



**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Bulk Manufacturer of Controlled Substances Application: AMPAC Fine Chemicals**

**LLC**

**[Docket No. DEA-392]**

**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 in accordance with 21 CFR 1301.33(a) on or before [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

**ADDRESS:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODXL, 8701 Morrisette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrisette 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 Deputy Assistant Administrator of the DEA Office of Diversion Control (“Deputy 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 20, 2015, AMPAC Fine Chemicals LLC, Highway 50 and Hazel Avenue, Building 05001, Rancho Cordova, California 95670 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

| <b><u>Controlled Substance</u></b> | <b><u>Schedule</u></b> |
|------------------------------------|------------------------|
| Methylphenidate (1724)             | II                     |
| Thebaine (9333)                    | II                     |
| Poppy Straw Concentrate (9670)     | II                     |
| Tapentadol (9780)                  | II                     |

The company is a contract manufacturer. In reference to Poppy Straw Concentrate the company will manufacture thebaine intermediates for sale to its customers for further manufacture. No other activity for this drug code is authorized for this registration.

Dated: June 25, 2015

Joseph T. Rannazzisi,  
*Deputy Assistant Administrator.*

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