

*United States v. Coastal Coal Company, et al.*, DOJ No. 90-5-1-4287-1.

Copies of the proposed Consent Decree may be examined at the Office of the United States Attorney, Northern District of West Virginia, 100 Main Street, Room 200, Wheeling, West Virginia 26003; EPA Region III, 1650 Arch Street, Philadelphia, PA 19103; and at the Consent Decree Library, 1120 G Street, N.W., 4th Floor, Washington, D.C. 20005, (202) 624-0892. A copy of the proposed Consent Decree may be obtained in person or by mail from the Consent Decree Library, 1120 G Street, N.W., 4th Floor, Washington, D.C. 20005. When requesting a copy of the proposed Consent Decree, please enclose a check to cover the twenty-five cents per page reproduction costs payable to the "Consent Decree Library" in the amount of \$10, and please reference *United States v. Coastal Coal Company, et al.*, DOJ No. 90-5-1-4287-1.

**Joel M. Gross,**

Chief, Environmental Enforcement Section, Environment and Natural Resources Division, U.S. Department of Justice.

[FR Doc. 98-27852 Filed 10-15-98; 8:45 am]

BILLING CODE 4410-15-M

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[DEA #179P]

**Controlled Substances: Proposed Aggregate Production Quotas for 1999**

**AGENCY:** Drug Enforcement Administration (DEA), Justice.

**ACTION:** Notice of proposed 1999 aggregate production quotas.

**SUMMARY:** This notice proposes initial 1999 aggregate production quotas for controlled substances in Schedules I and II of the Controlled Substances Act (CSA).

**DATES:** Comments or objections must be received on or before November 16, 1998.

**ADDRESSES:** Send comments or objections to the Acting Deputy Administrator, Drug Enforcement Administration, Washington, D.C. 20537, Attn: DEA Federal Register Representative (CCR).

**FOR FURTHER INFORMATION CONTACT:** Frank L. Sapienza, Chief, Drug and Chemical Evaluation Section, Drug Enforcement Administration, Washington, D.C. 20537, Telephone: (202) 307-7183.

**SUPPLEMENTARY INFORMATION:** Section 306 of the 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 Section 0.100 of Title 28 of the Code of Federal Regulations. The Administrator, in turn, has redelegated this function to the Deputy Administrator of the DEA pursuant to Section 0.104 of Title 28 of the Code of Federal Regulations.

The proposed 1999 aggregate production quotas represent those quantities of controlled substances that may be produced in the United States in 1999 to provide adequate supplies of each substance for: the estimated medical, scientific, research, and industrial needs of the United States; lawful export requirements; and the establishment and maintenance of reserve stocks. These quotas do not include imports of controlled

substances for use in industrial processes.

In determining the proposed 1999 aggregate production quotas, the Acting Deputy Administrator considered the following factors: total actual 1997 and estimated 1998 and 1999 net disposals of each substance by all manufacturers; estimates of 1998 year-end inventories of each substance and of any substance manufactured from it and trends in accumulation of such inventories; product development requirements of both bulk and finished dosage form manufacturers; projected demand as indicated by procurement quota applications filed pursuant to Section 1303.12 of Title 21 of the Code of Federal Regulations; and other pertinent information.

Pursuant to Section 1303 of Title 21 of the Code of Federal Regulations, the Acting Deputy Administrator of the DEA will, in early 1999, adjust aggregate production quotas and individual manufacturing quotas allocated for the year based upon 1998 year-end inventory and actual 1998 disposition data supplied by quotas recipients for each basic class of Schedule I or II controlled substance.

