnyvale, California 94085, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
3,4-Methylenedioxy- methamphetamine (7405).	I
Cocaine (9041)	II
Oxycodone	II
Hydrocodone (9193)	II
Methadone (9250)	II
Dextropropoxyphene, bulk, (9273)	II
Morphine (9300)	II

The company plans to manufacture the listed controlled substances as bulk reagents for use in drug abuse testing.

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 should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative/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 September 25, 2006.

Dated: July 19, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–11931 Filed 7–25–06; 8:45 am]

BILLING CODE 4410–09–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 21 CFR 1301.34(a), this is notice that on June 30, 2005, Meridian Medical Technologies, 255 Hermelin Drive, St. Louis, Missouri 63144, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of Morphine (9300), a basic class of controlled substance listed in Schedule II.

The company plans to import products for research experimentation or clinical use and analytical testing.

Any manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic class 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 written comments or objections being sent via regular mail