

Settlement Agreement may also be examined on the following Department of Justice Web site, http://www.usdoj.gov/enrd/Consent_Decrees.html. A copy of the Settlement Agreement may also be obtained by mail from the Consent Decree Library, U.S. Department of Justice, P.O. Box 7611, Washington, DC 20044-7611 or by faxing, or e-mailing a request to Tonia Fleetwood (tonia.fleetwood@usdoj.gov), fax no. (202) 512-0097, phone confirmation number (202) 514-1547. In requesting a copy from the Consent Decree Library, please refer to *United States v. Dean R. Soulliere and Colleen A. Soulliere, and Soulliere and Jackson, Inc.*, d/b/a One Hour Martinizing (Settlement Agreement with Dean R. Soulliere *et al.*, DOJ Ref. No. 90-11-2-07430), and enclose a check in the amount of \$2.00 (25 cents per page reproduction cost) payable to the U.S. Treasury or, if by e-mail or fax, forward a check in that amount to the Consent Decree Library at the stated address.

Robert E. Maher, Jr.,
Assistant Section Chief, Environmental
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117-0013]

Agency Information Collection Activities: Proposed Collection; Comments Requested

ACTION: 30-Day Notice of Information Collection Under Review; Application for Permit to Import Controlled Substances for Domestic and/or Scientific Purposes pursuant to 21 U.S.C. 952—DEA Form 357.

The Department of Justice (DOJ), Drug Enforcement Administration (DEA) will be submitting the following information collection request to the Office of Management and Budget (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. This proposed information collection was previously published in the **Federal Register** Volume 72, Number 71, page 18668 on April 13, 2007, allowing for a 60 day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until July 23, 2007. This

process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202) 395-5806.

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 agencies 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.

Overview of this Information Collection:

(1) *Type of Information Collection:* Renewal of an existing collection.

(2) *Title of the Form/Collection:* Application for Permit to Import Controlled Substances for Domestic and/or Scientific Purposes pursuant to 21 U.S.C. 952 (DEA Form 357).

(3) *Agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form number: DEA Form 357.

Component: Office of Diversion Control, Drug Enforcement Administration, U.S. Department of Justice.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:*

Primary: Business or other for-profit.

Other: None.

Abstract: Title 21, CFR, Section 1312.11 requires any registrant who desires to import certain controlled

substances into the United States to have an import permit. In order to obtain the permit, an application must be made to the Drug Enforcement Administration on DEA Form 357.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* 47 respondents, 406 responses, .25 hour per response. A respondent may submit multiple responses. A respondent will take an estimate of 15 minutes to complete each form.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 101.5 annual burden hours.

If additional information is required contact: Lynn Bryant, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Patrick Henry Building, Suite 1600, 601 D Street, NW., Washington, DC 20530.

Dated: June 15, 2007.

Lynn Bryant,
Department Clearance Officer, PRA
Department of Justice.

[FR Doc. E7-12035 Filed 6-20-07; 8:45 am]

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DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review: Comment Request

June 18, 2007.

The Department of Labor (DOL) has submitted the following public information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (Pub. L. 104-13, 44 U.S.C. Chapter 35). A copy of this ICR, with applicable supporting documentation, may be obtained at <http://www.reginfo.gov/public/do/PRAMain>, or contact Ira Mills on 202-693-4122 (this is not a toll-free number) or e-mail: Mills.Ira@dol.gov.

Comments should be sent to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for U.S. Department of Labor/Employment and Training Administration (ETA), Office of Management and Budget, Room 10235, Washington, DC 20503, 202-395-7316 (this is not a toll free number), within 30 days from the date of this publication in the **Federal Register**.

The OMB is particularly interested in comments which:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the