

SEE INSTRUCTIONS ON REVERSE SIDE	No procurement quota may be issued unless a completed application form has been received, 21 CFR 1303.12(b).	OMB APPROVAL No. 1117 - 0008
1. NAME OF BASIC CLASS DESIRED (<i>Only one basic class per DEA-250</i>)		2. SCHEDULE NUMBER
4. NAME AND ADDRESS OF REGISTRANT (<i>Include No., Street, City, State and ZIP Code</i>)		3. DEA DRUG CODE NUMBER
		5. YEAR FOR WHICH QUOTA IS REQUESTED
		6. DEA REGISTRATION NUMBER
7. NAME OF CONTACT PERSON	8. TELEPHONE NO. (<i>Include ext., if applicable</i>)	9. FAX NUMBER

NOTE: All quantities are to be expressed in grams of anhydrous acid, base or alkaloid (not as salts).

10. QUOTA HISTORY	QUOTAS PREVIOUSLY ISSUED BY DEA			QUOTA REQUESTED
	2nd PRECEDING YEAR	1st PRECEDING YEAR	CURRENT YEAR	
	_____ Grams	_____ Grams	_____ Grams	_____ Grams

11. PRODUCTION DATA	2nd PRECEDING YEAR	1st PRECEDING YEAR	ESTIMATE FOR CURRENT YEAR	ESTIMATE FOR YEAR FOR WHICH QUOTA IS REQUESTED
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I. INVENTORY AS OF DEC. 31				
a. Bulk controlled substance				
b. In-process material				
c. Contained in FINISHED Dosage Forms				
TOTAL (a + b + c)				

II. DISPOSITION / UTILIZATION				
a. Domestic				
b. Exports				
TOTAL (a + b)				

III. ACQUISITION / PRODUCTION				
a. Domestic Sources				
b. Importation				
TOTAL (a + b)				

12. IF THE PURPOSE IS TO MANUFACTURE ANOTHER SUBSTANCE (S), FURNISH THE FOLLOWING INFORMATION:					
NAME OF NEW SUBSTANCE	DEA DRUG CODE NUMBER	AMOUNT USED FOR THIS PURPOSE			% YIELD (Historical)
		2nd PRECEDING YEAR	1st PRECEDING YEAR	CURRENT YEAR	

13. IF THE PURPOSE IS TO MANUFACTURE THE BASIC CLASS INTO DOSAGE FORMS, FURNISH THE FOLLOWING INFORMATION:					
NAME OF DOSAGE FORM	SCHE- DULE	AMOUNT USED FOR THIS PURPOSE		ESTIMATE FOR CURRENT YEAR	ESTIMATE FOR YEAR FOR WHICH QUOTA IS REQUESTED
		2nd PRECEDING YEAR	1st PRECEDING YEAR		

SIGNATURE OF APPLICANT	PRINT or TYPE NAME and TITLE of SIGNER	DATE
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**INSTRUCTIONS FOR COMPLETING THE DEA FORM 250:
Application For Procurement Quota**

The DEA-250 must be filed on or before April 1 of the year preceding the calendar year for which the procurement quota is being applied. Regulations governing quotas are included in Title 21, Code of Federal Regulations, Part 1300 to end. Copies of these regulations may be ordered from: The Government Printing Office, Superintendent of Documents, Attn: New Orders, P.O. Box 371954, Pittsburgh, PA 15250-7954. Submit the completed form to:

Drug Enforcement Administration
Office of Diversion Control
Drug & Chemical Evaluation Section (ODE)
Washington, D.C. 20537

The following instruction is for that item which is not completely self-explanatory.

Item 11(I). This is to include all factory and branch stocks which have reached that point in manufacturing as to be identifiable, whether in bulk form, in the process of manufacture, in finished form, or otherwise (e.g., damaged, defective, or impure substances awaiting disposal, substances held in quarantine, or substances maintained for extemporaneous compounding), as a basic class of controlled substance manufactured or otherwise acquired by a registrant, whether in bulk, commercial containers, or contained in pharmaceutical preparations in the possession of the registrant (including stocks held by the registrant under separate registration as a manufacturer, importer, exporter or distributor).

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 1 hour 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 FOI and Records Management Section, Drug Enforcement Administration, Washington, D.C. 20537; and to the Office of Management and Budget, Paperwork Reduction Project No. 1117-0008, Washington, D.C. 20503