Online Quota System Year-End Reporting System

Diversion Control Division
United Nations Reporting and Quota Section (DRQ)
Supply Chain Conference
Houston, TX

Online Quota Applications



Quota Applications Must be Submitted Online

Benefits

- Provides secure web-based database system
- Verifies registration as active and current
- Preloads your available drug codes, historical data
- Performs calculations
- Provides confirmation of agency receipt and status
- Expedites process

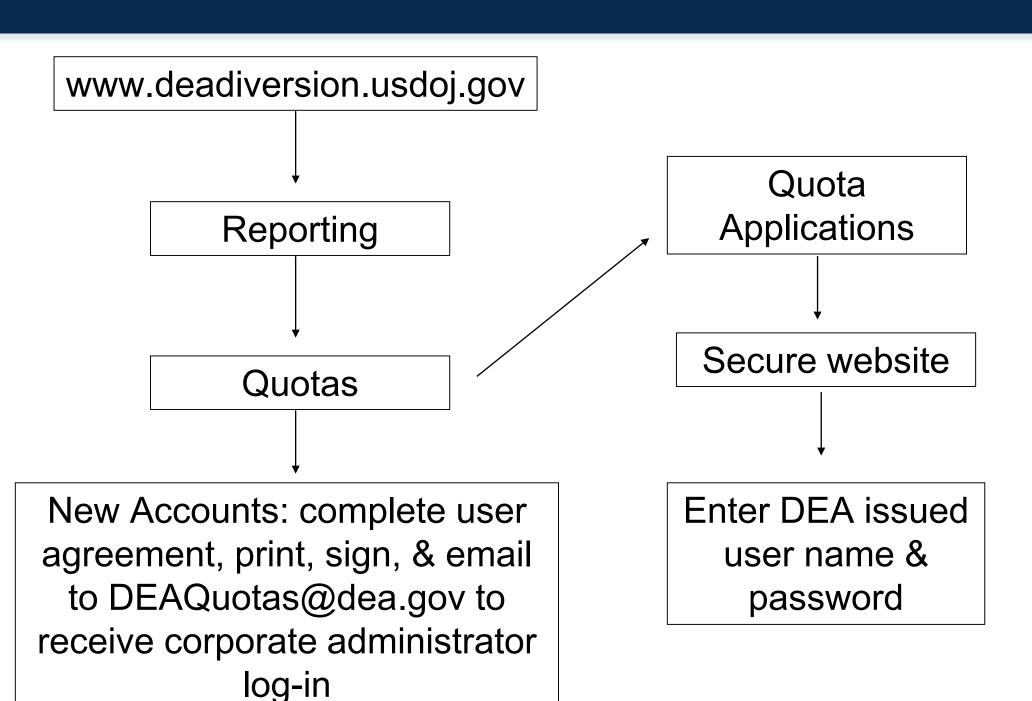
Code of Federal Regulations Requirements for Submitting Quota Requests



- Procurement applications due by April 1
- Import applications due by April 1
- Manufacturing applications due by May 1
- Adjustments to any quota can be requested throughout the year
 - Avoid duplicate entries, request withdrawal of erroneous entries
- Quantities should be expressed as:
 - Anhydrous acid, base or alkaloid
 - GRAMS, rounded to the nearest WHOLE gram

Accessing the Online Quota System





Corporate Administrator Login





Create New Corporate User Account



Corporate Administrator User Administration

You are the Corporate Administrator. These are the corporate users accounts for DEA Number

Create a New User Account

New User

Reset User Password

O USER ACCT 1

Delete User Account

USER ACCT 1

Click Back to return to the User Login screen. Click Next to process your selection.

Back

Next

Corporate Administrator Create New User rised of at least one, but not more the ten digits and the underscore ch

A password must be comprised of at least one, but not more than ten uppercase or lowercase letters, any of the ten digits and the underscore character, arranged in any order.

DEA Number:

Username:

New Password:

Confirm New Password:

Click Cancel to abort this action and return to the User Administration screen. Click Next to process your selection.

Cancel

Reset

Next

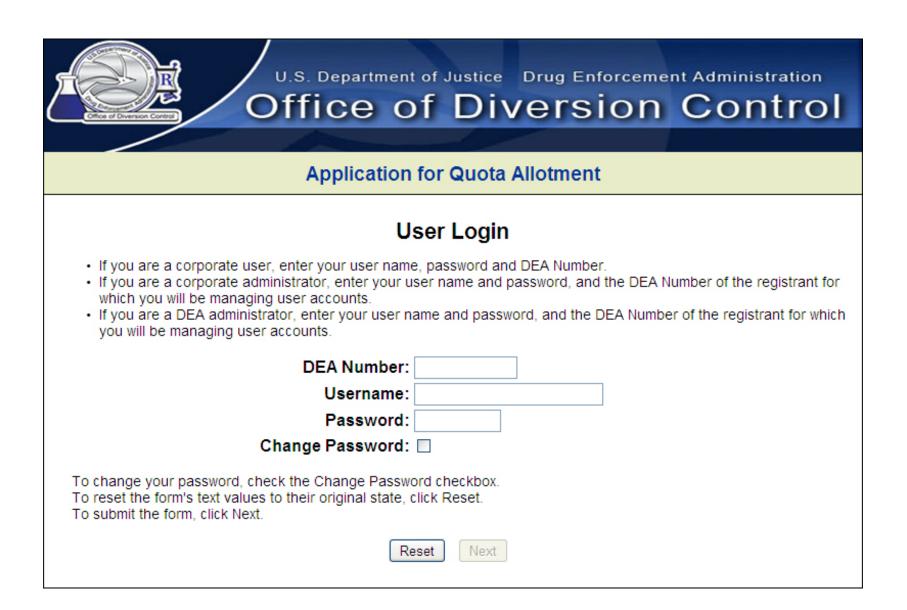
Corporate Administrator User Administration Confirmation

