burden of 80 hours. It is estimated that 250 persons complete DEA Form 224c electronically, at 15 minutes per form, for an annual burden of 62.5 hours.

(6) *An estimate of the total public burden (in hours) associated with the collection:* It is estimated that this collection will create a burden of 56,354 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: August 18, 2009.

**Lynn Bryant,**

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

[FR Doc. E9–20183 Filed 8–20–09; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[OMB Number 1117–0050]**

### Agency Information Collection Activities: Proposed Collection; Comments Requested: Reports of Dispensing of Controlled Substances by Online Pharmacies (DEA Form 332)

**ACTION:** 30-day notice of information collection under review.

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 74, Number 113, page 28275, on June 15, 2009, allowing for a 60-day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until September 21, 2009. 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 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.

**Overview of Information Collection 1117–0050**

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

(2) *Title of the Form/Collection:* Reports of Dispensing of Controlled Substances by Online Pharmacies (DEA Form 332).

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:*

*Form Number:* DEA Form 332; Office of Diversion Control, Drug Enforcement Administration, 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:* Not-for-Profit Institutions; State, Local or Tribal Government.

*Abstract:* The Controlled Substances Act (21 U.S.C. 827(d)(2)) requires online pharmacies to report to DEA the total quantity of controlled substances that the pharmacy has dispensed during each calendar month by any means, regardless of whether the controlled substances are dispensed by means of the Internet. Reports are required to be filed by every pharmacy that, at any time during a calendar month, holds a modification of registration authorizing it to operate as an online pharmacy, regardless of whether the online pharmacy dispenses any controlled substances by means of the Internet during the month. Such reporting is mandated by the Ryan Haight Act and permits DEA to monitor the dispensing of controlled substances by online pharmacies.

(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 250 persons complete DEA Form 332 electronically, at 15 minutes per form, for an annual burden of 750 hours.

(6) *An estimate of the total public burden (in hours) associated with the collection:* It is estimated that this collection will create a burden of 750 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: August 18, 2009.

**Lynn Bryant,**

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

[FR Doc. E9–20184 Filed 8–21–09; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Antitrust Division

### Notice Pursuant to the National Cooperative Research and Production Act of 1993—American Society of Mechanical Engineers

Notice is hereby given that, on July 20, 2009, pursuant to section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), American Society of Mechanical Engineers (''ASME'') has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing additions or changes to its standards development activities. The notifications were filed for the purpose of extending the Act's provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, since May 7, 2009, ASME has partitioned its existing standards writing committee on Boilers and Pressure Vessels into ten separate standards writing committees, each with its own individual charter; published two new standards; and initiated three new standards activities within the general nature and scope of ASME's standards development activities, as