Drug Enforcement Administration
Reports for Regulated Machines

Web Tracking No.:

Federal Regulations require a regulated person to submit a detailed report of any Import or Export of tableting or encapsulating machines. A tableting machine or encapsulating machine may not be imported or exported until a DEA transaction identification has been filed by the Administration. Federal regulations require a regulated person to submit a detailed report of all domestic regulated transactions in a tableting machine or encapsulating machine.

1. **Type of Request:**	OMB Approval No. 1117-0024 Expiration Date: 06/30/2018

2. **Type of Submission:** [ ] ORIGINAL [ ] AMENDED [ ] WITHDRAWAL

3. Purpose Need: [ ] Medical [ ] Commercial [ ] Scientific [ ] Other (please describe):

4. **Proposed Date of Export:** **Anticipated Foreign Port of Entry:** **Country of Entry:**	**DEA Transaction ID:**
5. **Regulated Person:** (Business Name, Business Address): Registration Number (required if registered): POC Name: Email address: Business Phone:	6. **Intermediate Consignee – *if any*:** (Business Name, Business Address): Country: POC Name: Email address: Business Phone:

7**. Consignee:** (Business Name, Business Address):

Country:
POC Name:
Email address:
Business Phone:

E-SIGNATURE OF AUTHORIZED INDIVIDUAL:	DATE:

PRIVACY ACT INFORMATION

AUTHORITY: Title 21 U.S.C. 802 and 830

PURPOSE: Report of Regulated Machines.

ROUTINE USES: The Controlled Substances Act authorizes the production of special reports required for statistical and analytical purposes. Disclosures of information from this system are made to the following categories of users for the purposes stated:

A. Other Federal law enforcement and regulatory agencies for law enforcement and regulatory purposes.

B. State and local law enforcement and regulatory agencies for law enforcement and regulatory purposes.

EFFECT: Failure to report may result in penalties under Section 402 and 403 of the Controlled Substances Act.

Under the Paperwork Reduction Act, a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

Public reporting burden for this collection of information is estimated to average 15 minutes per response for imports, exports, and domestic transactions, and 5 minutes per response for Return Declarations, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to the Drug Enforcement Administration, Attn: Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, VA 22152; and to the Office of Management and Budget, Paperwork Reduction Project No. 1117-0024, Washington, D.C. 20503.

Freedom of Information: Please prominently identify any confidential business information per 28 CFR 16.8(c) and Exemption 4 of the Freedom of Information Act (FOIA). In the event DEA receives a FOIA request to obtain such information, DEA will give written notice to the registrant to obtain such information. DEA will give written notice to the registrant to allow an opportunity to object prior to the release of information.

8. **Description of Each Machine** **DEA Transaction ID:**

Machine Type: [ ] Encapsulating [ ] Tableting Serial #: Make: Model: [ ] Electric [ ] Manual Description:	**Machine Type:** [ ] Encapsulating [ ] Tableting Serial #: Make: Model: [ ] Electric [ ] Manual Description:	**Machine Type:** [ ] Encapsulating [ ] Tableting Serial #: Make: Model: [ ] Electric [ ] Manual Description:
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