You have successfully created a user account for

Next

Corporate User Login





Corporate User Login Confirmation





Application for Quota Allotment

Login Confirmation

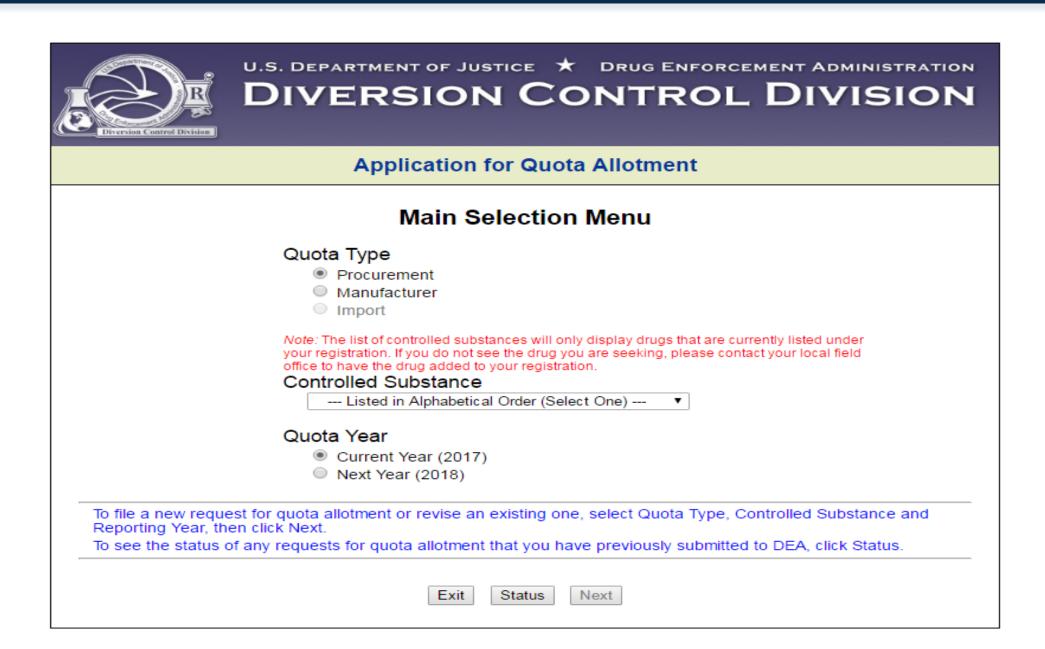
- You are logged in as
- As a Corporate User of the Quota Allotments system, you are authorized to submit requests for Quota Allotments for DEA Number (DEA MANUFACTURER TEST ACCOUNT).
- If this information is not correct, please contact your administrator immediately.



Click Return to return to the User Login screen. Click Next to advance to the next screen.

Report Selection Menu





Online Quota Application



Must request specific isomers for:

- Amphetamine
 - d-amphetamine
 - I-amphetamine
 - d,l-amphetamine
- Methamphetamine
 - d-methamphetamine
 - I-methamphetamine
 - d,l-methamphetamine
- Tetrahydrocannabinols
 - Delta-9-THC
 - All Other Tetrahydrocannabinols

 As of 2022, Methylphenidate and Dexmethylphenidate must be applied for separately

Substance Selection Identifiers



"For Sale" vs. "For Conversion"

<u>For Sale</u>: substance used to manufacture products containing same basic class which is bought from the bulk manufacturer.

- involves ~95% of the quota applications

ex: Codeine (for sale) codeine/APAP

For Conversion: substance used to synthesize a different basic class.

ex: Codeine (for conversion) \longrightarrow hydrocodone



Examples of "For Sale" vs. "For Conversion"

- Codeine
- D-amphetamine
- D-methamphetamine
- Dexmethylphenidate
- Diphenoxylate
- Hydrocodone

- Methylphenidate
- Morphine
- Noroxymorphone
- Oxycodone
- Oxymorphone

Procurement Quota Application





Application for Quota Allotment

Year-End Estimates for Reporting Year 2017

Current Selection

Quota Type	Quota Year	Drug Code	Drug Name	Pending Quota	Approved Quota
Procurement	2017	9193	HYDROCODONE (FOR SALE)		.000

DEA Form 250 -- Box 12

Enter your estimated year-end report projections in the fields provided.

Use the TAB and SHIFT-TAB keys to tab forward and backward among the fields.

All amounts are expressed as grams of anhydrous base or alkaloid, not as salt.

Reporting Year	2015 (actual)	2016 (actual)	2017 (estimate)			
Inventory (as of Dec 31)						
Bulk controlled substance (API)	0.000	0.000	0.000			
In-process material (in all forms)	0.000	0.000	0.000			
Finished dosage forms (bulk and/or packaged)	0.000	0.000	0.000			
Total	0.000	0.000	0.000			
Dispositions						
Domestic	0.000	0.000	0.000			
Exports	0.000	0.000	0.000			
Total	0.000	0.000	0.000			
Acquisitions						
Domestic	0.000	0.000	0.000			
Imports	0.000	0.000	0.000			
Total	0.000	0.000	0.000			

Click Next to advance to the next page.

