
The general area of SpeC’s planned activity is (1) minimize barriers to entry businesses and non-traditional vendors to work with the U.S. Government and to identify and realize teaming opportunities among entities to promote integrated research and prototyping efficiencies, and (2) reducing the cost of prototype development.

Suzanne Morris,
Chief, Premerger and Division Statistics Unit,
Antitrust Division.

[FR Doc. 2018–21431 Filed 10–1–18; 8:45 am]
BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Digital Manufacturing Design Innovation Institute

Notice is hereby given that, on September 19, 2018, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 et seq. (“the Act”), Digital Manufacturing Design Innovation Institute (“DMDII”) has 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 extending the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, The Coca-Cola Company, Atlanta, GA; Phynsys LLC, Cincinnati, OH; PUNDITAS LLC, Wakefield, MA; United Electric Corporation, Canonsburg, PA;
Correction

In the Federal Register on August 15, 2018, in FR Doc No: 2018–17605 (83 FR 158) on pages 40567 and 40568, correct the table to include the following basic class of controlled substance:

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fentanyl</td>
<td>9801</td>
<td>II</td>
</tr>
</tbody>
</table>


John J. Martin,
Assistant Administrator.
[FR Doc. 2018–21432 Filed 10–1–18; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration
[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: AMPAC Fine Chemicals, LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before December 3, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on March 14, 2018, AMPAC Fine Chemicals, LLC, Highway 50 and Hazel Avenue, Building 05001, Rancho Cordova, California 95670 applied to be registered.