

When respondents complete a fillable form, they may submit it by uploading it to a secure webserver, emailing it to the study team at mtbeffects@usitc.gov, faxing it, or mailing a hard copy to the Commission.

III. Request for Comments

Comments are invited on (1) whether the proposed collection of information is necessary; (2) the accuracy of the agency’s estimate of the burden (including hours and cost) of the proposed collection of information; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

The draft questionnaire and other supplementary documents may be downloaded from the USITC website at <https://www.usitc.gov/MTBEeffects>.

Comments submitted in response to this notice will be summarized and/or included in the request for OMB approval of this information collection;

they will also become a matter of public record.

By order of the Commission.
Issued: September 18, 2018.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2018–20657 Filed 9–21–18; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Halo Pharmaceutical, Inc.

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 November 23, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 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 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 July 18, 2018, Halo Pharmaceutical, Inc., 30 North Jefferson Road, Whippany, New Jersey 07981–1030 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Dihydromorphone	9145	I
Hydromorphone	9150	II

The company plans to manufacture Hydromorphone (9150) for distribution to its customers. Dihydromorphone (9145) as an intermediate in the manufacture of Hydromorphone and is not for commercial distribution.

Dated: September 14, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–20701 Filed 9–21–18; 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 Virginia, 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 November 23, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 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 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 April 19, 2018, AMPAC Fine Chemicals Virginia, LLC, 2820 North Normandy Drive, Petersburg, Virginia 23805–2380 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Phenylacetone	8501	II
Morphine	9300	II
Thebaine	9333	II
Noroxymorphone	9668	II