and complainant Trend Micro filed a contingent petition for review. The IA did not file a petition. On May 27, 2005, Fortinet filed a response to Trend Micro's contingent petition for review, and Trend Micro filed a response to Fortinet's petition for review. On June 2, 2005, the IA filed a response to Trend Micro and Fortinet's petition for review.

On July 8, 2005, the Commission issued a notice indicating that it had determined not to review the ALJ's final ID on violation, thereby finding a violation of section 337.70 FR 40731 (July 14, 2005). The Commission also invited the parties to file written submission regarding the issues of remedy, the public interest, and bonding, and provided a schedule for filing such submissions.

Having reviewed the record in this investigation, including the parties' written submissions and responses thereto, the Commission determined that the appropriate form of relief in this investigation is a limited exclusion order prohibiting the unlicensed entry of systems for detecting and removing viruses or worms, components thereof and products containing same covered by claims 4, 7, 8, and 11–15 of the '600 patent. The order covers systems for detecting and removing viruses or worms, components thereof and products containing same that are manufactured abroad by or on behalf of, or imported by or on behalf of the respondent, or any of their affiliated companies, parents, subsidiaries, or other related business entities, or their successors or assigns.

The Commission also determined to issue a cease and desist order prohibiting the respondent from importing, selling, marketing, advertising, distributing, offering for sale, transferring (except for exportation), and soliciting U.S. agents or distributors for systems for detecting and removing viruses or worms, components thereof and products containing same.

The Commission further determined that the public interest factors enumerated in sections 337(d)(1) and (f)(1), 19 U.S.C. 1337(d)(1) and (f)(1), do not preclude issuance of either the limited exclusion order or the cease and desist order. In addition, the Commission determined that the amount of bond to permit temporary importation during the Presidential review period shall be in the amount of 100 percent of the entered value of the imported articles. The Commission's orders and opinion in support thereof were delivered to the President on the day of their issuance.

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and section 210.50 of the Commission's Interim Rules of Practice and Procedure (19 CFR 210.50).

Issued: August 8, 2005.

By order of the Commission.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. 05–15934 Filed 8–10–05; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Agency Information Collection Activities: Proposed Collection; Comments Requested

ACTION: 60-Day notice of information collection under review: Application for registration (DEA Form 224); Application for registration renewal (DEA Form 224a); and Affidavit for chain renewal (DEA Form 224B).

The Department of Justice (DOJ), Drug Enforcement Administration (DEA), has submitted 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. Comments are encouraged and will be accepted for ''sixty days'' until October 11, 2005. 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 Patricia M. Good, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537.

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 a currently approved collection.

(2) *Title of the Form/Collection:* Application for Registration (DEA Form 224); Application for Registration Renewal (DEA Form 224a); and Affidavit for Chain Renewal (DEA Form 224B)

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: DEA Form 224, 224a and 224B; 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. All firms and individuals who distribute or dispense controlled substances must register with the DEA under the Controlled Substances Act. Registration is needed for control measures over legal handlers of controlled substances and is used to monitor their activities.

(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 13,510 persons complete DEA Form 224 on paper, at 12 minutes per form, for an annual burden of 2,702 hours. It is estimated that 41,839 persons complete DEA Form 224 electronically, at 8 minutes per form, for an annual burden of 5,579 hours. It is estimated that 159,009 persons complete DEA Form 224a on paper, at 12 minutes per form, for an annual burden of 31,820 hours. It is estimated that 178,884 persons complete DEA Form 224a electronically, at 4 minutes per form, for an annual burden of 11,926 hours. It is estimated that 72 persons complete DEA Form 224b, at 5 hours per form, for an annual burden of 360 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 52,387 annual burden hours.

If additional information is required contact: Brenda E. Dyer, 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 5, 2005.

Brenda E. Dyer,

Department Clearance Officer, Department of Justice.

[FR Doc. 05–15874 Filed 8–10–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Agency Information Collection Activities: Proposed Collection; Comments Requested:

ACTION: 60-day notice of information collection under review: Application for registration (DEA Form 225); Application for registration renewal (DEA Form 225a); Affidavit for chain renewal (DEA Form 225B).

The Department of Justice (DOJ), Drug Enforcement Administration (DEA), has submitted 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. Comments are encouraged and will be accepted for ''sixty days'' until October 11, 2005. 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 Patricia M. Good, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537.

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 a currently approved collection.

(2) *Title of the Form/Collection:* Application for Registration (DEA Form 225); Application for Registration Renewal (DEA Form 225a); Affidavit for Chain Renewal (DEA Form 225B).

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: DEA Form 225, 225a, and 225B; 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: The Controlled Substances Act requires all persons who manufacture, distribute, import, export, conduct research or dispense controlled substances to register with DEA. Registration provides a closed system of distribution to control the flow of controlled substances through the distribution chain.

(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 1,318 persons complete DEA Form 225 on paper, at 30 minutes per form, for an annual burden of 659 hours. It is estimated that 284 persons complete DEA Form 225 electronically, at 10 minutes per form, for an annual burden of 47 hours. It is estimated that 5,338 persons complete DEA Form 225a on paper, at 30 minutes per form, for an annual burden of 2,669 hours. It is estimated that 4 persons complete DEA Form 225B on paper, at one hour per form, for an annual burden of 4 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 3,379 annual burden hours.

If additional information is required contact: Brenda E. Dyer, 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 5, 2005.

Brenda E. Dyer,

Department Clearance Officer, Department of Justice.

[FR Doc. 05–15875 Filed 8–10–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Agency Information Collection Activities: Proposed Collection; Comments Requested

ACTION: 60-day notice of information collection under review: Application for registration (DEA Form 363) and application for registration renewal (DEA Form 363a).

The Department of Justice (DOJ), Drug Enforcement Administration (DEA), has submitted 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. Comments are encouraged and will be accepted for ''sixty days'' until October 11, 2005. 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 Patricia M. Good, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537.

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,