

provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, ALSAN Research, Ankey, IA; Gharda Chemicals Ltd., Washington, DC; and Tri Corporation, Houston, TX has become members. Additionally, the following companies have changed their names: (1) Ciba-Geigy Corporation, Greenboro, NC, owned by Ciba-Geigy Ltd. Group, Basel, SWITZERLAND and Sandoz AGRO, Des Plaines, IA have merged to become Novartis Crop Protection, Inc., Greensboro, NC; and (2) Akzo Nobel Chemicals b.v., Arnhem, NETHERLANDS has been acquired by NuFarm Limited, Laverton, North Victoria, AUSTRALIA.

No other changes have been made in either the membership, corporate name, or planned activities of the venture.

On May 15, 1990, the Spray Drift Task Force filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on July 5, 1990 (55 FR 27701). The last notification was filed with the Department on November 16, 1995. A notice was published in the

Federal Register on April 8, 1996 (61 FR 15522).

Constance K. Robinson,
Director of Operations, Antitrust Division.
[FR Doc. 97-15502 Filed 6-12-97; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated February 21, 1997, and published in the **Federal Register** on March 28, 1997, (62 FR 14946), Knight Seed Company, Inc., 151 W. 126th Street, Burnsville, Minnesota 55337, made application to the Drug Enforcement Administration to be registered as an importer of marihuana (7360), a basic class of controlled substance listed in Schedule I.

No comments or objections have been received. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of

Federal Regulations, Section 1311.42, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: May 13, 1997.

Terrance W. Woodworth,
Acting 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 Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 16, 1997, Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Pholcodine (9314)	I
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Benzoylcegonine (9180)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone-intermediate (9254)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Morphine (9300)	II
Thebaine (9333)	II
Opium extracts (9610)	II
Opium fluid extract (9620)	II
Opium tincture (9630)	II
Opium powdered (9639)	II
Opium granulated (9640)	II
Levo-alphaacetylmethadol (9648)	II
Oxymorphone (9652)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

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 Acting Deputy Assistant Administrator, Office of Diversion

Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than August 12, 1997.

Dated: May 7, 1997.

Terrance W. Woodworth,
Acting Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

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