No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and CMX intends to file additional written notifications disclosing all changes in membership.

On March 12, 2010, CMX 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 April 16, 2010 (75 FR 20003).

The last notification was filed with the Department on May 28, 2010. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on July 14, 2010 (75 FR 40851).

Patricia A. Brink,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 2010–25212 Filed 10–8–10; 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—Marine Well Containment Venture

Notice is hereby given that, on August 18, 2010, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 et seq. ("the Act"), Marine Well Containment Venture ("MWCV") has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing (1) the identities and nationalities of the parties to the production venture and any person who controls a party 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: Chevron USA, Inc., Houston, TX; ConocoPhillips Co., Houston, TX; ExxonMobil Development Co., Houston, TX; and Shell Offshore Inc., Houston, TX. The general area of MWCV's planned activity is (i) to design, produce (assemble and/or construct), operate, maintain, and own a system to provide emergency hydrocarbon well containment and related non-emergency services in the Gulf of Mexico and potentially in other regions; and (ii) to perform and sponsor related research and development activities.

Patricia A. Brink,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 2010–25206 Filed 10–8–10; 8:45 am] BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 10, 2008, and published in the **Federal Register** on March 19, 2008 (73 FR 14841), Chemica, Inc., 316 West 130th Street, Los Angeles, California 90061, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of methamphetamine (1105), a basic class of controlled substance listed in schedule II.

The above-listed controlled substance is an intermediate in the manufacture of Benzphetamine, a schedule III nonnarcotic controlled substance. The methamphetamine will not be sold as a commercial product in the domestic market.

A comment and objection was received. However, after a thorough review of this matter, DEA has concluded that issues raised in the comment and objection do not warrant the denial of this application.

DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Chemica, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Chemica, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: October 5, 2010.

Joseph T. Rannazzisi,

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

[FR Doc. 2010–25540 Filed 10–8–10; 8:45 am] BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated June 17, 2010, and published in the **Federal Register** on June 28, 2010 (75 FR 36683), Siegfried (USA), 33 Industrial Park Road, Pennsville, New Jersey 08070, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Gamma Hydroxybutyric Acid (2010), a basic class of controlled substance listed in schedule I.

The company plans to manufacture the listed controlled substance in bulk for sale to customers.

Three comments were received. Two of the three comments supported the granting of registration as a bulk manufacturer of the basic class of controlled substance listed to this applicant.

The third comment objected to the granting of registration. However, after a thorough review of this matter, DEA has concluded that the issues raised in the comment and objection do not warrant the denial of this application.

DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Siegfried (USA) to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Siegfried (USA) to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: October 5, 2010.

Joseph T. Rannazzisi,

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

[FR Doc. 2010–25539 Filed 10–8–10; 8:45 am]

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