if requested using the procedures found in 14 CFR 39.19. In accordance with 14 CFR 39.19, send your request to your principal inspector or local Flight Standards District Office, as appropriate. If sending information directly to the manager of the certification office, send it to the attention of the person identified in Related Information. You may email your request to *ANE-AD-AMOC@faa.gov*.

(2) Before using any approved AMOC, notify your appropriate principal inspector, or lacking a principal inspector, the manager of the local flight standards district office/certificate holding district office.

(j) Related Information

For more information about this AD, contact Nicholas Paine, Aviation Safety Engineer, ECO Branch, FAA, 1200 District Avenue, Burlington, MA 01803; phone: (781) 238–7116; fax: (781) 238–7199; email: *nicholas.j.paine@faa.gov*.

(k) Material Incorporated by Reference

None.

Issued on December 28, 2020.

Lance T. Gant,

Director, Compliance & Airworthiness Division, Aircraft Certification Service.

[FR Doc. 2021–00053 Filed 1–5–21; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF COMMERCE

Bureau of Industry and Security

15 CFR Part 774

[Docket No. 201215–0342]

RIN 0694–AH89

Technical Amendments to the Export Administration Regulations: Export Control Classification Number 0Y521 Series Supplement—Extension of Software Specially Designed To Automate the Analysis of Geospatial Imagery Classification

AGENCY: Bureau of Industry and Security, Commerce.

ACTION: Interim final rule; technical amendment.

SUMMARY: On January 6, 2020, the Bureau of Industry and Security (BIS) amended the Export Administration Regulations (EAR) to add Software Specially Designed to Automate the Analysis of Geospatial Imagery to the 0Y521 Temporary Export Control Classification Numbers (ECCN) Series as 0D521. In this action BIS extends that status for a year pursuant to the 0Y521 series extension procedures.

DATES: This rule is effective January 6, 2021.

FOR FURTHER INFORMATION CONTACT: Aaron Amundson, Director, Information Technology Division, Office of National Security and Technology Transfer Controls, at email *Aaron.Amundson@bis.doc.gov* or by phone at (202) 482–5299.

SUPPLEMENTARY INFORMATION:

Background

On January 6, 2020, the Bureau of Industry and Security (BIS) amended the Export Administration Regulations (EAR) with an interim final rule to add Software Specially Designed to Automate the Analysis of Geospatial Imagery to the 0Y521 Temporary Export Control Classification Numbers (ECCN) Series as 0D521. More specifically, the software was described as Geospatial imagery ''software'' ''specially designed'' for training a Deep Convolutional Neural Network to automate the analysis of geospatial imagery and point clouds. *See* 85 FR 459.

BIS established the ECCN 0Y521 series in a final rule published April 13, 2012 (72 FR 22191) (hereinafter ''April 13 rule'') to identify items that warrant control on the Commerce Control List (CCL) but are not yet identified in an existing ECCN. Items in the 0Y521 series of ECCNs are added upon a determination by the Department of Commerce, with the concurrence of the Departments of Defense and State, and other agencies as appropriate, that the items warrant control for export because the items may provide a significant military or intelligence advantage to the United States or because foreign policy reasons justify control. The ECCN 0Y521 series is a temporary holding classification.

Under the procedures established in the April 13 rule and codified at § 742.6(a)(8)(iii) of the EAR, items classified under ECCN 0Y521 remain so classified for one year from the date they are listed in supplement no. 5 to part 774 of the EAR, unless the items are re-classified under a different ECCN or the 0Y521 classification is extended.

BIS may extend an item's ECCN 0Y521 classification for two one-year periods, provided that the U.S. Government has submitted a proposal to the relevant multilateral regime(s) (*e.g.,* the Wassenaar Arrangement) to obtain multilateral controls over the item, with the understanding that multilateral controls are preferable when practical. Further extension beyond three years may occur only if the Under Secretary for Industry and Security makes a determination that such extension is in the national security or foreign policy interest of the United States. Any extension or re-extension of control of an ECCN 0Y521 item, including the determination by the Under Secretary, shall be published in the **Federal Register**.

In this action, BIS extends the status of an item classified under a 0Y521 ECCN for a year consistent with procedures that allow such an extension. Specifically, in this case the U.S. Government submitted a proposal for multilateral control of the 0D521 software specially designed to automate the analysis of geospatial imagery, as described in the January 6, 2020 interim final rule, to the relevant multilateral regime (the Wassenaar Arrangement) in a timely manner, within the first year of the item's 0D521 classification. However, due to the pandemic, the regime did not convene and therefore did not consider acceptance of the proposal. An extension of time is appropriate in order for the U.S. Government to continue its effort at the Wassenaar Arrangement in 2021.