Reset Re

Return



Quota Allotment Categories



Commercial Manufacturing

- Conversion to other substances
- Dosage form

Product Development

- All stages leading to FDA approval
- Laboratory scale
- Scale up
- Exhibit/Submission/Registration
- Validation
- Stability

Packaging

- Packaging/Repackaging
- Labeling/Relabeling

Replacement

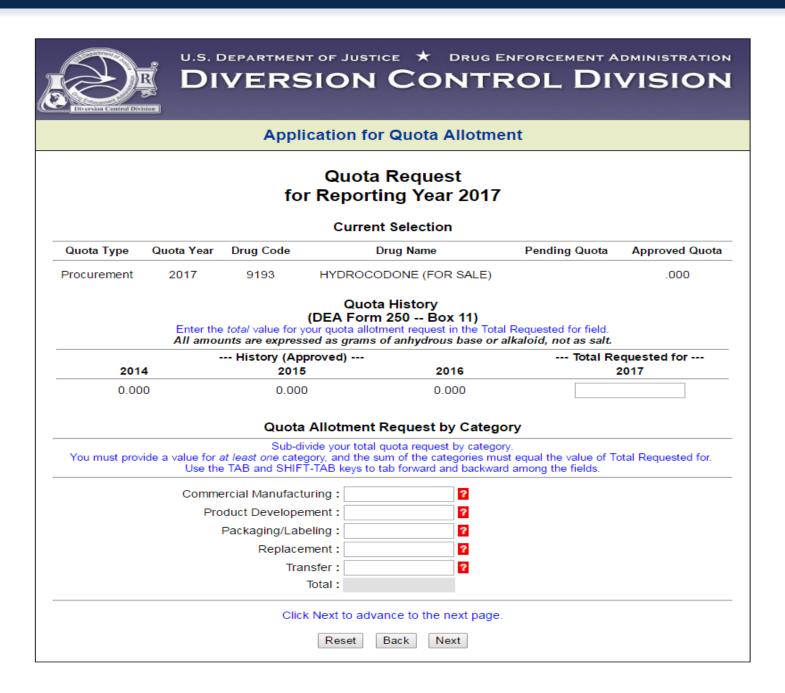
 Case-by-case basis for commercial manufacturing

Transfer

- Bulk API
- In-process material
- Finished dosage units

Quota Allotment Categories





Quota Allotment Categories





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Application for Quota Allotment

Report Selection Menu

Current Selection

Quota Type	Quota Year	Drug Code	Drug Name	Pending Quota	Approved Quota
Procurement	2017	9193	HYDROCODONE (FOR SALE)	1,000.000	.000

Sub-Reports Selection

In order to complete your application for quota allotment, you will need to provide additional information using the sub-reports below. To fill out a sub-report, click the Report Link to advance to the corresponding report form. If the color of the sub-report link is gray, it means there are sub-reports already on file. You may review and/or update these by clicking the link. You will not be able to advance to the next page until all required sub-reports have been completed.

Required / Optional	Current Request	Report Link
Optional	500.000	Commercial Manufacturing <u>Dosage Forms</u> (DEA Form 250 Box 14)
Required	200.000	Product Development (DEA Form 250 Box 14)
Required	100.000	Packaging/Labeling (DEA Form 250 Box 14)
Required	100.000	<u>Replacement</u>
Required	100.000	<u>Transfer</u>

If you have completed all required sub-reports, click Next to continue.

Back

Next

Commercial – Conversions



Basic Class To Be Manufactured (DEA Form 250 -- Box 13)

To convert PHENYLPROPANOLAMINE (FOR CONVERSION) to another substance, select the substance from the list. If the target substances is not in the list, either contact DEA to have the substance added to your registration, or select "Users Choice" (the last entry).

--- Listed in Alphabetical Order (Select One) ---

Drug Name	Drug	2009	2010	2011	% Yield
	Code	(actual)	(actual)	(estimated)	(estimated)

Fill in the fields above and click Submit.

Reset

Submit

Summary

Remove	Drug Name	Drug Code	2009 (actual)	2010 (actual)	2011 (estimated)	% Yield (estimated)
0	Vraxoin	0000	.000	.000	2,524.002	252.24

To remove an entry, click the Remove radio button.

Next

Commercial - Dosage Forms





Product Development





Labeling & Packaging





Replacement





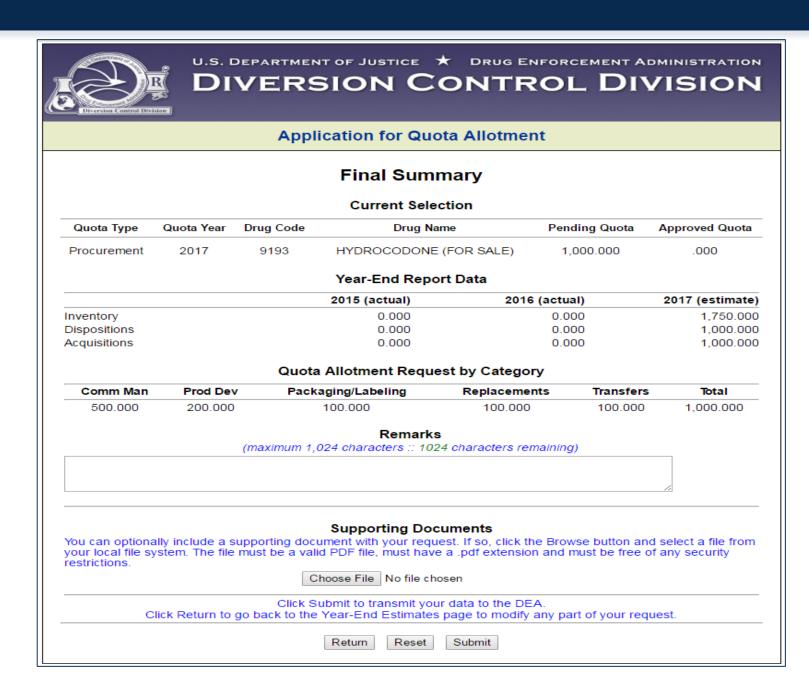
Transfer





Summary





Summary





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Application for Quota Allotment

Summary Confirmation

Current Selection

Quota Type	Quota Year	Drug Code	Drug Name	Pending Quota	Approved Quota
Procurement	2017	9193	HYDROCODONE (FOR SALE)	1,000.000	.000

Confirmation

The Quota Allotment request for the substance above has been submitted successfully.

