

which could result in fuel leakage and reduced structural integrity of the wing.

(f) Compliance

Comply with this AD within the compliance times specified, unless already done.

(g) Requirements

Except as specified in paragraphs (h) and (i) of this AD: Comply with all required actions and compliance times specified in, and in accordance with, Brazilian AD 2019–06–01.

(h) Optional Terminating Action

Accomplishing the installation of doublers reinforcement on the wing forward and rear lower skin panel, in accordance with the Accomplishment Instructions of Embraer Service Bulletin SB190–57–0056, dated December 5, 2019, terminates the repetitive inspections required by this AD, as specified in Brazilian AD 2019–06–01.

(i) Exceptions to Brazilian AD 2019–06–01

For purposes of determining compliance with the requirements of this AD:

(1) Where Brazilian AD 2019–06–01 refers to its effective date, this AD requires using the effective date of this AD.

(2) The “Alternative method of compliance (AMOC)” section of Brazilian AD 2019–06–01 does not apply to this AD.

(3) Where paragraph (a)(1) of Brazilian AD 2019–06–01 specifies an initial inspection time, this AD requires an initial inspection at the applicable time specified in paragraph (i)(3)(i) or (ii) of this AD, whichever occurs later.

(i) Before the accumulation of 17,000 total flight cycles or 27,000 total flight hours, whichever occurs first.

(ii) Within 680 flight cycles or 900 flight hours after the effective date of this AD, whichever occurs first.

(4) Where paragraph (a)(1)(ii) of Brazilian AD 2019–06–01 specifies to do a special detailed inspection (SDI) in case of any “signal” of cracks, this AD requires doing an SDI before further flight after the detection of any “sign” of structural cracks in the inspected area.

(j) Other FAA AD Provisions

The following provisions also apply to this AD:

(1) *Alternative Methods of Compliance (AMOCs)*: The Manager, International Section, Transport Standards Branch, FAA, has the authority to approve AMOCs for this AD, 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 International Section, send it to the attention of the person identified in paragraph (k) of this AD. Information may be emailed to: 9-ANM-116-AMOC-REQUESTS@faa.gov. 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.

(2) *Contacting the Manufacturer*: For any requirement in this AD to obtain instructions

from a manufacturer, the instructions must be accomplished using a method approved by the Manager, International Section, Transport Standards Branch, FAA; or ANAC; or ANAC’s authorized Designee. If approved by the ANAC Designee, the approval must include the Designee’s authorized signature.

(k) Related Information

For more information about this AD, contact Krista Greer, Aerospace Engineer, International Section, Transport Standards Branch, FAA, 2200 South 216th St., Des Moines, WA 98198; telephone and fax 206–231–3221; email krista.greer@faa.gov.

(l) Material Incorporated by Reference

(1) The Director of the Federal Register approved the incorporation by reference (IBR) of the service information listed in this paragraph under 5 U.S.C. 552(a) and 1 CFR part 51.

(2) You must use this service information as applicable to do the actions required by this AD, unless this AD specifies otherwise.

(i) Agência Nacional de Aviação Civil National Civil Aviation Agency (ANAC) Brazilian AD 2019–06–01, effective June 17, 2019.

(ii) Embraer Service Bulletin SB190–57–0056, dated December 5, 2019.

(3) For information about Brazilian AD 2019–06–01, contact National Civil Aviation Agency, Aeronautical Products Certification Branch (GGCP), Rua Laurent Martins, n° 209, Jardim Esplanada, CEP 12242–431—São José dos Campos—SP, Brazil; telephone 55 (12) 3203–6600; email pac@anac.gov.br; internet www.anac.gov.br/en/. You may find this IBR material on the ANAC website at <https://sistemas.anac.gov.br/certificacao/DA/DAE.asp>. For information about Embraer service information, contact Embraer S.A., Technical Publications Section (PC 060), Av. Brigadeiro Faria Lima, 2170—Putim—12227–901 São José dos Campos—SP—Brazil; telephone +55 12 3927–5852 or +55 12 3309–0732; fax +55 12 3927–7546; email distrib@embraer.com.br; internet <http://www.flyembraer.com>.

