of this notice and report are being provided to the Committee on Government Reform of the House of Representatives, the Committee on Homeland Security and Governmental Affairs of the Senate, and the Office of Management and Budget.

Authority: This matching program is being conducted under the authority of the Internal Revenue Code (IRC) 6103(m)(2). This provides for disclosure, upon written request, of a taxpayer's mailing address for use by officers, employees, or agents of a Federal agency for the purpose of locating such taxpayer to collect or compromise a Federal claim against the taxpayer in accordance with sections 3711, 3717, and 3718 of title 31 of the United States Code, statutory provisions which authorize DOJ to collect debts on behalf of the United States through litigation.

Objectives To Be Met by the Matching Program: The purpose of this program is to provide DOJ with the most current addresses of taxpayers to notify debtors of legal actions that may be taken by DOJ and the rights afforded them in the litigation to enforce collection of debts owed to the United States.

Records To Be Matched: DOJ will provide records from the Debt Collection Management System, JUSTICE/JMD–006, last published in its entirety at 58 FR 60058–60060 (November 12, 1993), and from the Debt Collection Enforcement System, JUSTICE/USA–015, last published in its entirety at 71 FR 42118–42122 (July 25, 2006). These systems of records contain information on persons who owe debts to the United States and whose debts have been referred to the DOJ for litigation and/or enforced collection. DOJ records will be matched against records contained in the Privacy Act System of Records: Customer Account Data Engine (CADE) Individual Master File (IMF), Treasury/IRS 24.030, last published at 73 FR 13304 (March 12, 2008). This system of records, among other information, contains the taxpayer's name, Social Security Number (SSN), and most recent address known by IRS.

Categories of Records/Individuals Involved: DOJ will submit the nine-digit Social Security Number (SSN) and four character Name Control (the first four letters of the surname) of each individual whose current address is requested. IRS will provide an address for each taxpayer whose SSN and Name Control matches the record submitted by DOJ or a code explaining that no match was found on the IMF.

Notice Procedures: IRS provides direct notice to taxpayers in the instructions to Forms 1040, 1040A, and 1040EZ, and constructive notice in the **Federal Register** system of records notice, that information provided on U.S. Individual Income Tax Returns may be given to other Federal agencies, as provided by law. For the records involved in this match, both IRS and DOJ have provided constructive notice to records subjects through the publication, in the **Federal Register**, of system of record notices that contain routine uses permitting disclosures for this matching program.

Address for Receipt of Public Comments or Inquiries: Interested persons are invited to submit written comments regarding this notice to Holley B. O'Brien, Director, Debt Collection Management Staff, Justice Management Division, 145 N St. NE., Rm 5E.101, Washington, DC 20530.

Lee Lofthus,

Assistant Attorney General for Administration.

[FR Doc. 2011–31673 Filed 12–8–11; 8:45 am]

BILLING CODE 4410–NW–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–343F]

Controlled Substances: Final Adjusted Aggregate Production Quotas for 2011

AGENCY: Drug Enforcement Administration (DEA), Department of Justice.

ACTION: Notice.

SUMMARY: This notice establishes final adjusted 2011 aggregate production quotas for controlled substances in Schedules I and II of the Controlled Substances Act (CSA).

FOR FURTHER INFORMATION CONTACT: John W. Partridge, Chief, Liaison and Policy Section, Drug Enforcement Administration, 8701 Morrissette Drive, Springfield, VA 22152, *Telephone:* (202) 307–4564.

SUPPLEMENTARY INFORMATION:

Background

Section 306 of the Controlled Substances Act (CSA) (21 U.S.C. 826) requires that the Attorney General establish aggregate production quotas for each basic class of controlled substance listed in Schedules I and II. This responsibility has been delegated to the Administrator of the DEA by 28 CFR 0.100. In accordance with 21 U.S.C. 826 and 21 CFR 1303.11, DEA published in the **Federal Register** on December 20, 2010, notice of the established 2011 aggregate production quotas for controlled substances in Schedules I and II (75 FR 79404). That notice stated that the Administrator would adjust, as needed, the established aggregate production quotas in 2011 as provided for in 21 CFR 1303.13. The 2011 proposed adjusted aggregate production quotas were subsequently published in the **Federal Register** on September 14, 2011, (76 FR 56810) in consideration of the outlined criteria. All interested persons were invited to comment on or object to the proposed adjusted aggregate production quotas on or before October 14, 2011.