Reference Number: 121583

Questions regarding this transaction may be emailed to DEA at ODE.quota@usdoj.gov. Please include your DEA registration number, the above reference number and your company name in the subject line of all correspondence regarding this request.

This request may be amended by selecting the same substance from the Main Selection Menu. All data entry fields will be loaded with the values from the latest completed request.

Printing Options

Click the Print button below to print the request or archive it as a PDF report.

Print

Return

Exit

Status Report





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Application for Quota Allotment

Quota Request Status

The following table lists substances for which you have formally requested a quota allotment, and the status of your application. Multiple requests for the same substance are sorted sequentially by Drug Name within Quota Type.

A date in the Date Approved field indicates the date the request was approved. Requests that have not been approved yet show as "Pending."

Requests may be amended by returning to the Main Selection Menu and selecting the substance from the list.

To view the request as a PDF report, click the Request ID link.

Туре	Year	Date Submitted			Quota	Date Approved	Request ID
PRO	2017	Mar 31, 2016	9200	HEROIN	50.000	Pending	121569
		Apr 40		HYDROCODONE (FOR SALE)	1,000.000	Pending	<u>121583</u>

Back

Print

Quota Amendments





Office of Diversion Control

Application for Quota Allotment

Quota Request Amendment

Notice

You have previously filed a quota allotment request for the substance noted above. The request was filed on Mar 02, 2011 in the amount of 585.000 grams. The Reference Number is 108019.

In the pages that follow, your original data values for the request will be displayed. You may freely remove, append or amend this data. Once you complete this amended request, it will be filed with its own reference number and will become the new statement of record for the requested quota allotment for this substance.

Current Selection

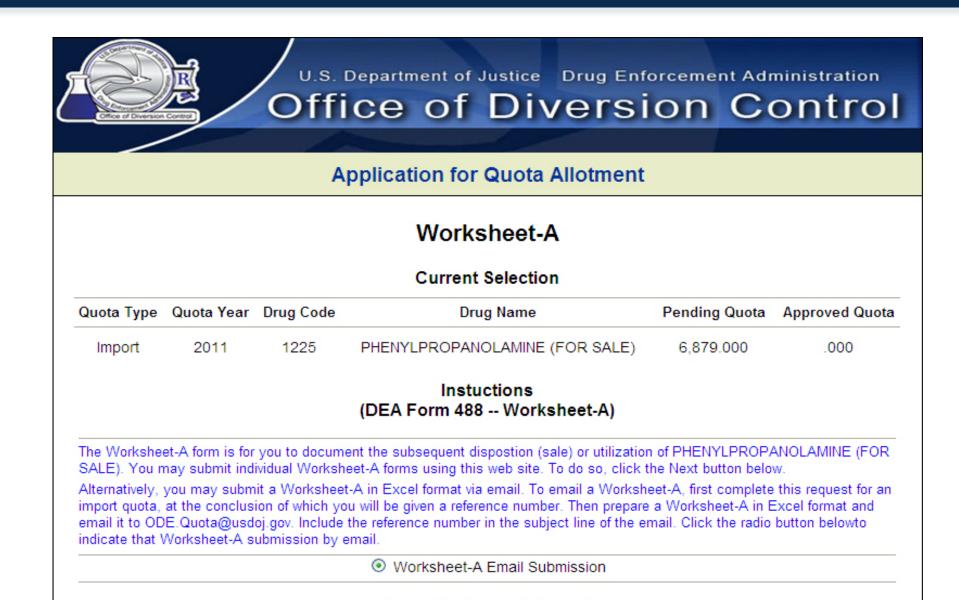
Quota Type	Quota Year	Drug Code	Drug Name	Pending Quota	Approved Quota
Procurement	2011	8113(B)	EPHEDRINE (FOR SALE)	585.000	.000

Back

Next

List 1 Import quota





Return

Next

Reset

List 1 Import quota



			Current Selection		
uota Type	Quota Year	Drug Code	Drug Name	Pending Quota	Approved Quot
Import	2011	1225	PHENYLPROPANOLAMINE (FOR CONVERSION)	12.000	.000
			Worksheet-A Data (DEA Form 488 Worksheet-A)		
⋆ DEA Nun ⋆ Address			★ Name of company		
* Address					~
* Phone:		*	Fax: * Email:		
★ Contact	Person:				
⋆ Quantity	to be sold (a	s grams of	anhydrous base) :		
★ Provide (documentatio	on for Quant	tity to be sold (purchase orders, supply agreement	s, etc):	
				<u>~</u>	
⋆ Intended	Use: OS	ale / Produc	ct Launch O Product Development / Research	Other (explain	n)
he ⋆ symb	ool denotes a	required fie	eld. Number fields must be a valid number between	0 and 999,999,9	999.999 inclusiv
			Fill in the fields above and click Submit.		

List 1 Import quota



Summary

Remove

DEA Num: ####

Company Name: dFGHFYT

Address: HFGGGYYJK,GYFJYYU CTGBD BGFSHDFJKSD

Phone: 256-515-4422

Fax: 121-142-4448

Email: NCBHXJH@GMAIL.COM

Contact: SHJAGYSG

Quantity to be sold: 5,226.000

Supporting docs: GJKEYU

Intended Use: Other

Intended Use Explain: FJDSIOUFRW

To remove an entry, click the Remove radio button.

To return to the Worksheet-A Selection Menu, click Return.

