DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—National Center for Manufacturing Sciences, Inc. (NCMS)

Notice is hereby given that, on March 3, 1997, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), the National Center for Manufacturing Sciences, Inc. (''NCMS'') has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing a change in the participants in a joint venture identified as ''Rapid Response Manufacturing.'' The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, Structural Dynamics Research Corporation, Milford, OH, has been added as participant in the joint venture.

No other changes have been made in the joint venture, and its nature and objectives remain unchanged. NCMS intends to file additional written notification disclosing all changes in the joint venture.

On February 20, 1987, NCMS 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 of March 17, 1987 (52 FR 8375).

The last notification was filed with the Department on February 26, 1997. This notice has not been published in the **Federal Register**.

Constance K. Robinson,

Director of Operations, Antitrust Division.

[FR Doc. 97–10950 Filed 4–28–97; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—National Industrial Information Infrastructure Protocols Solutions for Manufacturing—Adaptable Replicable Technology

Notice is hereby given that, on March 21, 1997, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), the National Industrial Information Infrastructure Protocols Solutions for Manufacturing—Adaptable Replicable Technology (''NIIIP–SMART'') filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. 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.

Specifically, the following organization has joined NIIIP–SMART: Pilot Industries, Inc. The following organizations have withdrawn their membership from NIIIP–SMART: Consilium; General Motors Corporation; and Promis.

No other changes have been made in either the membership or planned activities of NIIIP–SMART. Membership remains open and NIIIP–SMART intends to file additional written notifications disclosing all changes in membership.

On May 1, 1996, NIIIP–SMART 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 June 13, 1996 (61 FR 30098).

Constance K. Robinson,

Director of Operations, Antitrust Division.

[FR Doc. 97–10949 Filed 4–28–97; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on February 12, 1997, Celgene Corporation, 7 Powder Horn Drive, Warren, New Jersey 07059, made application to the Drug Enforcement Administration (DEA) for registration by letter as a bulk manufacturer of the Schedule II controlled substance methylphenidate (1724).

The firm plans to manufacture methylphenidate for clinical trials, formulation studies, and product research and development.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the above application.

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 June 30, 1997.

Dated: March 31, 1997.

Terrance W. Woodworth,

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

[FR Doc. 97–11036 Filed 4–28–97; 8:45 am]

BILLING CODE 4410–09–M

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 January 29, 1997, Celgene Corporation, 7 Powder Horn Drive, Warren, New Jersey 07059, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the Schedule I controlled substance 4-Methoxyamphetamine (7411).

The firm plans to manufacture 4-methoxyamphetamine which is used as an intermediate in the manufacture of a non-controlled substance.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the above application.

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 June 30, 1997.

Dated: April 8, 1997.

Terrance W. Woodworth,

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

[FR Doc. 97–11037 Filed 4–28–97; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Samuel Wise Chang, M.D.; Revocation of Registration

On October 24, 1996, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA) issued an Order