(OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. Comments are encouraged and will be accepted for ''sixty days'' until March 31, 2014. This process is conducted in accordance with 5 CFR 1320.10.

If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact April Carroll, Chief, Law Enforcement Support Branch, National Tracing Center, 244 Needy Road, Martinsburg, WV 25405.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

—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;

—Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

—Enhance the quality, utility, and clarity of the information to be collected; and

—Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

*Summary of Information Collection:*

(1) *Type of Information Collection:* Extension of a currently approved collection.

(2) *Title of the Form/Collection:* Federal Firearms Licensee Firearms Inventory Theft/Loss Report.

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: ATF F 3310.11. Bureau of Alcohol, Tobacco, Firearms and Explosives.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Individuals or households. Other: Business or other for-profit.

*Need for Collection:*

Authorization of this form is requested as the Violent Crime Control and Law Enforcement Act requires Federal firearms licensees to report to the Bureau of Alcohol,Tobacco, Firearms and Explosives and to the appropriate local authorities any theft or loss of a firearm from the licensee's inventory or collection, within a specific time frame after the theft or loss is discovered.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* It is estimated that 4,000 respondents will complete a 24 minute form.

(6) *An estimate of the total public burden (in hours) associated with the collection:* There are an estimated 1,600 annual total burden hours associated with this collection.

If additional information is required contact: Jerri Murray, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, Department of Justice, Two Constitution Square, 145 N Street NE., Room 3W–1407B, Washington, DC 20530.

Dated: January 27, 2014.

**Jerri Murray,**

*Department Clearance Officer, PRA, U.S. Department of Justice.*

[FR Doc. 2014–01845 Filed 1–29–14; 8:45 am]

**BILLING CODE 4410–FY–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–387P]**

### Controlled Substances: 2014 Proposed Aggregate Production Quota for Three Temporarily Controlled Synthetic Phenethylamines

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

**ACTION:** Notice of a proposed 2014 aggregate production quota for three synthetic phenethylamines.

**SUMMARY:** Three synthetic phenethylamines 2-(4-iodo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl)ethanamine (25I-NBOMe; 2C-I-NBOMe; 25I; Cimbi-5), 2-(4-chloro-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl)ethanamine (25C-NBOMe; 2C-C-NBOMe; 25C; Cimbi-82), and 2-(4-bromo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl)ethanamine (25B-NBOMe; 2C-B-NBOMe; 25B; Cimbi-36) were temporarily placed in schedule I of the Controlled Substances Act (CSA) by a final order published by the Drug Enforcement Administration (DEA) on November 15, 2013 (78 FR 68716). This means that any manufacturer that wishes to manufacture 25I-NBOMe, 25C-NBOMe, or 25B-NBOMe after November 15, 2013, must be registered with the DEA and have obtained a manufacturing quota for 25I-NBOMe, 25C-NBOMe, or 25B-NBOMe pursuant to 21 CFR part 1303. The DEA cannot issue individual manufacturing quotas for 25I-NBOMe, 25C-NBOMe, or 25B-NBOMe unless and until it establishes an aggregate production quota. Therefore, this notice proposes a 2014 aggregate production quota for 25I-NBOMe, 25C-NBOMe, and 25B-NBOMe.

**DATES:** Comments or objections should be received on or before March 3, 2014.

**ADDRESSES:** To ensure proper handling of comments, please reference ''Docket No. DEA–387P'' on all electronic and written correspondence. The DEA encourages that all comments be submitted electronically through *www.regulations.gov* using the electronic comment form provided on that site. An electronic copy of this document is also available at *www.regulations.gov* for easy reference. Paper comments that duplicate the electronic submission are not necessary as all comments submitted to *www.regulations.gov* will be posted for public review and are part of the official docket record. Written comments submitted via regular or express mail should be sent to the Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

**FOR FURTHER INFORMATION CONTACT:** Ruth A. Carter, Acting Chief, Policy Evaluation and Analysis Section, Office of Diversion Control, Drug Enforcement Administration; Mailing Address: 8701 Morrissette Drive, Springfield, Virginia 22152, Telephone: (202) 598–6812.

**SUPPLEMENTARY INFORMATION:**

#### Posting of Public Comments

The Freedom of Information Act applies to all comments received. All comments received are considered part of the public record and made available for public inspection online at *www.regulations.gov* and in the DEA's public docket. Such information includes personal identifying information (such as your name, address, etc.) voluntarily submitted by the commenter.

If you want to submit personal identifying information (such as your name, address, etc.) as part of your comment, but do not want it to be posted online or made available in the public docket, you must include the

phrase ''PERSONAL IDENTIFYING INFORMATION'' in the first paragraph of your comment. You must also place all the personal identifying information you do not want posted online or made available in the public docket in the first paragraph of your comment and identify what information you want redacted.