Next

How to Improve Your Applications



- Provide YTD sales in grams base when applying for commercial quota
- If requested, provide customers' DEA#'s and the amount sold to each customer in grams base
- Provide product development in the following format: batch purpose, strength (mg) x Units x batches x conversion factor = total grams base. Please note amounts for overages, stability, retains, and waste.
- Include DEA#, application# on all correspondence
 - POC/direct phone line/extension if different than registration
- Too much or too little information will slow your quota review down
- Do not submit a copy of this/past applications, registration certificate, past quota letters with your supporting documentation
- If you're going to submit a large package of information please provide a summary page that provides a synopsis of your activity and the calculations you used to determine your quota requirements

Year-End Reporting System (YERS)

Pre-YERS History



- ARCOS system was developed to track the distribution of controlled substances
 - Manufacturer >> Distributor >> Various Retail Outlets
- Timeliness of registrants completing reports makes it less suitable for certain DEA functions
- It was decided that ARCOS data could not be used to comply with UN mandates

YERS History



In the early 90's, the DEA created year-end reports to facilitate:

- Establishment & Revision of
 - Aggregate Production Quotas (APQ) CI, CII
 - Assessment of Annual Needs (AAN) CMEA List I Chemicals
 - Individual registrant quotas (IQ,MQ,PQ)
- Monitoring the "80/20" Narcotic Raw Material import rule
- UN-treaty mandated reporting

YERS Benefits



- Provides a secure web-based database system
- Preloads your available drug codes, historical data
- Performs calculations
- Generates error messages for possible incorrect entries or when "specify" fields are not completed
- Moves submitted reports to completed list

Year-end Reporting



Reports need to be completed by:

- DEA REGISTERED MANUFACTURERS
 - Schedule I and II controlled substances
 - Psychotropic drugs on UN's "green list"
- DEA REGISTERED CMEA LIST I CHEMICAL IMPORTERS
 - Ephedrine
 - Pseudoephedrine
 - Phenylpropanolamine

Important Things To Remember



- Due by January 31 after end of reporting calendar Year (CY)
- Separate report required for each substance if
 - Quota was issued for that reporting CY
 - Inventory processed or carried over from an earlier CY
 - Psychotropic drug on UN's 'green list'
 - Newly scheduled CI or CII drugs
- Quantities reported should be expressed as:
 - Anhydrous acid, base or alkaloid
 - -GRAMS

Accessing YERS Online



Www.deadiversion.dea.gov Reporting

Year-End
Reports

Secure
New Accounts: Complete user

Reporting

agreement, print, sign, & email to DEAQuotas@dea.gov to receive corporate administrator log-in

Enter DEA issued username & password

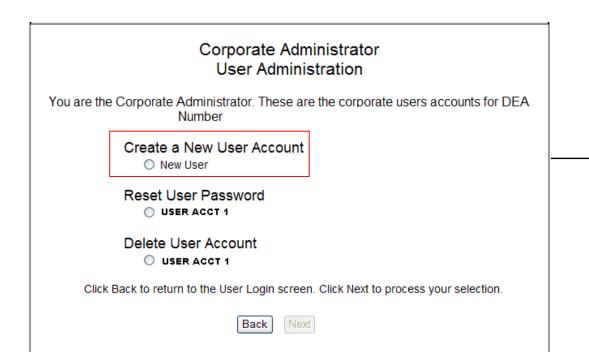
Corporate Administrator Login





Create New Corporate User Account







Corporate Administrator User Administration Confirmation

You have successfully created a user account for

Corporate User Login





Corporate User Login Confirmation





Year-End Reports Login Confirmation

Corporate User Login Confirmation

- You are logged in as
- You are authorized as a corporate user to submit Year-End Inventory Reports for DEA Number
- If this information is not correct, please contact your administrator immediately.

Click Back to return to the User Login screen. Click Next to advance to the Main Selection Menu screen.

Back

Report Selection Menu





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OFFICE OF DIVERSION CONTROL

Year-End Reports

Report Selection Menu

Year-End Report for Procured Substances

- Registered Substances With Quotas and / or Outstanding Inventory
- Other registered substances
- View Null Reports
- Drugs Converted to Other Substances

Year-End Report for Manufactured Substances

- Registered Substances With Quotas and / or Outstanding Inventory
- Other registered substances
- View Null Reports
- Drugs Converted to Other Substances

Year-End Report for Psychotropic Substances

Psychotropic Drugs





Report Selection





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Year-End Report for Procured Substances Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Pending Reports

These are controlled substances or listed chemicals (ephedrine, pseudoephedrine and phenylpropanolamine) (arranged in drug code order) for which you were granted a Procurement quota for reporting year 2016 and / or for which there was outstanding inventory for reporting year 2015. Click the *File* radio button to file a year-end report.

File	Drug Code	Drug Name	Quota Granted (grams)
•	1100	D,L-AMPHETAMINE	N/A
	1100	D-AMPHETAMINE (FOR SALE)	N/A
	1237	METHCATHINONE	N/A
	1724	METHYLPHENIDATE	N/A
	2125	AMOBARBITAL	N/A

Rows 1 through 5 of 6
Use the Back / Forward links to scroll the list. Use the Back / Next buttons below to advance the screen.

<< Back Scroll Forward >>

Completed Reports

These are year-end Procurement reports (arranged in drug code order) that meet the criteria above and that you have already filed for reporting year 2016. Click the *Update* radio button to review or make changes to the report. Click the *Delete* radio button to delete the report.

Update	Delete	Print	Drug Code	Drug Name	Quota Granted (grams)

You have not yet filed any Procurement reports for reporting year 2016.

To scroll the list backward or forward, click one of the Back or Forward links above.

To make a selection, click a radio button in one of the the lists.

To submit your selection for processing, click the Next button below.

