aircraft in air commerce by prescribing regulations for practices, methods, and procedures the Administrator finds necessary for safety in air commerce. This regulation is within the scope of that authority.

Regulatory Findings

The FAA determined that this AD will not have federalism implications under Executive Order 13132. This AD will not have a substantial direct effect on the States, on the relationship between the national government and the States, or on the distribution of power and responsibilities among the various levels of government.

For the reasons discussed above, I certify that this AD:

(1) Is not a ''significant regulatory action'' under Executive Order 12866,

(2) Will not affect intrastate aviation in Alaska, and

(3) Will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 39

Air transportation, Aircraft, Aviation safety, Incorporation by reference, Safety.

The Amendment

Accordingly, under the authority delegated to me by the Administrator, the FAA amends 14 CFR part 39 as follows:

PART 39—AIRWORTHINESS DIRECTIVES

■ 1. The authority citation for part 39 continues to read as follows:

Authority: 49 U.S.C. 106(g), 40113, 44701.

§ 39.13 [Amended]

■ 2. The FAA amends § 39.13 by:

■ a. Removing Airworthiness Directive (AD) 2010–26–05, Amendment 39–16544 (75 FR 79952, December 21, 2010), and

■ b. Adding the following new AD:

2010–26–05R1 Dassault Aviation: Amendment 39–22621; Docket No. FAA–2023–1719; Project Identifier AD–2008–NM–202–AD.

(a) Effective Date

This AD is effective December 6, 2023.

(b) Affected AD

This AD replaces AD 2010–26–05, Amendment 39–16544 (75 FR 79952, December 21, 2010) (AD 2010–26–05).

(c) Applicability

This action applies to the airplanes identified in paragraphs (c)(1) and (2) of this AD, certificated in any category.

(1) Dassault Aviation Model Falcon 10 airplanes; Model FAN JET FALCON, FAN JET FALCON SERIES C, D, E, F, and G airplanes; and Model MYSTERE–FALCON 20–C5, 20–D5, 20–E5, and 20–F5 airplanes; all serial numbers, equipped with Liebherr or ABG-Semca pressurization outflow valves.

(2) Dassault Aviation Model MYSTERE–FALCON 200 airplanes, Model MYSTERE–FALCON 50 and MYSTERE–FALCON 900 airplanes, and FALCON 900EX airplanes; and Model FALCON 2000 and FALCON 2000EX airplanes; all serial numbers.

(d) Subject

Air Transport Association (ATA) of America Code 21, Air conditioning.

(e) Terminating Action

This AD terminates all requirements of AD 2010–26–05.

(f) Related Information

For more information about this AD, contact Tom Rodriguez, Aviation Safety Engineer, FAA, 1600 Stewart Avenue, Suite 410, Westbury, NY 11590; telephone: 206–231–3226; email: *tom.rodriguez@faa.gov.*

(g) Material Incorporated by Reference

None.

Issued on November 29, 2023.

Victor Wicklund,

Deputy Director, Compliance & Airworthiness Division, Aircraft Certification Service.

[FR Doc. 2023–26662 Filed 12–5–23; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 73

[Docket No. FAA–2023–2220; Airspace Docket No. 23–AWP–59]

RIN 2120–AA66

Amendment of Restricted Area R–2512 Holtville, CA

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule; correction.

SUMMARY: This action corrects a final rule published by the FAA in the **Federal Register** on November 16, 2023, that amends restricted area R–2512 in the vicinity of Holtville, CA.

DATES: Effective date 0901 UTC, January 25, 2024. The Director of the Federal Register approves this incorporation by reference action under 1 CFR part 51, subject to the annual revision of FAA Order 7400.10 and publication of conforming amendments.

ADDRESSES: A copy of the Notice of Proposed Rulemaking (NPRM), all comments received, the final rule, this final rule correction, and all background material may be viewed online at *www.regulations.gov* using the FAA Docket number. Electronic retrieval help and guidelines are available on the website. It is available 24 hours each day, 365 days each year.

FAA Order JO 7400.11H, Airspace Designations and Reporting Points, and subsequent amendments can be viewed online at *www.faa.gov/air_traffic/publications/.* You may also contact the Rules and Regulations Group, Office of Policy, Federal Aviation Administration, 800 Independence Avenue SW, Washington, DC 20591; telephone: (202) 267–8783.

FOR FURTHER INFORMATION CONTACT: Steven Roff, Rules and Regulations Group, Office of Policy, Federal Aviation Administration, 800 Independence Avenue SW, Washington, DC 20591; telephone: (202) 267–8783.

