

termination of this investigation does not impose any undue burdens on the public health and welfare, competitive conditions in the United States economy, production of like or directly competitive articles in the United States, or United States consumers. No petitions for review of the ID were received.

The Commission has determined not to review the subject ID and has determined to issue a consent order. The Commission has further determined that Hyundai's declaration is now moot given the termination of DTL, the final remaining respondent in this investigation. Finally, the Commission has determined to request briefing on the issues of remedy, bonding, and the public interest.

Section 337(g)(1) (19 U.S.C. 1337(g)(1)) and Commission Rule 210.16(c) (19 CFR 210.16(c)) authorize the Commission, upon request, to issue a limited exclusion order or a cease and desist order or both against a respondent found in default, unless after consideration of the public interest factors in Section 337(g)(1), it finds that such relief should not issue. Accordingly, in connection with the final disposition of this investigation, the Commission is interested in receiving written submissions that address the form of remedy, if any, that should be ordered with respect to the Defaulting Respondents, identified above. If a party seeks exclusion of an article from entry into the United States for purposes other than entry for consumption, the party should so indicate and provide information establishing that activities involving other types of entry either are adversely affecting it or likely to do so. For background, see *Certain Devices for Connecting Computers via Telephone Lines*, Inv. No. 337-TA-360, USITC Pub. No. 2843, Comm'n Op. at 7-10 (December 1994).

The statute requires the Commission to consider the effects of that remedy upon the public interest. The public interest factors the Commission will consider include the effect that an exclusion order and/or cease and desist orders would have on (1) the public health and welfare, (2) competitive conditions in the U.S. economy, (3) U.S. production of articles that are like or directly competitive with those that are subject to investigation, and (4) U.S. consumers. The Commission is therefore interested in receiving written submissions that address the aforementioned public interest factors in the context of this investigation.

If the Commission orders some form of remedy, the U.S. Trade

Representative, as delegated by the President, has 60 days to approve, disapprove, or take no action on the Commission's action. See Presidential Memorandum of July 21, 2005, 70 FR 43251 (July 26, 2005). During this period, the subject articles would be entitled to enter the United States under bond, in an amount determined by the Commission and prescribed by the Secretary of the Treasury. The Commission is therefore interested in receiving submissions concerning the amount of the bond that should be imposed if a remedy is ordered.

Written Submissions: Complainants, interested government agencies, and any other interested persons are encouraged to file written submissions on the issues of remedy, the public interest, and bonding.

In their initial submission, complainants are requested to identify the form of the remedy sought and to submit proposed remedial orders for the Commission's consideration. Complainants are also requested to state the date that the asserted patents expire, the HTSUS subheadings under which the products at issue are imported, and to supply the identification information for all known importers of the products at issue in this investigation. Initial written submissions regarding remedy, bonding, and the public interest and proposed remedial orders must be filed no later than close of business on April 9, 2020. Reply submissions must be filed no later than the close of business on April 16, 2020. No further submissions on these issues will be permitted unless otherwise ordered by the Commission.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above. The Commission's paper filing requirements are currently waived. Submissions should refer to the investigation number ("Inv. No. 337-TA-1160") in a prominent place on the cover page and/or the first page. (See Handbook on Filing Procedures, https://www.usitc.gov/documents/handbook_on_filing_procedures.pdf). Persons with questions regarding filing should contact the Secretary at (202) 205-2000.

Any person desiring to submit a document to the Commission in confidence must request confidential treatment unless the information has already been granted such treatment during the proceedings. All such requests should be directed to the Secretary of the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 210.6. Documents for which confidential treatment by the

Commission is sought will be treated accordingly. A redacted non-confidential version of the document must also be filed simultaneously with any confidential filing. All information, including confidential business information and documents for which confidential treatment is properly sought, submitted to the Commission for purposes of this Investigation may be disclosed to and used: (i) By the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel¹ solely for cybersecurity purposes. All non-confidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.

The authority for the Commission's determination is contained in Section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.

Issued: March 26, 2020.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2020-06713 Filed 3-31-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Bureau of Alcohol, Tobacco, Firearms and Explosives

[OMB Number 1140-0018]

Agency Information Collection Activities; Proposed eCollection Activities; Proposed eComments Requested; Revision of a Currently Approved Collection Application for Federal Firearms License—ATF Form 7 (5310.12)/7 CR (5310.16)

AGENCY: Bureau of Alcohol, Tobacco, Firearms and Explosives, Department of Justice.

ACTION: 60-Day notice.

SUMMARY: The Department of Justice (DOJ), Bureau of Alcohol, Tobacco, Firearms and Explosives (ATF), will submit the following information collection request to the Office of

¹ All contract personnel will sign appropriate nondisclosure agreements.

Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed collection OMB 1140–0018 (Application for Federal Firearms License—ATF Form 7 (5310.12)/7 CR (5310.16), is being revised to include modifications to the verbiage used in Part B and the Instructions/Definitions section of the form.

DATES: Comments are encouraged and will be accepted for 60 days until June 1, 2020.

FOR FURTHER INFORMATION CONTACT: If you have additional comments, regarding the estimated public burden, associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions, or additional information, please contact: Tracey Robertson, ATF Federal Firearms Licensing Center, either by mail at 244 Needy Road, Martinsburg, WV 25405, by email at Tracey.Robertson@atf.gov, or by telephone at 304–616–4647.

SUPPLEMENTARY INFORMATION: 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;
- Evaluate whether and if so how the quality, utility, and clarity of the information to be collected can be enhanced; 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* (check justification or form 83): Revision of a currently approved collection.

2. *The Title of the Form/Collection:* Application for Federal Firearms License.

3. *The agency form number, if any, and the applicable component of the*

Department sponsoring the collection: Form number (if applicable): ATF Form 7 (5310.12)/7 CR (5310.16). *Component:* Bureau of Alcohol, Tobacco, Firearms and Explosives, 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 (if applicable): Individuals or households.

Abstract: The Application for Federal Firearms License—ATF Form 7 (5310.12)/7 CR (5310.16) is used by members of the public to apply for all types of federal firearm licenses (FFLs). The information requested on the form is used to determine the eligibility of the applicant to obtain a FFL, and verify the identity of a responsible person (RP).

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 13,000 respondents will utilize the form annually, and it will take each respondent approximately one (1) hour to complete their responses.

6. *An estimate of the total public burden (in hours) associated with the collection:* The estimated annual public burden associated with this collection is 13,000 hours, which is equal to 13,000 (# of respondents) * 1 (# of responses per respondent) * 1 (60 minutes).

7. *An Explanation of the Change in Estimates:* The adjustments associated with this information collection include a decrease in the total respondents and responses by 2,000, since the last renewal in 2017. However, due to an increase in the postal rate, the total mailing costs for this IC has also risen by \$100 since 2017.

If additional information is required contact: Melody Braswell, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, 3E.405A, Washington, DC 20530.

Dated: March 27, 2020.

Melody Braswell,

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

[FR Doc. 2020–06805 Filed 3–31–20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–612]

Importer of Controlled Substances Application: Wildlife Laboratories, Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before May 1, 2020. Such persons may also file a written request for a hearing on the application on or before May 1, 2020.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All request for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on February 11, 2020, Wildlife Laboratories, Inc., 1230 W Ash Street Unit D Windsor, Colorado 80550, applied to be registered as an importer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Etorphine HCL ..	9059	II
Thiafentanil	9729	II

The company plans to import the listed controlled substances for distribution to its customers.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–06759 Filed 3–31–20; 8:45 am]

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