To return to the previous page, click the Back button below.

Back

Year-end Reporting – UN Mandated Reports



UN mandated individual reports for the following optical isomers:

- Amphetamine
 - d-amphetamine
 - I-amphetamine
 - d,l-amphetamine
- Methamphetamine
 - d-methamphetamine
 - I-methamphetamine
 - d,l-methamphetamine
- Tetrahydrocannabinols
 - Delta-9-THC
 - All Other Tetrahydrocannabinols

Substance Selection Identifiers



"For Sale" vs. "For Conversion"

<u>For Sale</u>: substance used to manufacture products containing same basic class which is bought from the bulk manufacturer.

- involves ~95% of the quota applications

ex: Codeine (for sale) codeine/APAP

For Conversion: substance used to synthesize a different basic class.

ex: Codeine (for conversion) \longrightarrow hydrocodone



Examples of "For Sale" vs. "For Conversion"

- Codeine
- D-amphetamine
- D-methamphetamine
- Dexmethylphenidate
- Diphenoxylate
- Hydrocodone

- Methylphenidate
- Morphine
- Noroxymorphone
- Oxycodone
- Oxymorphone

Report Selection Confirmation





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Year-End Report for Procured Substances Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Confirmation to File a New Report

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)
2016	1100	D,L-AMPHETAMINE	N/A

All amounts are expressed as grams of anhydrous base or alkaloid, not as salt.

Please note that the quota value for this controlled substance or listed chemical (ephedrine, pseudoephedrine and phenylpropanolamine) reflects your established Procurement quota for reporting year 2016 and does not include any granted replacement quota. If this figure does not agree with your records, please contact <u>ODEQ</u>.





Year-end Reporting: Inventory



- Inventories should include amounts held by the registrant and be reported for each manufacturer, importer, exporter or distributor registration at that registration's <u>physical location</u>
- "Bulk Controlled Substance" is the amount of raw material in inventory
- "In-process" i.e. beads not yet encapsulated
- "Non-saleable Materials" includes any non-saleable inventory i.e. samples, product development

Inventory





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Year-End Report for Procured Substances Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Inventory

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)
2016	1100	D,L-AMPHETAMINE	N/A

All amounts are expressed as grams of anhydrous base or alkaloid, not as salt.

Use the Tab key to navigate between fields.

Inventory (as of December 31)	2015	2016	
Bulk controlled substance (API)	600.000	.000	
In-process material (in all forms)	200.000	.000	
Finished dosage forms (in bulk and/or packaged)	300.000	.000	
Non-saleable material			
retains / stability samples	100.000	.000	
product development material	100.000	.000	
pending destruction	100.000	.000	
other (specify):	.000	.000	
Total non-saleable material	300.000	.000	
Total Ending Inventory (grams)	1,400.000	.000	Sum

Click Return to abandon this report and start over with a new selection.

Click Next to advance to the Acquisitions page.

Return

Year-end Reporting: Acquisitions



 Only material which was obtained against a quota should be reported under "Domestic Procurement" or "Manufactured/ Produced/ Synthesized/ Extracted"

 Material acquired through <u>authorized replacement</u> of destroyed material is reported under "Other Acquisitions to be Accounted For"

Acquisitions



Year-End Report for Procured Substances Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Acquisitions

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)
2016	1100	D,L-AMPHETAMINE	N/A

All amounts are expressed as grams of anhydrous base or alkaloid, not as salt.

Use the Tab key to navigate between fields.

Acquisitions (during the year)	2016	
Domestic procurements, acquisitions, transfers, etc		
bulk API	.000	
non-bulk API material (in process, blends, beads, dosage forms, etc.)	.000	
Total domestic procurements	.000	
Imports		
bulk API	.000	
non-bulk API material (in process, blends, beads, dosage forms, etc.)	.000	
Total imports	.000	
Returns by customer for credit, salvage, rework, etc.		
bulk API	.000	
non-bulk API material (in process, blends, beads, dosage forms, etc.)	.000	
Total returns	.000	
Other acquisitions		
Replacement quota	.000	
other (specify):	.000	
Total other acquisitions	.000	
Total acquisitions (grams)	.000	Sum

Click Back to return to the Inventory page. Click Next to advance to the Dispositions page.

Back

Year-end Reporting: Dispositions



- Material acquired with current year's quota
- Material from previous year's inventory
 - Domestic sales / transfers
 - Exempted products
 - Exports
 - Returns
 - Authorized destructions

Dispositions



Year-End Report for Procured Substances Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Dispositions

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)
2016	1100	D,L-AMPHETAMINE	N/A

All amounts are expressed as grams of anhydrous base or alkaloid, not as salt.

Use the Tab key to navigate between fields.

Dispositions (during the year)	2016	
Domestic Sales or Transfers to:		
Pharmacies, Doctors, Hospitals, Clinics	.000	
Other manufacturers	.000	
Distributors for commercial sale	.000	
Distributors for destruction	.000	
Researchers, analytical labs	.000	
Other federal, state or county agencies	.000	
Total domestic sales or transfers	.000	
Put in process for conversion to bulk substances	.000	
Exports	.000	
Returns from you to your suppliers (specify):	.000	
Quantity used to make exempted or excluded products	.000	
Losses (specify):	.000	
Authorized destructions via DEA Form 41	.000	
Other dispositions (specify):	.000	
Total dispositions (grams)	.000	Sum

Click Back to return to the Acquisitions page.

Click Next to save your work and view a Summary page for this report.

If you wish to put in process for conversion, enter an amount in to textbox and click Next.

Back

Summary Totals





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Year-End Report for Procured Substances Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Summary Totals

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)
2016	1100	D,L-AMPHETAMINE	N/A

All amounts are expressed as grams of anhydrous base or alkaloid, not as salt.