If you want to submit confidential business information as part of your comment, but do not want it to be posted online or made available in the public docket, you must include the phrase ''CONFIDENTIAL BUSINESS INFORMATION'' in the first paragraph of your comment. You must also prominently identify confidential business information to be redacted within the comment. If a comment has so much confidential business information that it cannot be effectively redacted, all or part of that comment may not be posted online or made available in the public docket.

Personal identifying information and confidential business information identified and located as set forth above will be redacted, and the comment, in redacted form, will be posted online and placed in the DEA's public docket file.

If you wish to inspect the DEA's public docket file in person by appointment, please see the **FOR FURTHER INFORMATION CONTACT** paragraph.

**Background**

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 each year. This responsibility has been delegated to the Administrator of the DEA by 28 CFR 0.100. The Administrator, in turn, has redelegated this function to the Deputy Administrator, pursuant to 28 CFR 0.104.

The DEA established the 2014 aggregate production quotas for substances in schedules I and II on September 9, 2013 (78 FR 55099). Subsequently, on October 10, 2013, the DEA published in the **Federal Register** a notice of intent to temporarily place three synthetic phenethylamines (25I-NBOMe, 25C-NBOMe, and 25B-NBOMe) in schedule I of the CSA (78 FR 61991). On November 15, 2013, the DEA published in the **Federal Register** a final order to temporarily place these three synthetic phenethylamines in schedule I of the CSA (78 FR 68716), making all regulations pertaining to schedule I controlled substances applicable to the manufacture of these three synthetic phenethylamines, including the establishment of an aggregate production quota pursuant to 21 CFR 1303.11.

25I-NBOMe, 25C-NBOMe, and 25B-NBOMe were non-controlled substances when the aggregate production quotas for schedule I and II substances were established, therefore, no aggregate production quotas for 25I-NBOMe, 25C-NBOMe, and 25B-NBOMe were established at that time.

In determining the 2014 aggregate production quotas of these three phenethylamines, the Deputy Administrator considered the following factors in accordance with 21 U.S.C. 826(a) and 21 CFR 1303.11: (1) Total estimated net disposal of each substance by all manufacturers; (2) estimated trends in the national rate of net disposal; (3) total estimated inventories of the basic class and of all substances manufactured from the class; (4) projected demand for each class as indicated by procurement quotas requested pursuant to 21 CFR 1303.12; and (5) other factors affecting medical, scientific, research, and industrial needs of the United States and lawful export requirements, as the Deputy Administrator finds relevant. These quotas do not include imports of controlled substances for use in industrial processes.

The Deputy Administrator, therefore, proposes that the year 2014 aggregate production quotas for the following temporarily controlled schedule I controlled substances, expressed in grams of anhydrous acid or base, be established as follows:

| Basic class–schedule I | Proposed 2014 quota |
|---|---|
| 2-(4-bromo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl)ethanamine (25B-NBOMe; 2C-B-NBOMe; 25B; Cimbi-36) | 15 g |
| 2-(4-chloro-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl)ethanamine (25C-NBOMe; 2C-C-NBOMe; 25C; Cimbi-82) | 15 g |
| 2-(4-iodo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl)ethanamine (25I-NBOMe; 2C-I-NBOMe; 25I; Cimbi-5) | 15 g |

**Comments**

Pursuant to 21 CFR 1303.11, any interested person may submit written comments on or objections to these proposed determinations. Based on comments received in response to this Notice, the Deputy Administrator may hold a public hearing on one or more issues raised. In the event the Deputy Administrator decides in his sole discretion to hold such a hearing, the Deputy Administrator will publish a notice of any such hearing in the **Federal Register**. After consideration of any comments and after a hearing, if one is held, the Deputy Administrator will publish in the **Federal Register** a Final Order establishing the 2014 aggregate production quota for 25I-NBOMe, 25C-NBOMe, and 25B-NBOMe.

Dated: January 17, 2014.

**Thomas M. Harrigan,**

*Deputy Administrator.*

[FR Doc. 2014–01778 Filed 1–29–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. 13–40]**

### House of Medicine; Decision and Order

On October 2, 2013, Administrative Law Judge (ALJ) Christopher B. McNeil issued the attached Recommended Decision (R.D.). Therein, the ALJ found that there was no dispute over the material fact that Respondent does not possess authority under the laws of California, the State in which it has applied for a DEA Certificate of Registration as a Retail Pharmacy, to dispense controlled substances. R.D. at 5–6. Accordingly, the ALJ held that Applicant does not meet the statutory definition of a practitioner, *see* 21 U.S.C. 802(21), and therefore is not entitled to be registered under 21 U.S.C. 823(f). *Id.* at 6. The ALJ thus granted the Government's Motion for Summary Disposition and recommended that the Administrator deny Respondent's application. *Id.* at 7. Neither party filed exceptions to the Recommended Decision.

Having reviewed the record, I have decided to adopt the ALJ's Recommended Decision in its entirety except as discussed below.[1]

[1] In the R.D., the ALJ found that the Order to Show Cause was issued on August 6, 2013. R.D. at 2. The ALJ then found that ''[o]n December 26, 2012, Respondent . . . filed a timely request for

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