(4) You may view this material at the FAA, Transport Standards Branch, 2200 South 216th St., Des Moines, WA. For information on the availability of this material at the FAA, call 206–231–3195. This material may be found in the AD docket on the internet at <https://www.regulations.gov> by searching for and locating Docket No. FAA–2019–0701.

(5) You may view this material that is incorporated by reference at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, email fedreg.legal@nara.gov, or go to: <https://www.archives.gov/federal-register/cfr/ibr-locations.html>.

Issued on February 25, 2020.

Lance T. Gant,

Director, Compliance & Airworthiness Division, Aircraft Certification Service.

[FR Doc. 2020–06793 Filed 4–1–20; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF COMMERCE

Bureau of Industry and Security

15 CFR Parts 732 and 734

[Docket No. 200312–0076]

RIN 0694–AF47

Control of Firearms, Guns, Ammunition and Related Articles the President Determines No Longer Warrant Control Under the United States Munitions List (USML); Notifying the Public of the Bureau’s Interim Measures With Respect to March 6, 2020 Court Order

AGENCY: Bureau of Industry and Security, Department of Commerce.

ACTION: Notification of court order.

SUMMARY: The Bureau of Industry and Security (BIS) is publishing this notification to alert the public of the Bureau’s interim measures with respect to a court order issued on March 6, 2020.

DATES: The court order was effective March 6, 2020.

FOR FURTHER INFORMATION CONTACT:

Steven Clagett, Office of Nonproliferation Controls and Treaty Compliance, Nuclear and Missile Technology Controls Division, tel. (202) 482–1641 or email steven.clagett@bis.doc.gov.

SUPPLEMENTARY INFORMATION: On March 6, 2020, the Honorable Richard A. Jones, District Judge of the U.S. District Court for the Western District of Washington issued an order enjoining the U.S. Department of State from implementing or enforcing the regulation entitled International Traffic In Arms Regulations: U.S. Munitions List Categories I, II, and III, 85 FR 3819 (Jan. 23, 2020) “insofar as it alters the status quo restrictions on technical data and software directly related to the production of firearms or firearm parts using a 3D-printer or similar equipment.” (Case No. 2:20–cv–00111–RA)).

As a result, any request for licenses of items that would otherwise fall under the U.S. Department of Commerce regulation, 15 CFR 732.2(b) and 734.7(c) (added by the final rule, entitled, Control of Firearms, Guns, Ammunition and Related Articles the President Determines No Longer Warrant Control Under the United States Munitions List (USML); 85 FR 4136, Jan. 23, 2020), should instead be directed to the U.S. Department of State.

BIS posted information on its website to alert the public of the Bureau’s

interim measures with respect to this court order. See https://www.bis.doc.gov/index.php/component/docman/?task=doc_download&gid=2535. For additional information about the court ordered injunction pertaining to revisions to the U.S. https://www.pmdc.state.gov/ddtc_public?id=ddtc_public_portal_news_and_events&timeframe=week.

Dated: March 16, 2020.

Matthew S. Borman,

Deputy Assistant Secretary for Export Administration.

[FR Doc. 2020-05934 Filed 4-1-20; 8:45 am]

BILLING CODE 3510-33-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 5, 801, 803, 807, 814, 820, 821, 822, 830, 860, 884, 900, and 1002

[Docket No. FDA-2020-N-0011]

Medical Devices; Technical Amendments

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is amending its medical device regulations. These revisions are necessary to reflect changes to the Agency's Center for Devices and Radiological Health's organizational structure, including the reorganization of its offices. The revisions replace references to the obsolete offices and positions with the current information, update the physical addresses for such offices, and correct inaccurate citations. In addition, as part of this effort we made other editorial non-substantive changes to correct other addresses, references, and citations, as appropriate. The rule does not impose any new regulatory requirements on affected parties. This action is editorial in nature and is intended to improve the accuracy of the Agency's regulations.

DATES: This rule is effective April 1, 2020.

