

U.S. DEPARTMENT OF JUSTICE – DRUG ENFORCEMENT ADMINISTRATION
APPLICATION FOR PERMIT TO EXPORT CONTROLLED SUBSTANCES
FOR SUBSEQUENT REEXPORT
PURSUANT TO SECTION 1003(f), Title III, PL 109-57
(See Instructions and Privacy Act Information)

DATE:		EXPORTER APPLICATION NUMBER (If applicable)	
1. NAME OF CONSIGNEE IN FIRST COUNTRY		2. ADDRESS OF CONSIGNEE IN FIRST COUNTRY	
3. BUSINESS OF CONSIGNEE IN FIRST COUNTRY		4. FOREIGN PORT OF ENTRY (City & Country)	
5a. PORT OF EXPORTATION (City & state of last U.S. Customs port)	5b. NAME OF EXPORTING CARRIER OR VESSEL (Air, Ship)	5c. APPROX. DATE OF EXPORTATION	
6. FOREIGN IMPORT LICENSE OR PERMIT NO.		ISSUE DATE:	EXPIRE DATE:
7a. NAME AND QUANTITY OF DRUG OR PREPARATION TO BE EXPORTED (Enter names as shown on labels; numbers and sizes of packages; bulk or tablets/capsules, strength of tablets, capsules, etc. CSA Drug Code, and NDC Number)	7b. CONTROLLED SUBSTANCE CONTENT OF DRUG OR PREPARATION TO BE EXPORTED expressed as acid, base, or alkaloid. (Enter name of controlled substance contained in the drug, compound, or preparation.)	7c. DATE EXPORT OCCURED AND ACTUAL QUANTITY (Completed and signed by registrant at time of export and returned within 30 days to DEA.)	
		DEA PERMIT NO:	
		DATE ACTUALLY SHIPPED:	
		SIGNATURE OF RESPONSIBLE COMPANY OFFICIAL:	
8a. NAME OF CONSIGNEE IN SECOND COUNTRY	8b. ADDRESS OF CONSIGNEE IN SECOND COUNTRY	8c. AMOUNT TO BE REEXPORTED TO SECOND COUNTRY (Enter name of controlled substances (7b) and net weight in gms)	

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**APPLICATION FOR PERMIT TO EXPORT CONTROLLED SUBSTANCES
FOR SUBSEQUENT REEXPORT (page 2)**

9a. NAME OF CONSIGNEE IN SECOND COUNTRY	9b. ADDRESS OF CONSIGNEE IN SECOND COUNTRY	9c. AMOUNT TO BE REEXPORTED TO SECOND COUNTRY (Enter name of controlled substances (7b) and net weight in gms)
10a. NAME OF CONSIGNEE IN SECOND COUNTRY	10b. ADDRESS OF CONSIGNEE IN SECOND COUNTRY	10c. AMOUNT TO BE REEXPORTED TO SECOND COUNTRY (Enter name of controlled substances (7b) and net weight in gms)
11a. NAME OF CONSIGNEE IN SECOND COUNTRY	11b. ADDRESS OF CONSIGNEE IN SECOND COUNTRY	11c. AMOUNT TO BE REEXPORTED TO SECOND COUNTRY (Enter name of controlled substances(7b) and net weight in gms)

PLEASE ATTACH ADDITIONAL SHEETS OF FORM DEA 161-R OR DOCUMENTATION PER TITLE 21 CFR 1312.22.

AFFIDAVIT

To the best of my knowledge and belief (1) both the first country to which the controlled substance(s) are exported from the United States and the second country to which the controlled substances are exported are parties to the Single Convention on Narcotic Drugs, 1961, and the Convention on Psychotropic Substances, 1971; (2) the first and second countries have each instituted and maintain a system of controls of imports in conformity with such Conventions; (3) the drugs will be consigned to a holder of such permits or licenses as may be required in the country of import and that a permit or license for importation will be issued for such import into the second country; (4) **that the controlled substances will be reexported from the first country to the second country no later than 90 days after exportation from the United States;** (5) the packages are labeled in conformance with the Single Convention on Narcotic Drugs, 1961 and the Convention on Psychotropic Substances, 1971, and any amendments to these treaties; (6) the controlled substances are to be applied exclusively to medical, scientific, or other legitimate uses within the second country; and (7) the controlled substances will not be exported from the second country.

NAME OF EXPORTER		ADDRESS OF EXPORTER
EXPORTER'S TELEPHONE NO.	EXPORTER'S DEA REGISTRATION NO.	PRINTED NAME & SIGNATURE AND TITLE OF PERSON MAKING APPLICATION

NOTICE: Controlled Substances may not be exported by mail or parcel post.

