The company plans to import the FDA approved drug product in finished dosage form for distribution to its customers.

Dated: July 10, 2015.
Joseph T. Rannazzisi,
Deputy Assistant Administrator.
[FR Doc. 2015–17514 Filed 7–16–15; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Mallinckrodt, LLC

ACTION: Notice of registration.

SUMMARY: Mallinckrodt, LLC applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Mallinckrodt, LLC registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated January 21, 2015, and published in the Federal Register on January 28, 2015, 80 FR 4592, Mallinckrodt LLC, 3600 North Second Street, St. Louis, Missouri 63147 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted to this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Mallinckrodt, LLC to manufacture the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company’s maintenance of effective controls against diversion by inspecting and testing the company’s physical security systems, verifying the company’s compliance with state and local laws, and reviewing the company’s background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above-named company is granted registration as a bulk manufacturer of the following basic classes of controlled substances:

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gamma Hydroxybutyric Acid (2010)</td>
<td>I</td>
</tr>
<tr>
<td>Lisdexamfetamine (1205)</td>
<td>II</td>
</tr>
<tr>
<td>Oripavine (9330)</td>
<td>II</td>
</tr>
<tr>
<td>Tapentadol (9780)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to manufacturer bulk active pharmaceutical ingredients (API) for distribution and product development to its customers.

Dated: July 10, 2015.
Joseph T. Rannazzisi,
Deputy Assistant Administrator.
[FR Doc. 2015–17523 Filed 7–16–15; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Navinta LLC

ACTION: Notice of registration.

SUMMARY: Navinta LLC applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Navinta LLC registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated February 11, 2015, and published in the Federal Register on February 19, 2015, 80 FR 8901, Navinta LLC, 1499 Lower Ferry Road, Ewing, New Jersey 08618–1414 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Navinta LLC to manufacture the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company’s maintenance of effective controls against diversion by inspecting and testing the company’s physical security systems, verifying the company’s compliance with state and local laws, and reviewing the company’s background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the above-named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed:

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Remifentanil (9739)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to initially manufacture API quantities of the listed controlled substances for validation purposes and FDA approval, and then produce commercial size batches for distribution to dosage form manufacturers upon FDA approval.

Dated: July 10, 2015.
Joseph T. Rannazzisi,
Deputy Assistant Administrator.
[FR Doc. 2015–17525 Filed 7–16–15; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
[Docket No. DEA–392]

Importer of Controlled Substances Registration: Siegfried USA, LLC

ACTION: Notice of registration.

SUMMARY: Siegfried USA, LLC applied to be registered as an importer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Siegfried USA, LLC registration as an importer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated April 14, 2015, and published in the Federal Register on April 22, 2015, 80 FR 22561, Siegfried USA, LLC, 33 Industrial Park Road, Pennsville, New Jersey 08070 applied to be registered as an importer of certain basic classes of controlled substances. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 23, 2007).

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Siegfried USA, LLC to import the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company’s maintenance of effective controls against diversion by inspecting and testing the company’s physical security systems, verifying the company’s compliance with state and local laws, and reviewing the company’s background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above-named company is granted registration as an
importer of the basic classes of controlled substances:

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Opium, raw (9600)</td>
<td>II</td>
</tr>
<tr>
<td>Poppy Straw Concentrate (9670)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances to bulk manufacture API’s for distribution to its customer.

Dated: July 10, 2015.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

FOR FURTHER INFORMATION CONTACT:

Lashon Hilliard, Department of Justice, Community Oriented Policing Services (COPS) Office, 145 N Street NE., 3E.405B, Washington, DC 20530.

Dated: July 14, 2015.

Jerri Murray,
Department Clearance Officer for PRA, U.S. Department of Justice.

DEPARTMENT OF JUSTICE

[OMB Number 1103–0093]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Revision of a Previously Approved Collection COPS Extension Request Form

AGENCY Community Oriented Policing Services (COPS) Office, Department of Justice.

ACTION 30-day notice.

SUMMARY: The Department of Justice (DOJ), Community Oriented Policing Services (COPS) Office, will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. This proposed information collection was previously published in 80 FR 9750, on February 24, 2015, allowing for a 60 day comment period.

DATES: Comments are encouraged and will be accepted for an additional 30 days until August 17, 2015.

FOR FURTHER INFORMATION CONTACT: If you have additional comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Lashon Hilliard, Department of Justice, Community Oriented Policing Services (COPS) Office, 145 N Street NE., Washington, DC 20530 (202–514–6563). Written comments and/or suggestions can also be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20530 or sent to OIRA_submissions@omb.eop.gov.

SUPPLEMENTARY INFORMATION: Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

1. Type of Information Collection: Revision of a currently approved collection, with change; comments requested.
2. The Title of the Form/Collection: COPS Extension Request Form.
3. The agency form number: None. U.S. Department of Justice, Community Oriented Policing Services (COPS) Office.
4. Affected public who will be asked or required to respond, as well as a brief abstract: Law enforcement agencies and other COPS grants recipients that have grants expiring within 90 days of the date of the form/request. The extension request form will allow recipients of COPS grants the opportunity to request a "no-cost" time extension in order to complete the federal funding period and requirements for their grant/cooperative agreement award. Requesting and/or receiving a time extension will not provide additional funding.

5. An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond: It is estimated that approximately 2,700 respondents annually will complete the form within 30 minutes.
6. An estimate of the total public burden (in hours) associated with the collection: 350 total annual burden hours (0.5 hours x 2700 respondents + 1,350 total burden hours).

If additional information is required contact: Jerri Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE., 3E.405B, Washington, DC 20530.

Dated: July 14, 2015.

DEPARTMENT OF JUSTICE

[OMB Number 1123–0011]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Revision and Extension of a Currently Approved Collection; Department of Justice Equitable Sharing Agreement and Certification

AGENCY: Asset Forfeiture and Money Laundering Section, Department of Justice.

ACTION: 60-day notice.

SUMMARY: The Department of Justice (DOJ), Criminal Division, Asset Forfeiture and Money Laundering Section, will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995.

DATES: Comments are encouraged and will be accepted for 60 days until September 15, 2015.

FOR FURTHER INFORMATION CONTACT: If you have additional comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Jennifer Bickford, Acting Assistant Deputy Chief, Asset Forfeiture and Money Laundering Section, 1400 New York Avenue NW., Washington, DC 20005 (phone: 202–514–1263).

SUPPLEMENTARY INFORMATION: Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Bureau of Justice