FOR FURTHER INFORMATION CONTACT: Madhusoodana Nambiar, Office of Policy, Center for Devices and Radiological Health, 10903 New Hampshire Ave., Bldg. 66, Rm. 5519, Silver Spring, MD 20993-0002, 301-796-5837.

SUPPLEMENTARY INFORMATION:

I. Background

The FDA Center for Devices and Radiological Health (CDRH) has reorganized (84 FR 22854, May 20, 2019) to create an agile infrastructure that can adapt to future organizational, regulatory, and scientific needs. The goal of this change is to implement more efficient, consistent work processes across CDRH that better support and advance CDRH's public health mission and vision. The reorganization will integrate CDRH's premarket and postmarket program functions along product lines, allowing experts to leverage their knowledge to optimize decision making across the product life cycle. Implementation took a phased approach starting on March 18, 2019, and was completed on September 30, 2019.

Historically, CDRH has been organized according to the stage of the product's life cycle, e.g., premarket review, postmarket surveillance, and compliance, rather than by the type of product regulated. The reorganization integrates these functions by product type within the Office of Product Evaluation and Quality (OPEQ). OPEQ was formed by combining the Office of Compliance, the Office of Device Evaluation, the Office of Surveillance and Biometrics, and the Office of In Vitro Diagnostics and Radiological Health into one super office focused on a Total Product Lifecycle approach to medical device oversight. Within OPEQ, there are offices divided by product type, referred to as Offices of Health Technology (OHT), as well as cross-cutting offices focusing on specific policy and programmatic needs including the Office of Regulatory Programs and the Office of Clinical Evidence and Analysis. In addition, the reorganization established the Office of Policy, which includes two teams, the Guidance, Legislation and Special Projects Team and the Regulatory Documents and Special Projects Team, with no changes in the functions for CDRH Policy. The reorganization also established the Office of Strategic Partnerships and Technology Innovation (OST), which combined the Science and Strategic Partnerships, Digital Health, Health Informatics and Innovation teams. There are no changes in functions within the different OST teams. CDRH reorganization also realigned Management Services within the Center to ensure administrative functions in CDRH are optimally aligned, structured, and deliver excellent service. The reorganization streamlined the Center's communication functions, by combining the internal

and external communication functions, including CDRH Executive Secretary and Speaker Liaison, into the renamed Division of Communication in the Office of Communication and Education, and created an Internal Communication Branch. The structure of the Office of Science and Engineering Laboratories remains unchanged.

As part of this effort, we are also making other editorial non-substantive changes to correct other addresses, references, and citations, as appropriate.

II. Description of the Technical Amendments

The regulations specified in this rule have been revised to replace all references to "Office of Device Evaluation", "Office of Compliance", "Office of Surveillance and Biometrics" with "Office of Product Evaluation and Quality," and where, appropriate, we have used the term "Office," "Division," "Team" or "Office of Health Technology" to reflect the responsible unit within CDRH. We have also made conforming edits, as appropriate. In addition, because of the reorganization, the physical location for many of the offices changed, and thus, we have made non-substantive amendments to ensure that the room numbers and addresses reflect the current information, and other changes as necessary to update outdated addresses, references, and citations in the regulations pertaining to medical devices. The rule does not impose any new regulatory requirements on affected parties. The amendments are editorial in nature and should not be construed as modifying any substantive standards or requirements.

III. Notice and Public Comment

Publication of this document constitutes final action of these changes under the Administrative Procedure Act (5 U.S.C. 553). Section 553 of the Administrative Procedure Act (APA) exempts "rules of agency organization, procedure, or practice" from proposed rulemaking (i.e., notice and comment rulemaking). 5 U.S.C. 553(b)(3)(A). Rules are also exempt when an agency finds "good cause" that notice and comment rulemaking procedures would be "impracticable, unnecessary, or contrary to the public interest." 5 U.S.C. 553(b)(3)(B).

FDA has determined that this rulemaking meets the notice and comment exemption requirements in 5 U.S.C. 553(b)(3)(A) and (B). FDA's revisions make technical or non-substantive changes that pertain solely to the CDRH reorganization and office move and do not alter any substantive