DEA USE ONLY	APPROVED EXPORT PERMIT NUMBER	DATE EXPORT PERMIT NUMBER ISSUED
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INSTRUCTIONS AND INFORMATION, FORM DEA-161-R

This application must be completed in triplicate. The original is sent to DEA HQs: Drug Enforcement Administration, Office of Diversion Control, Import/Export Unit, Washington, D.C. 20537. See Instructions 5a and 9 for copies two and three.

- (1) The name and address of the consignee as shown on this application and on the permit to export must correspond with that shown on the foreign import certificate.
- (2) To avoid delays in clearance at the port of export, be sure to enter the correct port on this application. A copy of your export permit is sent directly to the District Director of Customs at the port indicated on the application for comparison with the permit presented for clearance of the shipment. The shipment will not clear at any other port without an amendment of the permit indicating a change to that effect.
- (3) The original or an authentic signed and/or notarized copy of the foreign import certificate must accompany this application. If this certificate is in a foreign language, a translation must accompany the application. If this certificate is needed to accomplish entry of the drug into the country of destination, your request for its return to you should accompany the application.
- (4) Application should be made in the name of the registered legal entity, as shown on the DEA registration certificate, and signed by a responsible authorized official if a corporation, by a partner, or by the person registered as an individual. Only persons registered to export may be issued export permits. The registrations of manufacturers, distributors, practitioners, researchers, etc. do not entitle them to export controlled substances.
- (5) Controlled substances in Schedule I or II, or a narcotic drug in Schedule III or IV may be exported from the United States to a country for subsequent export from that country to another country:
 - a. Thirty days from the date of exportation from the United States to the first country, the exporter must return Copy 2 of Form DEA 161-R with 7c completed, that is, with the actual date of export and actual quantity shipped.
 - b. The controlled substances must be exported from the first country to the second country no later than 90 days from the date of exportation from the United States
 - c. Within 30 days after the controlled substance is exported from the first country to the second country or the order is canceled by the second country, the person who exported the controlled substance from the United States must deliver to DEA Headquarters documentation certifying that such export from the first country to the second country has occurred or was refused.
 - i. The company must provide on company letterhead signed by the responsible company official the following information: (1) Name of the second country, (2) actual quantity shipped, (3) actual date shipped, and (4) DEA export permit number for the original export to the first country.
 - ii. For refused shipments, the company must file a written request with DEA for return, a brief summary of the facts warranting the return, and a DEA Form 357, Application for Import Permit. DEA will evaluate the request and return a response in writing.
- (6) Permits will be mailed to the exporter at the address shown at the bottom of the application unless contrary instructions are attached to and made a part of this application.
- (7) Identification of drugs to be exported and the controlled substance content should be entered on the application in the following manner:

8a. NAME AND QUANTITY OF DRUG OR PREPARATIONS TO BE EXPORTED	8b. CONTROLLED SUBSTANCE CONTENT OF DRUG OR PREPARATION TO BE EXPORTED (expressed as acid, base or alkaloid, not salt)
3 bottles x 100 Secobarbital Sodium capsules (100 mg/capsule)	Secobarbital 24.47 gm
2 boxes x 100 Meperidine HCL ampules (5%, 2ml ampules)	Meperidine 17.43 gm
1 box x 100 Meperidine HCL vials (10%, 20 ml, vials)	Meperidine 174.30 gm
2 x 1 pt. Meperidine HCL Syrup (50 mg/5ml, pints)	Meperidine 8.24 gm
1 box x 100 gm Dextroamphetamine Sulfate powder	Dextroamphetamine 73.38 gm.
1 bottle x 500 Hydromorphone HCl tablets (4 mg/tablets)	Hydromorphone 1.77 gm

- (9) Copy 3 of the Application for Permit to Export Controlled Substances for Subsequent Reexport (Form DEA 161-R) is retained by the registrant.

PRIVACY ACT INFORMATION

AUTHORITY: Section 1003 of the Controlled Substances Act of 1970 (PL-513)

PURPOSE: Control exportation of certain Controlled Substances from the United States.

ROUTINE USES: The Controlled Substances Act Registration Records produces special reports as required for statistical analytical purposes. Disclosures of information from this system are made to the following categories of users for the purposed states:

- 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.
- C. Persons registered under the Controlled Substances Act (Public Law 91-513).

EFFECT: No permit will be issued.

According to the Paperwork Reduction Act of 1995, no persons are required to respond to a Collection of Information unless it displays a valid OMB control number. The valid OMB control number for this Information Collection is 1117-0004. The time required to complete this information collection is estimated to average 15 minutes per response, including the time to review instruction, search existing data resources, gather the data needed, and complete and review the information collection.

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