Use the Tab key to navigate between fields.

Total Ending Inventory for 2015	1,400.000
Total Ending Inventory for 2016	.000
Total Acquisitions for 2016	.000
Total To Be Accounted For (Total Ending Inventory For 2015 plus Total Acquisitions For 2016)	1,400.000
Total Dispositions for 2016	.000
Total Accounted For (Total Ending Inventory For 2016 plus Total Dispositions For 2016)	.000
Difference (Total To Be Accounted For minus Total Accounted For)	1,400.000
Please provide an explanation for any difference: (maximum 1,024 characters :: 1024 characters remaining)	
Remarks: (maximum 1,024 characters :: 1024 characters remaining)	

To save this report to permanent storage, click Submit.

To return to the beginning of this report in order to review or change the entered data, click Back.

To return to the Report Selection List, click Return.

Return

Back

Submit

Confirmation





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Year-End Report for Procured Substances

Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Summary Confirmation

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)
2016	1100	D,L-AMPHETAMINE	N/A

You have successfully created the year-end report for the substance noted above. The report selection has been moved to the Completed Reports section of the QA/OI Selection List.

To return to the Report Selection Menu, click Main.
To return to the Report Selection List, click Return.
To exit this application, click Exit.

Main | F

Return

Exit

Completed Reports





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Year-End Report for Procured Substances Drugs for Which a Quota Was Granted or Outstanding Inventory Exists

Pending Reports

These are controlled substances or listed chemicals (ephedrine, pseudoephedrine and phenylpropanolamine) (arranged in drug code order) for which you were granted a Procurement quota for reporting year 2016 and / or for which there was outstanding inventory for reporting year 2015. Click the *File* radio button to file a year-end report.

File	Drug Code	Drug Name	Quota Granted (grams)
	1100	D-AMPHETAMINE (FOR SALE)	N/A
	1237	METHCATHINONE	N/A
	1724	METHYLPHENIDATE	N/A
	2125	AMOBARBITAL	N/A
	2270	PENTOBARBITAL	N/A

Completed Reports

These are year-end Procurement reports (arranged in drug code order) that meet the criteria above and that you have already filed for reporting year 2016. Click the *Update* radio button to review or make changes to the report. Click the *Delete* radio button to delete the report.

Update	Delete	Print	Drug Code	Drug Name	Quota Granted (grams)
		•	1100	D,L-AMPHETAMINE	N/A

You no longer hold a valid registration for the substance outlined in red above. For that reason, your only option is to print a copy of the report.

To scroll the list backward or forward, click one of the Back or Forward links above.

To make a selection, click a radio button in one of the the lists.

To submit your selection for processing, click the Next button below.

To return to the previous page, click the Back button below.

Back

Year-End Report Manufacturing Selection Menu





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Year-End Report for Psychotropic Substances

Psychotropic Drugs





Year-end Reporting



- Narcotic Raw Materials reporting is required by 1961 Single Convention on Narcotic Drugs:
 - Opium
 - Concentrated poppy straw (CPS)
- Alkaloids in CPS which are reported:
 - Codeine
 - Morphine
 - Oripavine
 - Thebaine

Year-end Report for Bulk Manufacturers



- To be completed by a manufacturer who manufactures, produces, synthesizes
 or extracts a bulk Schedule I or II controlled substance
- In-process inventories are to be reported as end-product inventories, not as precursor inventories
 - based on actual assay of in-process material, or
 - based on historical yields

Conversion Reporting



- To be completed by a manufacturer who utilized a basic class for conversion to another controlled or non-controlled substance
 - Example 1 If a manufacturer used phenylacetone (II) to synthesize amphetamine (II), the amount of phenylacetone used and the amount of amphetamine synthesized would be reported on this worksheet
 - Example 2 Manufacturer received a "codeine (for conversion)" manufacturing quota and utilized it for purposes of making "hydrocodone (for sale)".
- This is NOT to be used to report:
 - the manufacture of dosage forms
 - the manufacture of exempt or excluded products
 - the "conversion" of a schedule II to a III, IV or V

Dispositions – Bulk Conversions



Dispositions **Current Selection** Reporting Year **Drug Code** Drug Name Quota Granted (grams) 2007 PENTOBARBITAL 2270 10.000.000 All amounts are expressed as grams of anhydrous base or alkaloid, not as salt Use the Tab key to navigate between fields. Dispositions (during the year) 2007 Domestic Sales or Transfers to: Pharmacies, Doctors, Hospitals, Clinics 1.000 Other manufacturers 1.000 Distributors for commercial sale 1.000 Distributors for destruction 1.000 Researchers, analytical labs 1.000 Other federal, state or county agencies 1.000 6.000 Total domestic sales or transfers Put in process for conversion to bulk substances. Yes No .000 1.000 Exports 1.000 Returns from you to your suppliers (specify): TEST Quantity used to make exempted or excluded products 1.000 Losses (specify): TEST2 1.000 Authorized destructions via DEA Form 41 1.000 Other dispositions (specify): TEST3 1.000 Total dispositions (grams) 12.000 Click Back to return to the Acquisitions page. Click Next to save your work and view a Summary page for this report. Next Back

Drugs Converted to Other Substances Converted-From Substance **Drug Code Drug Name** Quantity (grams) 2270 PENTOBARBITAL 1.000 Converted-To Substance You are converting the drug listed above to one or more other substances. If the substance to be converted is in the drop down list, click the radio button and select the substance from the list. Then fill in the values for Qty Converted and Qty Produced. If the substance is not in the list, click the radio button for Non-Controlled Substance and type the name of the substance in the Non-Controlled Substance field and fill in the values for Qty Converted and Qty Produced. Controlled Substance : Qty Converted: Qty Produced: Non-Controlled Substance : Qtv Converted: Qty Produced: To record a converted substance, make your selection and click Submit. To see a summary of all the converted substance entries you've made far so far, click Summary. Submit Summary

Dispositions - Bulk Conversions Summary



Drugs Converted to Other Substances

Converted-From Substance

Drug Code	Drug Name	Quantity (grams)
2270	PENTOBARBITAL	1.000

Summary

Remove	Drug Produced	Qty Converted	Qty Produced
0	5-METHOXY-3,4-METHYLENEDIOXY-AMPHETAMINE	1.000	1.000
\circ	4-BROMO-2,5-DIMETHOXYPHENETHYLAMINE	11.000	1.000
Total		12.000	2.000

Click Continue to return to the Converted Substances Selection page. Click Finish to return to the Report Dispositions page.