The September 14, 2011, proposed adjusted aggregate production quotas also included proposed aggregate production quotas for five newly scheduled substances. On March 1, 2011, the DEA Administrator published a final order (76 FR 11075) which temporarily placed five synthetic cannabinoids in Schedule I: 1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl)indole (JWH-200); 1-Butyl-3-(1-naphthoyl)indole (JWH-073); 1-Pentyl-3-(1-naphthoyl)indole (JWH-018); 5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol; and 5-(1,1-Dimethyloctyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol. That final order stated that quotas for the five substances would be ''established based on registrations granted and quota applications received pursuant to part 1303 of Title 21 of the Code of Federal Regulations.'' 76 FR 11077. Aggregate production quotas for these newly scheduled substances had not been previously established and were initially proposed in the above referenced notice published in the **Federal Register** on September 14, 2011 (76 FR 56810). All interested persons were invited to comment on or object to the proposed aggregate production quotas on or before October 14, 2011.

Analysis for Final Adjusted 2011 Aggregate Production Quotas

Consideration has been given to the criteria outlined in the September 14, 2011, notice of proposed adjusted aggregate production quotas in accordance with 21 CFR 1303.13. In addition, six companies, four DEA registered manufacturers and two non-registrants, submitted timely comments regarding a total of 22 Schedule I and II controlled substances. Comments received proposed that the aggregate production quotas for 4-anilino-N-phenethyl-4-piperidine (ANPP), alfentanil, amphetamine (for sale), diphenoxylate, fentanyl, gamma hydroxybutyric acid, hydrocodone, meperidine, methadone, methadone

intermediate, methylphenidate, morphine (for conversion), morphine (for sale), noroxymorphone (for conversion), noroxymorphone (for sale), oxycodone (for sale), oxymorphone (for conversion), oxymorphone (for sale), pentobarbital, secobarbital, sufentanil, and thebaine were insufficient to provide for the estimated medical, scientific, research, and industrial needs of the United States, for export requirements, and for the establishment and maintenance of reserve stocks. DEA has taken into consideration the above comments along with the relevant 2010 year-end inventories, initial 2011 manufacturing quotas, 2011 export requirements, actual and projected 2011 sales, research and product development requirements, and additional applications received.

Based on all of the above, the Administrator has determined that the proposed adjusted 2011 aggregate production quotas for alfentanil, diphenoxylate, gamma hydroxybutyric acid, meperidine, and pentobarbital required additional consideration and hereby further adjusts the 2011 aggregate production quotas for those substances.

Regarding 4-anilino-N-phenethyl-4-piperidine (ANPP), amphetamine (for sale), fentanyl, hydrocodone, methadone, methadone intermediate, methylphenidate, morphine (for conversion), morphine (for sale), noroxymorphone (for conversion), noroxymorphone (for sale), oxycodone (for sale), oxymorphone (for conversion), oxymorphone (for sale), secobarbital, sufentanil, and thebaine, the Administrator hereby determines that the proposed adjusted 2011 aggregate production quotas for these substances as published on September 14, 2011, at 76 FR 56810 are sufficient to meet the current 2011 estimated medical, scientific, research, and industrial needs of the United States and to provide for adequate inventories.

Pursuant to the above, the Administrator hereby establishes the 2011 final aggregate production quotas for Schedule I and II controlled substances, including the five newly scheduled substances previously referenced, expressed in grams of anhydrous acid or base, as follows:

Basic class—Schedule I	Final adjusted 2011 quotas
1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl)indole (JWH–200)	45 g
1-Butyl-3-(1-naphthoyl)indole (JWH–073)	45 g
1-Methyl-4-phenyl-4-propionoxypiperidine	2 g
1-Pentyl-3-(1-naphthoyl)indole (JWH–018)	45 g
2,5-Dimethoxyamphetamine	2 g
2,5-Dimethoxy-4-ethylamphetamine (DOET)	2 g
2,5-Dimethoxy-4-n-propylthiophenethylamine	2 g
3-Methylfentanyl	2 g
3-Methylthiofentanyl	2 g
3,4-Methylenedioxyamphetamine (MDA)	22 g
3,4-Methylenedioxy-N-ethylamphetamine (MDEA)	15 g
3,4-Methylenedioxymethamphetamine (MDMA)	22 g
3,4,5-Trimethoxyamphetamine	2 g
4-Bromo-2,5-dimethoxyamphetamine (DOB)	2 g
4-Bromo-2,5-dimethoxyphenethylamine (2–CB)	2 g
4-Methoxyamphetamine	77 g
4-Methylaminorex	2 g
4-Methyl-2,5-dimethoxyamphetamine (DOM)	2 g
5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol	68 g
5-(1,1-Dimethyloctyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol	53 g
5-Methoxy-3,4-methylenedioxyamphetamine	2 g
5-Methoxy-N,N-diisopropyltryptamine	2 g
Acetyl-alpha-methylfentanyl	2 g
Acetyldihydrocodeine	2 g
Acetylmethadol	2 g
Allylprodine	2 g
Alphacetylmethadol	2 g
Alpha-ethyltryptamine	2 g
Alphameprodine	2 g
Alphamethadol	2 g
Alpha-methylfentanyl	2 g
Alpha-methylthiofentanyl	2 g
Alpha-methyltryptamine (AMT)	2 g
Aminorex	2 g
Benzylmorphine	2 g
Betacetylmethadol	2 g
Beta-hydroxy-3-methylfentanyl	2 g
Beta-hydroxyfentanyl	2 g
Betameprodine	2 g
Betamethadol	2 g
Betaprodine	2 g
Bufotenine	3 g
Cathinone	4 g
Codeine-N-oxide	602 g
Diethyltryptamine	2 g
Difenoxin	50 g
Dihydromorphine	3,608,000 g
Dimethyltryptamine	7 g
Gamma-hydroxybutyric acid	5,772,000 g
Heroin	20 g

Basic class—Schedule I	Final adjusted 2011 quotas
Hydromorphinol	2 g
Hydroxypethidine	2 g
Ibogaine	5 g
Lysergic acid diethylamide (LSD)	16 g
Marihuana	21,000 g
Mescaline	5 g
Methaqualone	10 g
Methcathinone	4 g
Methyldihydromorphine	2 g
Morphine-N-oxide	605 g
N-Benzylpiperazine	2 g
N,N-Dimethylamphetamine	2 g
N-Ethylamphetamine	2 g
N-Hydroxy-3,4-methylenedioxyamphetamine	2 g
Noracymethadol	2 g
Norlevorphanol	52 g
Normethadone	2 g
Normorphine	18 g
Para-fluorofentanyl	2 g
Phenomorphan	2 g
Pholcodine	2 g
Psilocybin	2 g
Psilocyn	2 g
Tetrahydrocannabinols	393,000 g
Thiofentanyl	2 g
Tilidine	10 g
Trimeperidine	2 g

Basic class—Schedule II	Final adjusted 2011 quotas
1-Phenylcyclohexylamine	2 g
1-Piperdinocyclohexanecarbonitrile	2 g
4-Anilino-N-phenethyl-4-piperidine (ANPP)	1,800,000 g
Alfentanil	12,800 g
Alphaprodine	2 g
Amobarbital	40,007 g
Amphetamine (for conversion)	8,500,000 g
Amphetamine (for sale)	25,300,000 g
Cocaine	216,000 g
Codeine (for conversion)	65,000,000 g
Codeine (for sale)	39,605,000 g
Dextropropoxyphene	7 g
Dihydrocodeine	255,000 g
Diphenoxylate	730,000 g
Ecgonine	83,000 g
Ethylmorphine	2 g
Fentanyl	1,428,000 g
Glutethimide	2 g
Hydrocodone (for sale)	59,000,000 g
Hydromorphone	3,455,000 g
Isomethadone	2 g
Levo-alphacetylmethadol (LAAM)	3 g
Levomethorphan	2 g
Levorphanol	3,600 g
Lisdexamfetamine	10,400,000 g
Meperidine	5,500,000 g
Meperidine Intermediate-A	3 g
Meperidine Intermediate-B	7 g
Meperidine Intermediate-C	3 g
Metazocine	5 g
Methadone (for sale)	20,000,000 g
Methadone Intermediate	26,000,000 g
Methamphetamine	3,130,000 g
Methylphenidate	56,000,000 g
Morphine (for conversion)	70,000,000 g
Morphine (for sale)	39,000,000 g
Nabilone	10,502 g
Noroxymorphone (for conversion)	7,200,000 g
Noroxymorphone (for sale)	401,000 g
Opium (powder)	63,000 g
Opium (tincture)	1,000,000 g
Oripavine	8,000,000 g

Basic class—Schedule II	Final adjusted 2011 quotas
Oxycodone (for conversion)	5,600,000 g
Oxycodone (for sale)	98,000,000 g
Oxymorphone (for conversion)	12,800,000 g
Oxymorphone (for sale)	3,070,000 g
Pentobarbital	34,000,000 g
Phenazocine	5 g
Phencyclidine	24 g
Phenmetrazine	2 g
Phenylacetone	8,000,000 g
Racemethorphan	2 g
Remifentanil	2,500 g
Secobarbital	336,002 g
Sufentanil	5,000 g
Tapentadol	403,000 g
Thebaine	116,000,000 g

Aggregate production quotas for all other Schedule I and II controlled substances included in 21 CFR 1308.11 and 1308.12 remain at zero.