Export Control Reform Act of 2018

On August 13, 2018, the President signed into law the John S. McCain National Defense Authorization Act for Fiscal Year 2019, which included the Export Control Reform Act of 2018 (ECRA) (50 U.S.C. 4801–4852) that provides the legal basis for BIS's principal authorities and serves as the authority under which BIS issues this rule.

Rulemaking Requirements

1. Executive Orders 13563 and 12866 direct agencies to assess all costs and benefits of available regulatory alternatives and, if regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety effects, distribute impacts, and equity). Executive Order 13563 emphasizes the importance of quantifying both costs and benefits, of reducing costs, of harmonizing rules, and of promoting flexibility. This interim final rule has been designated to be not significant for purposes of Executive Order 12866. The requirements of Executive Order 13771 do not apply because the rule is not significant.

2. Notwithstanding any other provision of law, no person is required to respond to, nor is subject to a penalty for failure to comply with a collection of information, subject to the requirements of the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*) (PRA), unless that collection of information displays a currently valid OMB control number. This rule does not involve any collection of information.

3. This rule does not contain policies associated with Federalism as that term is defined under Executive Order 13132.

4. Pursuant to section 1762 of ECRA (see 50 U.S.C. 4821), this action is exempt from the Administrative Procedure Act requirements (under 5 U.S.C. 553) for notice of proposed rulemaking, opportunity for public participation, and delay in effective date. This rule only updates Supplement No. 5 to Part 774 to the EAR by extending the date of the period of validity of 0D521 software in Supplement No. 5 to Part 774 for one year. This revision is merely technical and in accordance with established 0Y521 ECCN series procedure and purpose, which was proposed to the public and subject of comment. This rule clarifies information, which serves to avoid confusing readers about the 0D521 item's status. It does not alter any right, obligation or prohibition that applies to any person under the EAR.

5. Because a notice of proposed rulemaking and an opportunity for public comment are not required to be given for this rule by 5 U.S.C. 553, or by any other law, the analytical requirements of the Regulatory Flexibility Act, 5 U.S.C. 601, *et seq.,* are not applicable. Accordingly, no regulatory flexibility analysis is required, and none has been prepared.

List of Subjects in 15 CFR Part 774

Exports, Reporting and recordkeeping requirements.

Accordingly, part 774 of the Export Administration Regulations (15 CFR parts 730 through 774) is amended as follows:

PART 774—THE COMMERCE CONTROL LIST

■ 1. The authority citation for part 774 continues to read as follows:

Authority: 50 U.S.C. 4801–4852; 50 U.S.C. 4601 *et seq.;* 50 U.S.C. 1701 *et seq.;* 10 U.S.C. 8720; 10 U.S.C. 8730(e); 22 U.S.C. 287c, 22 U.S.C. 3201 *et seq.;* 22 U.S.C. 6004; 42 U.S.C. 2139a; 15 U.S.C. 1824; 50 U.S.C. 4305; 22 U.S.C. 7201 *et seq.;* 22 U.S.C. 7210; E.O. 13026, 61 FR 58767, 3 CFR, 1996 Comp., p. 228; E.O. 13222, 66 FR 44025, 3 CFR, 2001 Comp., p. 783.

Supplement No. 5 to Part 774 [Amended]

■ 2. In Supplement No. 5 to part 774, amend the table, under the heading ''0D521. Software'' entry No 1, by revising the date in the third column to read: ''January 6, 2022''.

Matthew S. Borman,

Deputy Assistant Secretary for Export Administration.

[FR Doc. 2020–28776 Filed 1–5–21; 8:45 am]

BILLING CODE 3510–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 101

[Docket No. FDA–2000–N–0011]

Uniform Compliance Date for Food Labeling Regulations

AGENCY: Food and Drug Administration, Department of Helath and Human Services (HHS).

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA or we) is establishing January 1, 2024, as the uniform compliance date for food labeling regulations that are published on or after January 1, 2021, and on or before December 31, 2022. We periodically announce uniform compliance dates for new food labeling requirements to minimize the economic impact of labeling changes.

DATES: This rule is effective January 6, 2021. Submit either electronic or written comments on the final rule by March 8, 2021.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

• *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*

• If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

• *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

• For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2000–N–0011 for ''Uniform Compliance Date for Food Labeling Regulations.'' Received comments, those filed in a timely manner, will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

• Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' We will review this copy, including the claimed confidential information, in our consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments, and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://*