

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993 Wireless Industrial Technology Konsortium Inc.

Correction

In notice document E8-21742 appearing on page 54170 in the issue of Thursday, September 18, 2008, make the following corrections:

In the second column, in the first full paragraph, the 12th and 13th lines are corrected to read as follows: "Siemens AG, Karlsruhe, GERMANY; and ABBAutomation Productions GmbH, Alzenau, GERMANY."

[FR Doc. Z8-21742 Filed 10-7-08; 8:45 am]

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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 1, 2008, Clinical Supplies Management, Inc., 342 42nd Street, South Fargo, North Dakota 58103, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Sufentanil (9740), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance for clinical trials, research, and analytical purposes.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic class of controlled substance 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 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than November 7, 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 schedules 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: October 2, 2008.

Joseph T. Rannazzisi,
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 September 8, 2008, National Center for Natural Products Research—NIDA MProject, University of Mississippi, 135 Coy Waller Complex, University, Mississippi 38677, 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:

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

The company plans to cultivate marihuana for the National Institute on Drug Abuse for research approved by the Department of Health and Human Services.

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 Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than December 8, 2008.

Dated: October 2, 2008.

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

[FR Doc. E8-23816 Filed 10-7-08; 8:45 am]

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DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-63,691]

Newpage Corporation, Niagara Mill, Including On-Site Leased Workers From PSI, Naico, Gunville Trucking, Advanced Service Providers and Scott Vancourt, Inc., Niagara, WI; Amended Certification Regarding Eligibility To Apply for Worker Adjustment Assistance and Alternative Trade Adjustment Assistance

In accordance with Section 223 of the Trade Act of 1974 (19 U.S.C. 2273), and Section 246 of the Trade Act of 1974 (26 U.S.C. 2813), as amended, the Department of Labor issued a Certification of Eligibility to Apply for Worker Adjustment Assistance and Alternative Trade Adjustment Assistance on July 31, 2008, applicable to workers of NewPage Corporation, Niagara Mill, including on-site leased workers from PSI, Naico and Gunville Trucking, Niagara, Wisconsin. The notice was published in the **Federal Register** on August 12, 2008 (73 FR 46923). The certification was amended on August 20, 2008 to include on-site leased workers from Advanced Service Providers. The notice of published in the **Federal Register** on August 29, 2008 (73 FR 51001)

At the request of the State agency, the Department reviewed the certification for workers of the subject firm. The workers are engaged in the production of coated mechanical printing paper.

New information shows that workers leased from Scott VanCourt, Inc. were employed on-site at the Niagara, Wisconsin location of NewPage Corporation, Niagara Mill. The Department has determined that these workers were sufficiently under the