Chelmsford, MA; Manufacturing Techniques, Inc., Kilmarnock, VA; QorTek, Inc., Williamsport, PA; Resodyn Acoustic Mixers, Butte, MT; Rockwell Collins, Cedar Rapids, IA; Sabre Consulting and Training, LLC, Wharton, NJ; and UTRON, Inc., Manassas, VA, have been added as a party to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and NWEC intends to file additional written notifications disclosing all changes in membership.

On June 29, 2000, NWEC filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to section 6(b) of the Act on June 30, 2000 (65 FR 40693).

The last notification was filed with the Department on April 16, 2009. A notice was published in the **Federal Register** pursuant to section 6(b) of the Act on May 22, 2009 (74 FR 24035).

Patricia A. Brink,

Deputy Director of Operations, Antitrust Division.

[FR Doc. 2010–22221 Filed 9–7–10; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117–0047]

Agency Information Collection Activities: Proposed Collection; Comments Requested; Application for Import Quota for Ephedrine, Pseudoephedrine, and Phenylpropanolamine (DEA Form 488)

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 75, Number 128, Page 38834 on July 6, 2010, allowing for a 60 day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until October 8, 2010. 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 1117–0013

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

(2) *Title of the Form/Collection:* Application for Import Quota for Ephedrine, Pseudoephedrine, and Phenylpropanolamine (DEA Form 488).

(3) *Agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form number: DEA Form 488, 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: None.

Abstract: 21 U.S.C. 952 and 21 CFR 1315.34 require that persons who desire to import the List I chemicals ephedrine, pseudoephedrine, and phenylpropanolamine during the next calendar year shall apply on DEA Form 488 for import quota for such List I chemicals.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* DEA estimates that fifty-seven (57) individual respondents will submit eighty (80) individual import quota applications. DEA estimates that each response will take one hour.

(6) *An estimate of the total public burden (in hours) associated with the collection:* DEA estimates that this collection will involve eighty (80) annual public burden hours.

IF ADDITIONAL INFORMATION IS REQUIRED CONTACT: Lynn Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street, NE., Suite 2E–502, Washington, DC 20530.

Dated: August 2, 2010.

Lynn Murray,

Department Clearance Officer, PRA, United States Department of Justice.

[FR Doc. 2010–22384 Filed 9–7–10; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117–0013]

Agency Information Collection Activities: Proposed Collection; Comments Requested; Application for Permit To Import Controlled Substances for Domestic and/or Scientific Purposes Pursuant to 21 U.S.C. 952; DEA Form 357

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 75, Number 128, page 38835 on July 6, 2010, allowing for a 60 day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until October 8, 2010. This process is conducted in accordance with 5 CFR 1320.10.