SUPPLEMENTARY INFORMATION:

History

The FAA published a final rule in the **Federal Register** for Docket No. FAA–2023–2220 (88 FR 78636; November 16, 2023), that amended restricted area R–2512 in the vicinity of Holtville, CA. The section of 14 CFR part 73 to be amended by the final rule was incorrectly stated as 73.22. The correct section of 14 CFR part 73 to be amended is 73.25. This rule corrects this typographical error.

Correction to Final Rule

Accordingly, pursuant to the authority delegated to me, the airspace amendment reflected in Docket No. FAA–2023–2220, as published in the **Federal Register** of November 16, 2023 (88 FR 78636), FR Doc. 2023–25347, is corrected as follows:

§ 73.25 [Amended]

■ 2. Section 73.25 is amended as follows:

* * * * *

R–2512 Holtville, CA [Amended]

Boundaries. Beginning at lat. 33°05′00″ N, long. 115°17′33″ W; to lat. 33°00′00″ N, long. 115°13′33″ W; to lat. 32°51′00″ N, long. 115°05′33″ W; to lat. 32°51′00″ N, long. 115°17′03″ W; to lat. 32°58′00″ N, long. 115°17′33″ W; to lat. 33°05′00″ N, long. 115°20′03″ W; to the point of beginning.

Designated altitudes. Surface to 23,000 feet MSL.

Time of designation. 0600–2300 local time daily; other times by NOTAM 24 hours in advance.

Controlling agency. FAA, Los Angeles ARTCC.

Using Agency. U.S. Marine Corps, Commanding Officer, Marine Corps Air Station Yuma, Yuma, AZ.

* * * * *

Issued in Washington, DC, on November 30, 2023.

Karen Chiodini,

Acting Manager, Rules and Regulations Group.

[FR Doc. 2023–26706 Filed 12–5–23; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510, 516, 520, 522, 524, and 558

[Docket No. FDA–2023–N–0002]

New Animal Drugs; Approval of New Animal Drug Applications; Withdrawal of Approval of New Animal Drug Applications; Change of Sponsor; Change of Sponsor Address

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendments.

SUMMARY: The Food and Drug Administration (FDA or we) is amending the animal drug regulations to reflect application-related actions for new animal drug applications (NADAs), abbreviated new animal drug applications (ANADAs), and conditionally approved new animal drug applications (CNADAs) during July, August, and September 2023. The animal drug regulations are also being amended to improve their accuracy and readability.

DATES: This rule is effective December 6, 2023.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–402–5689, *George.Haibel@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION:

I. Approvals

FDA is amending the animal drug regulations to reflect approval actions for NADAs, ANADAs, and CNADAs during July, August, and September 2023, as listed in table 1. In addition, FDA is informing the public of the availability, where applicable, of documentation of environmental review required under the National Environmental Policy Act (NEPA) and, for actions requiring review of safety or effectiveness data, summaries of the basis of approval (FOIA Summaries) under the Freedom of Information Act (FOIA). These documents, along with marketing exclusivity and patent information, may be obtained at *AnimalDrugs@FDA: https://animaldrugsatfda.fda.gov/adafda/views/#/search*.

TABLE 1—ORIGINAL AND SUPPLEMENTAL NADAS, ANADAS, AND CNADAS APPROVED DURING JULY, AUGUST, AND SEPTEMBER 2023 REQUIRING EVIDENCE OF SAFETY AND/OR EFFECTIVENESS

Approval date	File No.	Sponsor	Product name	Effect of the action	21 CFR section
July 6, 2023	200–752	Cronus Pharma Specialties India Private Ltd., Sy No-99/1, M/s GMR Hyderabad Aviation SEZ Ltd., Mamidipalli Village, Shamshabad Mandal, Ranga Reddy, Hyderabad, Telangana, 501218, India.	DEXMEDVET (dexmedetomidine hydrochloride) Injectable Solution.	Original approval as a sedative, analgesic, and preanesthetic in dogs and cats as a generic copy of NADA 141–267.	522.558
July 11, 2023	200–753	Do	CROPAMEZOLE (atipamezole hydrochloride) Injectable Solution.	Original approval for reversal of sedation and analgesia in dogs as a generic copy of NADA 141–033.	522.147
July 19, 2023	141–554	Boehringer Ingelheim Animal Health USA, Inc., 3239 Satellite Blvd., Duluth, GA30096.	NEXGARD PLUS (afoxolaner, moxidectin, and pyrantel chewable tablets).	Original approval for the prevention, treatment, and control of internal and external parasites in dogs.	520.35
August 3, 2023	200–755	Felix Pharmaceuticals Pvt. Ltd., 25–28 North Wall Quay, Dublin 1, Ireland.	Firocoxib Chewable Tablets.	Original approval for the control of pain and inflammation associated with osteoarthritis and for the control of postoperative pain and inflammation associated with soft-tissue and orthopedic surgery as a generic copy of NADA 141–230.	520.928