Continue

Finish

Summary Totals



Year-End Report for Substances Procured Accountability for Outstanding Inventory Summary Totals Current Selection Reporting Year Drug Code Quota Granted (grams) Drug Name 2008 PENTOBARBITAL 10,000.000 2270 All amounts are expressed as grams of anhydrous base or alkaloid, not as salt. Use the Tab key to navigate between fields. **Total Ending Inventory for 2007** 177.494 **Total Ending Inventory for 2008** .000 **Total Acquisitions for 2008** .000 Total To Be Accounted For 177.494 (Total Ending Inventory For 2007 plus Total Acquisitions For 2008) **Total Dispositions for 2008** .000 **Total Accounted For** .000 (Total Ending Inventory For 2008 plus Total Dispositions For 2008) Difference 177.494 (Total To Be Accounted For minus Total Accounted For) Please provide an explanation for any difference: Remarks: To save this report to permanent storage, click Submit. To return to the beginning of this report in order to review or change the entered data, click Back.

To return to the Report Selection List, click Return.

Back

Submit

Return

Confirmation



Year-End Report for Substances Procured Accountability for Outstanding Inventory

Summary Confirmation

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)
2008	2270	PENTOBARBITAL	10,000.000

You have successfully created the year-end report for the controlled substance noted above. The report selection has been moved to the Completed Reports section of the Outstnading Inventory Selection List.

To return to the Report Selection Menu, click Main.

To return to the Report Selection List, click Return.

To exit this application, click Exit.



Return



UN Psychotropic Substance Report



- Specified psychotropic substances in UN Schedules III and IV, member countries voluntarily submit statistics to UN
 - Check the DEA Diversion web site (www.deadiversion.usdoj.gov)
 - Check the INCB web site (www.incb.org)
 - Check the "green list" attached
- Bulk and non-bulk manufacturers should report
- DEA attempted to codify a regulation which would have required such reporting (circa 1992), however never promulgated the rule

Year End Report Psychotropic Selection Menu





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Year-End Report for Psychotropic Substances

Psychotropic Drugs



Psychotropic Substances to be Reported



71 Schedule III:

Buprenorphine

Butalbital

Cathine

Cyclobarbital

Flunitrazepam

Pentazocine

71 Schedule IV:

Allobarbital

Alprazolam

Barbital

Benzphetamine

Bromazepam

Brotizolam

Camazepam

Chlordiazepoxide

Clobazam

Clonazepam

Clorazepate

Clotiazepam

Cloxazolaml

Delorazepam

Diazepam

Diethylpropion

Estazolam

Ethchlorvynol

Ethinamate

Ethyl loflazepate

Fencamfamin

Fenproporex

Fludiazepam

Flurazepam

Halazepam

Haloxazolam

Ketazolam

Loprazolam

Lorazepam

Lormetazepam

Mazindol

Medazepam

Mefenorex

Meprobamate

Mesocarb

Methylphenobarbital

Methyprylon

Midazolam

Nimetazepam

Nitrazepam

Nordazepam

Oxazepam

Oxazolam

Pemoline

Phendimetrazine

Phenobarbital

Phentermine

Pinazepam

Pipradrol

Prazepam

Pyrovalerone

Secbutabarbital

SPA (lefetamine)

Temazepam

Tetrazepam

Triazolam

Vinylbital

Zolpidem

Psychotropic Report





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UN Psychotropic Substances

Psychotropic Drugs

Psychotropic Drugs

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)	
2016	2765	DIAZEPAM		N/A

All amounts are expressed as grams of anhydrous base or alkaloid, not as salt.

	2015	2016			
Inventory (as of December 31)	.000	.000			
Bulk Manufactured		.000			
Dispositions (during 2016)					
Quantity Sold as Finished Dosage Forms		.000			
Quantity Used to Manufacture Non-Psychotropic Substances		.000			
Quantity Used to Manufacture Excluded, Excepted or Exempted Substances		.000			
Total Dispositions		.000	Sum		
Estimates for Future Usage	2017	2018			
Expected Future Usage (Sale) of Finished Dosage Forms	.000	.000			
Remarks: (maximum 255 characters :: 255 characters remaining)					

To save this report to permanent storage, click *Submit*.

To abandon this report and return to the Report Selection List, click *Return*.

Return

Submit

Psychotropic Report





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UN Psychotropic Substances
Psychotropic Drugs

Summary Confirmation

Current Selection

Reporting Year	Drug Code	Drug Name	Quota Granted (grams)	
2016	2765	DIAZEPAM		N/A

You have successfully filed the year-end report for the substance noted above. The report selection has been moved to the Completed Reports section of the Psychotropic Drugs Selection List.

To return to the Report Selection Menu, click Main.

To return to the Report Selection List, click Return.

To exit this application, click Exit.







Year-End Reporting



Thank you!

If you have questions we can be reached via email and phone.

United Nations Reporting and Quota Section (DRQ)

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