Therefore, under the authority vested in the Attorney General by Section 306 of the CSA of 1970 (21 U.S.C. 826), delegated to the Administrator of the DEA by Section 0.100 of Title 28 of the Code of Federal Regulations, and redelegated to the Deputy Administrator pursuant to Section 0.104 of Title 28 of the Code of Federal Regulations, the Acting Deputy Administrator hereby proposes that the 1999 aggregate production quotas for the following controlled substances, expressed in grams of anhydrous acid or base, be established as follows:

Basic class	Proposed 1999 quotas
Schedule I:	
2,5-Dimethoxyamphetamine .....	10,001,000
2,5-Dimethoxy-4-ethylamphetamine (DOET) .....	2
3-Methylfentanyl .....	14
3-Methylthiofentanyl .....	2
3,4-Methylenedioxyamphetamine (MDA) .....	20
3,4-Methylenedioxy-N-ethylamphetamine (MDEA) .....	30
3,4-Methylenedioxymethamphetamine (MDMA) .....	20
3,4,5-Trimethoxyamphetamine .....	2
4-Bromo-2,5-Dimethoxyamphetamine (DOB) .....	2
4-Bromo-2,5-Dimethoxyphenethylamine (2-CB) .....	2
4-Methoxyamphetamine .....	17
4-Methylaminorex .....	3
4-Methyl-2,5-Dimethoxyamphetamine (DOM) .....	2
5-Methoxy-3,4-Methylenedioxyamphetamine .....	2
Acetyl-alpha-methylfentanyl .....	2
Acetyldihydrocodeine .....	2
Acetylmethadol .....	7
Allylprodine .....	2
Alpha-acetylmethadol .....	7

Basic class	Proposed 1999 quotas
Alpha-ethyltryptamine .....	2
Alphameprodine .....	2
Alpha-methadol .....	2
Alpha-methylfentanyl .....	2
Alpha-methylthiofentanyl .....	2
Alphaprodine .....	2
Aminorex .....	7
Benzylmorphine .....	2
Beta-acetylmethadol .....	2
Beta-hydroxy-3-methylfentanyl .....	2
Beta-hydroxyfentanyl .....	2
Betameprodine .....	2
Beta-methadol .....	2
Betaprodine .....	2
Bufotenine .....	2
Cathinone .....	9
Codeine-N-oxide .....	2
Diethyltryptamine .....	3
Difenoxin .....	9,000
Dihydromorphine .....	7
Dimethyltryptamine .....	3
Heroin .....	2
Hydroxypethidine .....	2
Lysergic acid diethylamide (LSD) .....	57
Mescaline .....	8
Methaqualone .....	17
Methcathinone .....	11
Morphine-N-oxide .....	2
N,N-Diemethylamphetamine .....	7
N-Ethyl-1-Phenylcyclohexylamine (PCE) .....	5
N-Ethylamphetamine .....	7
N-Hydroxy-3,4-Methylenedioxyamphetamine .....	4
Noracymethadol .....	2
Norlevorphanol .....	2
Normethadone .....	7
Normorphine .....	7
Para-fluorofentanyl .....	2
Pholcodine .....	2
Propiram .....	415,000
Psilocin .....	2
Psilocybin .....	2
Tetrahydrocannabinols .....	52,000
Thiofentanyl .....	2
Trimeperidine .....	2
Schedule II:	
1-Phenylcyclohexylamine .....	12
1-Piperidinocyclohexanecarbonitrile (PCC) .....	12
Alfentanil .....	2
Amobarbital .....	12
Amphetamine .....	5,554,000
Cocaine .....	251,000
Codeine (for sale) .....	60,641,000
Codeine (for conversion) .....	22,950,000
Desoxyephedrine—662,000 grams of levodesoxyephedrine for use in a non-controlled, non-prescription product and 35,000 grams for methamphetamine .....	697,000
Dextropropoxyphene .....	109,500,000
Dihydrocodeine .....	121,000
Diphenoxylate .....	1,240,000
Ecgonine .....	151,000
Ethylmorphine .....	13
Fentanyl .....	228,000
Glutethimide .....	2
Hydrocodone (for sale) .....	16,314,000
Hydrocodone (for conversion) .....	1,300,000
Hydromorphone .....	856,000
Isomethadone .....	12
Levo-alpha-acetylmethadol (LAAM) .....	201,000
Levomethorphan .....	2
Levorphanol .....	15,000
Meperidine .....	10,294,000
Methadone (for sale) .....	4,992,000
Methadone (for conversion) .....	267,000
Methadone Intermediate .....	7,223,000

