

CONTROLLED SUBSTANCES IMPORT / EXPORT DECLARATION

(Read Instructions on reverse before completing)

OMB APPROVAL
No. 1117 - 0009

See reverse for Privacy Act

1. CHECK ONE <input type="checkbox"/> IMPORT DECLARATION Nonnarcotic Substances in Schedules III, IV, V <input type="checkbox"/> EXPORT DECLARATION Nonnarcotic Substances in Schedules III, and IV and all substances in Schedule V	BROKER OR FORWARDING AGENT, IF USED (Name and Address)	U.S. CUSTOMS CERTIFICATION Date of Departure / Arrival Name of Carrier / Vessel Date of Certification Signature of Customs Official
IMPORTER/EXPORTER (Name and Address) DEA REGISTRATION NO.		

2. CONTROLLED SUBSTANCES TO BE IMPORTED OR EXPORTED		
2a. NAME AND QUANTITY OF DRUG or PREPARATION <i>(Enter names as shown on labels; numbers and sizes of packages; strength of tablets, capsules, etc., CSA Drug Code and NDC Number)</i>	2b. CONTROLLED SUBSTANCE CONTENT OF DRUG OR PREPARATION expressed as acid, base or alkaloid. <i>(Enter names of controlled substances contained in the drug; compound, or preparation)</i>	2c. DATE IMPORTED/EXPORTED AND ACTUAL QUANTITY <i>(Completed by registrant at time of transaction)</i>

3. <input type="checkbox"/> FOREIGN <input type="checkbox"/> DOMESTIC PORT OF EXPORTATION (last U.S. Customs Port) AND APPROX. DEPARTURE DATE	<input type="checkbox"/> FOREIGN <input type="checkbox"/> DOMESTIC PORT OF IMPORTATION (first U.S. Customs Port) AND APPROX. ARRIVAL DATE
4. MODE OF TRANSPORT; NAME OF VESSEL / CARRIER (if known)	NAME OF ALL INTERMEDIATE CARRIERS
5. NAME AND ADDRESS OF FOREIGN CONSIGNEE/CONSIGNOR	

I hereby certify that the above named substance(s) to be Imported, Exported, are intended for Legitimate medical need, Scientific research, Other (If intended for reexport beyond the country of destination described in block 5 above, attach documentation per Title 21, CFR 1312.27.)

If used as "Export Declaration", attach documentation that importation is not contrary to the laws or regulations of the country of destination.

SIGNATURE OF AUTHORIZED INDIVIDUAL OF IMPORTER/EXPORTER, BROKER OR FORWARDING AGENT	DATE	NAME OF FIRM AND TELEPHONE NUMBER
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DEA REGISTRATION NO.		

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<p>IMPORTER/EXPORTER (Name and Address)</p>	<p>DATE OF DEPARTURE / ARRIVAL</p>
<p>DEA REGISTRATION NO.</p>	<p>NAME OF CARRIER / VESSEL</p>
<p>2. CONTROLLED SUBSTANCES TO BE IMPORTED OR EXPORTED</p>	
<p>2a. NAME AND QUANTITY OF DRUG or PREPARATION <i>(Enter names as shown on labels; numbers and sizes of packages; strength of tablets, capsules, etc., CSA Drug Code and NDC Number)</i></p>	<p>2b. CONTROLLED SUBSTANCE CONTENT OF DRUG OR PREPARATION expressed as acid, base or alkaloid. <i>(Enter names of controlled substances contained in the drug; compound, or preparation)</i></p>
<p>2c. DATE IMPORTED/EXPORTED AND ACTUAL QUANTITY <i>(Completed by registrant at time of transaction)</i></p>	
<p>BROKER OR FORWARDING AGENT, IF USED (Name and Address)</p>	
<p>DATE OF CERTIFICATION</p>	
<p>SIGNATURE OF CUSTOMS OFFICIAL</p>	

<p>3. <input type="checkbox"/> FOREIGN <input type="checkbox"/> DOMESTIC PORT OF EXPORTATION (last U.S. Customs Port) AND APPROX. DEPARTURE DATE</p>	<p><input type="checkbox"/> FOREIGN <input type="checkbox"/> DOMESTIC PORT OF IMPORTATION (first U.S. Customs Port) AND APPROX. ARRIVAL DATE</p>
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- IMPORT DECLARATION Nonnarcotic Substances in Schedules III, IV, V
- EXPORT DECLARATION Nonnarcotic Substances in Schedules III, and IV and all substances in Schedule V

U.S. CUSTOMS CERTIFICATION

Date of Departure / Arrival

Name of Carrier / Vessel

Date of Certification

Signature of Customs Official

IMPORTER/EXPORTER (Name and Address)

DEA REGISTRATION NO.

