Caryville Public Library, Caryville, TN.
Jacksboro Public Library, Jacksboro, TN.
Jellico Public Library, Jellico, TN.
LaFollette Public Library, LaFollette, TN.
Huntsville Public Library, Huntsville, TN.
Oneida Public Library, Oneida, TN.
Winfield Public Library, Winfield, TN.
Coalfield Public Library, Coalfield, TN.
Deer Lodge Public Library, Deer Lodge, TN.
Oakdale Public Library, Oakdale, TN.
Petros Public Library, Petros, TN.
Sunbright Public Library, Sunbright, TN.
Wartburg Public Library, Wartburg, TN.
Art Circle Public Library, Crossville, TN.
Fentress County Public Library, Jamestown, TN.

Virginia

Buchanan County Public Library, Grundy, VA.
Wise County Public Library, Wise, VA.
Russell County Public Library, Lebanon, VA.
Tazewell County Public Library, Tazewell, VA.
Scott County Public Library, Gate City, VA.
Lee County Public Library, Pennington Gap, VA.

West Virginia

Ansted Public Library, Ansted, WV.
Boone—Madison Public Library, Madison, WV.
Bradshaw Public Library , Davy, WV.
Clay Co. Public Library, Clay, WV.
Fort Gay Public Library, Fort Gay, WV.
Gilbert Public Library, Gilbert, WV.
Glasgow Public Library, Glasgow, WV.
Graigsville Public Library, Graigsville, WV.
Fayetteville Public Library, Fayetteville, WV.
Fayette County Public Libraries, Oak Hill, WV.
Hamlin—Lincoln Co., Hamlin WV.
Kanawha Co. Public, Charleston, WV.
Kermit Public Library, Kermit, WV.
Logan Area Public Library, Logan, WV.
Mingo County Public Library, Delbarton, WV.
McDowell County Public Library, Welch, WV.
Oceana Public Library, Oceana, WV.
Raleigh Public Library, Beckley, WV.
Sutton Public Library, Sutton, WV.
Wayne County Public Library, Kenova, WV.
Branch of Wayne County Public Library, Wayne, WV.
Whitesville Public Library, Whitesville, WV.

Dated: July 27, 2007.

Sterling Rideout,

Assistant Director, Program Support.

[FR Doc. E7–16628 Filed 8–23–07; 8:45 am]

BILLING CODE 4310–05–P

DEPARTMENT OF JUSTICE

Antitrust Division

Notice Pursuant to the National Cooperative Research and Production Act of 1993—Network Centric Operations Industry Consortium, Inc.

Notice is hereby given that, on July 25, 2007, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (''the Act''), Network Centric Operations Industry Consortium, Inc. has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. 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, BT Ltd., Diegem, BELGIUM; and SRA International, Fairfax, VA have been added as parties 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 Network Centric Operations Industry Consortium, Inc. intends to file additional written notifications disclosing all changes in membership.

On November 19, 2004, Network Centric Operations Industry Consortium, Inc. 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 February 2, 2005 (70 FR 5486).

The last notification was filed with the Department on May 18, 2007. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on July 13, 2007 (72 FR 38618).

J. Robert Kramer, II,

Director of Operations, Antitrust Division.

[FR Doc. 07–4146 Filed 8–23–07; 8:45 am]

BILLING CODE 4410–11–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117–0006]

Agency Information Collection Activities: Proposed Collection; Comments Requested

ACTION: 30-day notice of information collection under review: Application for individual manufacturing quota for a basic class of controlled substance and for ephedrine, pseudoephedrine, and phenylpropanolamine DEA Form189.

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 117, page 33774 on June 19, 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 September 24, 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:* Extension of an existing collection.

(2) *Title of the Form/Collection:* Application for Individual Manufacturing Quota for a Basic Class of Controlled Substance and for Ephedrine, Pseudoephedrine, and Phenylpropanolamine.

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

Form number: DEA Form 189.

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: 21 U.S.C. 826 and 21 CFR 1303.22 and 1315.22 require that any person who is registered to manufacture any basic class of controlled substances listed in Schedule I or II and who desires to manufacture a quantity of such class, or who desires to manufacture using the List I chemicals ephedrine, pseudoephedrine, or phenylpropanolamine, must apply on DEA Form 189 for a manufacturing quota for such quantity of such class or List I chemical.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* DEA estimates that each form takes 0.5 hours (30 minutes) to complete. In total, 37 firms submit 298 responses, with each response taking 0.5 hours (30 minutes) to complete. This results in a total public burden of 149 hours annually.

(6) *An estimate of the total public burden (in hours) associated with the collection:* In total, 37 firms submit 298 responses, with each response taking 0.5 hours (30 minutes) to complete. This results in a total public burden of 149 hours annually.

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 20, 2007.

Lynn Bryant,

Department Clearance Officer, PRA, Department of Justice.

[FR Doc. E7–16718 Filed 8–23–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117–0008]

Agency Information Collection Activities: Proposed Collection; Comments Requested

ACTION: 30-day notice of information collection under review: Application for procurement quota for controlled substances and ephedrine, pseudoephedrine, and phenylpropanolamine.

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 117, page 33775 on June 19, 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 September 24, 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:* Revision of an existing collection.

(2) *Title of the Form/Collection:* Application for Procurement Quota for Controlled Substances and Ephedrine, Pseudoephedrine, and Phenylpropanolamine.

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

Form number: DEA Form 250.

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: 21 U.S.C. 826 and 21 CFR 1303.12 and 1315.32 require that U.S. companies who desire to use any basic class of controlled substances listed in Schedule I or II or the List I chemicals ephedrine, pseudoephedrine, and phenylpropanolamine for purposes of manufacturing during the next calendar year shall apply on DEA Form 250 for procurement quota for such class or List I chemical.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* DEA estimates that each form takes 1 hour to complete. DEA estimates that 240 individual respondents will respond to this form.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 240 individual respondents will spend one hour annually completing this form for the List I chemicals ephedrine, pseudoephedrine, and phenylpropanolamine. This results in an annual public burden of 240 hours.

This form is already used to collect information regarding controlled