Basic class	Proposed 1999 quotas
Methamphetamine (for conversion) .....	723,000
Methylphenidate .....	14,442,000
Morphine (for sale) .....	12,445,000
Morphine (for conversion) .....	76,300,000
Nabilone .....	2
Noroxymorphone (for sale) .....	25,000
Noroxymorphone (for conversion) .....	2,067,000
Opium .....	640,000
Oxycodone (for sale) .....	12,118,000
Oxycodone (for conversion) .....	51,000
Oxymorphone .....	166,000
Pentobarbital .....	17,356,000
Phencyclidine .....	40
Phenmetrazine .....	2
Phenylacetone .....	10
Secobarbital .....	25
Sufentanil .....	752
Thebaine .....	17,695,000

The Acting Deputy Administrator further proposes that aggregate production quotas for all other Schedules I and II controlled substances included in Sections 1308.11 and 1308.12 of Title 21 of the Code of Federal Regulations be established at zero.

All interested persons are invited to submit their comments and objections in writing regarding this proposal. A person may object to or comment on the proposal relating to any of the above-mentioned substances without filing comments or objections regarding the others. If a person believes that one or more of these issues warrant a hearing, the individual should so state and summarize the reasons for this belief.

In the event that comments or objections to this proposal raise one or more issues which the Acting Deputy Administrator finds warrant a hearing, the Acting Deputy Administrator shall order a public hearing by notice in the Federal Register, summarizing the issues to be heard and setting the time for the hearing.

The Office of Management and Budget has determined that notices of aggregate production quotas are not subject to centralized review under Executive Order 12866. This action has been analyzed in accordance with the principles and criteria contained in Executive Order 12612, and it has been determined that this matter does not have sufficient federalism implications to warrant the preparation of a Federalism Assessment.

The Acting Deputy Administrator hereby certifies that this action will have no significant impact upon small entities whose interests must be considered under the Regulatory Flexibility Act, 5 U.S.C. 601 et seq. The establishment of aggregate production

quotas for Schedules I and II controlled substance is mandated by law and by international treaty obligations. Aggregate production quotas apply to approximately 200 DEA registered bulk and dosage form manufacturers of Schedules I and II controlled substances. The quotas are necessary to provide for the estimated medical, scientific, research and industrial needs of the United States, for export requirements and the establishment and maintenance of reserve stocks. While aggregate production quotas are of primary importance to large manufacturers, their impact upon small entities is neither negative nor beneficial. Accordingly, the Acting Deputy Administrator has determined that this action does not require a regulatory flexibility analysis.

Dated: October 8, 1998.

**Donnie R. Marshall,**  
*Acting Deputy Administrator.*  
 [FR Doc. 98-27740 Filed 10-15-98; 8:45 am]  
 BILLING CODE 4410-09-M

**DEPARTMENT OF JUSTICE**  
**Immigration and Naturalization Service**

**Agency Information Collection**  
**Activities: Proposed Collection;**  
**Comment Request**

**ACTION:** Extension of existing collection; reengineered foreign students pilot program.

The Department of Justice, Immigration and Naturalization Service (INS) has submitted the following information request to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995. The information collection was

previously published in the **Federal Register** on July 9, 1998 at 63 FR 37142, allowing for a 60-day comment period. No comments were received by the Immigration and Naturalization Service.

The purpose of this notice is to allow an additional 30 days for public comments. Comments are encouraged and will be accepted until November 16, 1998. This process is conducted in accordance with 5 CFR Part 1320.10.

Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Office of Information and Regulatory Affairs, Attention: OMB Desk Officer for the Immigration and Naturalization Service, Office of Management and Budget, Room 10235, Washington, DC 20530 (202-395-7316). Additionally, comments may be submitted to OMB via facsimile to 202-395-7285. Comments may also be submitted to the Department of Justice (DOJ), Justice Management Division, Information Management and Security Staff, Attention: Stuart Shapiro, Department Clearance Officer, Suite 850, 1001 G Street, NW., Washington, DC 20530. Additionally, comments may be submitted to DOJ via facsimile to 202-514-1534.

Written comments and suggestions from the public should address one or more of the following points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies/components estimate of the burden of the proposed collection of