BROKER OR FORWARDING AGENT, IF USED (Name and Address)

2. CONTROLLED SUBSTANCES TO BE IMPORTED OR EXPORTED

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2c. DATE IMPORTED/EXPORTED AND ACTUAL QUANTITY
(Completed by registrant at time of transaction)

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DATE

NAME OF FIRM AND TELEPHONE NUMBER

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INSTRUCTIONS FOR COMPLETING FORM DEA-236

This form is to be used in notifying DEA of all Imports or Exports as required by Title III, PL91-513, Sections 1002 and 1003, as amended, (Controlled Substances Import and Export Act, 21 U.S.C. 952 and 953). This form may be prepared and signed by the actual Importer or Exporter or by the Broker or Forwarding Agent used. The following instructions supplement the parts of the DEA-236 which are not completely self-explanatory.

Part 1. "IMPORTER" means the authorized DEA registrant who receives the controlled substance; "EXPORTER" means the authorized DEA registrant who ships the controlled substance.

Part 2. Typical entries might read:

Strength:	10 mg tablets
Size or	1,000 tablets/bottle
Weight (Bulk):	100 kilo/drum
Quantity:	100 bottles, 2 drums

If needed, use additional forms and distribute in the prescribed manner after the required documents are attached to each copy.

Part 3. Self-explanatory.

Part 4. Insert name of vessel or airline and flight number, together with all intermediate carriers. Furnish all information concerning the transportation of the goods known at the time of preparing form DEA-236.

Part 5. Enter DEA registration number, if known, for "Import Declaration", or foreign registration number, if applicable, for "Export Declaration".

INSTRUCTIONS FOR DISTRIBUTING FORM DEA-236

If this form is prepared as a **Controlled Substance Import Declaration**, distribute as follows:

Copies 1, 2, and 3 must be forwarded to the foreign shipper. These copies will accompany the shipment to certain points.

Upon receipt of **Copies 1, 2, and 3**, the foreign shipper will present **Copy 1** to the proper foreign government agency or authority, if required, as a prerequisite to export authorization. **Copy 1** shall then accompany the shipment to its final destination and shall be retained in the files of the importer for a period of at least two years.

Copy 2 shall be detached by the customs official at the foreign port.

Copy 3 shall be removed by an official of the United States Customs Service at the port of entry, certified and signed by the customs official (after noting any discrepancies), and forwarded to the Drug Enforcement Administration, Office of Diversion Control, International Drug Unit (ODOI), Washington, D.C. 20537.

Copy 4 must be forwarded at least 15 days prior to importation to the Drug Enforcement Administration, Office of Diversion Control, International Drug Unit (ODOI), Washington, D.C. 20537.

Copy 5 must be retained by the importer until receipt of **Copy 1**.

If this form is prepared as a **Controlled Substance Export Declaration**, distribute as follows:

Copies 1, 2, and 3 shall accompany the shipment to certain points.

Copy 1 shall remain with the shipment to its final destination.

Copy 2 shall remain with the shipment, to be detached and retained by the customs official of the foreign port of importation.

Copy 3 shall be removed by an official of the United States Customs Service at the domestic port of exportation, certified and signed by the customs official (after noting any discrepancies), and forwarded to the Drug Enforcement Administration, Office of Diversion Control, International Drug Unit (ODOI), Washington, D.C. 20537.

Copy 4 shall be forwarded at least 15 days prior to exportation to the Drug Enforcement Administration, Office of Diversion Control, International Drug Unit (ODOI), Washington, D.C. 20537. In cases where the 15 day notice cannot be given, a special waiver may be requested from the Administration.

Copy 5 shall be retained by the exporter as part of his records for a period of at least two years.

PRIVACY ACT INFORMATION

AUTHORITY: Sections 1002 and 1003 of the Controlled Substances Act of 1970 (PL91-513), as amended (21 U.S.C. 952 and 953).

PURPOSE: To obtain information regarding the importation of nonnarcotic substances in Schedules III, IV, and V and the exportation of nonnarcotic substances in Schedules III and IV and all substances in Schedule V.

ROUTINE USES: The Controlled Substances Import/Export Declaration produces special reports as required for statistical and law enforcement purposes. Disclosure of information from this system are made to the following categories of users for the purposes stated.

- Other Federal law enforcement and regulatory agencies for law enforcement purposes.
- State and local law enforcement and regulatory agencies for law enforcement and regulatory purposes.

EFFECT: Failure to complete this form will preclude requested importation or exportation of the referenced controlled substances.

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, 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, FOI and Records Management Section, Washington, D.C. 20537; and to the Office of Management and Budget, Paperwork Reduction Project No. 1117-0009, Washington, D.C. 20503.