Dated: December 1, 2011.

Michele M. Leonhart,

Administrator.

[FR Doc. 2011–31621 Filed 12–8–11; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA 350F]

Final Adjusted Assessment of Annual Needs for the List I Chemicals: Ephedrine, Pseudoephedrine, and Phenylpropanolamine for 2011

AGENCY: Drug Enforcement Administration (DEA), Department of Justice.

ACTION: Notice.

SUMMARY: This notice establishes the Final Adjusted 2011 assessment of annual needs for the List I chemicals ephedrine, pseudoephedrine, and phenylpropanolamine.

DATES: *Effective Date:* December 9, 2011.

FOR FURTHER INFORMATION CONTACT: John W. Partridge, Chief, Liaison and Policy Section, Drug Enforcement Administration (DEA), Springfield, Virginia 22152, *Telephone:* (202) 307–4564.

SUPPLEMENTARY INFORMATION: The 2011 assessment of annual needs represents those quantities of ephedrine, pseudoephedrine, and phenylpropanolamine which may be manufactured domestically and imported into the United States in 2011 to provide adequate supplies of each chemical for the estimated medical, scientific, research, and industrial needs of the United States, lawful export requirements, and the establishment and maintenance of reserve stocks of such chemicals. Section 306 of the Controlled Substances Act (CSA) (21 U.S.C. 826) requires that the Attorney General establish an assessment of annual needs for ephedrine, pseudoephedrine, and phenylpropanolamine. This responsibility has been delegated to the Administrator of the DEA by 28 CFR 0.100.

On September 14, 2011, a notice entitled ''Proposed Adjustment of the Assessment of Annual Needs for the List I Chemicals Ephedrine, Pseudoephedrine, and Phenylpropanolamine for 2011'' was published in the **Federal Register** (76 FR 56807). That notice proposed to adjust the 2011 assessment of annual needs for ephedrine (for sale), ephedrine (for conversion), pseudoephedrine (for sale), phenylpropanolamine (for sale) and phenylpropanolamine (for conversion). All interested persons were invited to comment on or object to the proposed assessments on or before October 14, 2011.

Comments Received

DEA did not receive any comments to the proposed adjustment of the assessment of annual needs for ephedrine (for sale), ephedrine (for conversion), pseudoephedrine (for sale), phenylpropanolamine (for sale) and phenylpropanolamine (for conversion).

Conclusion

In determining the adjusted 2011 assessments, DEA used the calculation methodology previously described in the 2010 and 2011 assessment of annual needs (74 FR 60294 and 75 FR 79407 respectively). DEA considered changes in demand, changes in the national rate of net disposal, and changes in the rate of net disposal by the registrants holding individual manufacturing or import quotas for the chemical; whether any increased demand or changes in the national and/or individual rates of net disposal are temporary, short term, or long term; whether any increased demand could be met through existing inventories, increased individual manufacturing quotas, or increased importation without increasing the assessment of annual needs; whether any decreased demand would result in excessive inventory accumulation by all persons registered to handle the particular chemical; and other factors affecting the medical, scientific, research, industrial, and importation needs in the United States, lawful export requirements, and reserve stocks, as found relevant.

Other factors that DEA considered include trends as derived from information provided in applications for import, manufacturing, and procurement quotas and in import and export declarations. The inventory, acquisition (purchases), and disposition (sales) data as provided by DEA registered manufacturers and importers reflects the most current information available to DEA at the time of publication of this Notice. The underlying data used to determine the final 2011 assessment of annual needs is the same as that used in determining the proposed 2011 assessment of annual needs, as published on September 14, 2011, at 76 FR 56807.

In accordance with 21 U.S.C. 826(a) and 21 CFR 1315.13, the Administrator hereby orders that the 2011 assessment of annual needs for ephedrine, pseudoephedrine, and phenylpropanolamine, expressed in kilograms of anhydrous acid or base, is adjusted and established as follows: