§ 39.13 [Amended]

2. Section 39.13 is amended by adding the following new airworthiness directive:

2001–26–10 Airbus Industrie: Amendment 39–12574. Docket 2001–NM–354–AD.

Applicability: Model A319, A320, and A321 series airplanes, certificated in any category, having manufacturer serial numbers 1035 and 1384 inclusive.

Note 1: This AD applies to each airplane identified in the preceding applicability provision, regardless of whether it has been modified, altered, or repaired in the area subject to the requirements of this AD. For airplanes that have been modified, altered, or repaired so that the performance of the requirements of this AD is affected, the owner/operator must request approval for an alternative method of compliance in accordance with paragraph (c) of this AD. The request should include an assessment of the effect of the modification, alteration, or repair on the unsafe condition addressed by this AD; and, if the unsafe condition has not been eliminated, the request should include specific proposed actions to address it.

Compliance: Required as indicated, unless accomplished previously.

To prevent failure of the oxygen containers to deliver oxygen to the passengers in the event of a rapid decompression or cabin depressurization; accomplish the following:

Inspection, Installation, and Other Actions

(a) Within 600 flight hours after the effective date of this AD, do an in-situ one-time detailed visual inspection of Dräeger Type I (three/four mask) oxygen containers, located in the passenger service units, and Drä eger Type II (two-mask) oxygen containers, located in the utility areas, for the presence of foam pads, per Airbus Service Bulletin A320–35–1022, dated June 27, 2001.

Note 2: For the purposes of this AD, a detailed visual inspection is defined as: ''An intensive visual examination of a specific structural area, system, installation, or assembly to detect damage, failure, or irregularity. Available lighting is normally supplemented with a direct source of good lighting at intensity deemed appropriate by the inspector. Inspection aids such as mirror, magnifying lenses, etc., may be used. Surface cleaning and elaborate access procedures may be required.''

(1) If all foam pads are installed, before further flight, complete the other actions (including repacking the masks in the correct position; checking the masks, tubes, and lanyards for correct stowage; and doing a manual release test and an operational test) specified in the Accomplishment Instructions of the service bulletin to ensure proper operation of the masks.

(2) If any foam pad is missing, before further flight, install a foam pad in the applicable oxygen container, and complete the other actions (including repacking the masks in the correct position; checking the masks, tubes, and lanyards for correct stowage; and doing a manual release test and an operational test) specified in the Accomplishment Instructions of the service bulletin to ensure proper operation of the masks.

Spares

(b) As of the effective date of this AD, no person shall install on any airplane a Dräeger Type I or Dräeger Type II oxygen container unless it has been inspected and other actions done per Airbus Service Bulletin A320–35–1022, dated June 27, 2001.

Alternative Methods of Compliance

(c) An alternative method of compliance or adjustment of the compliance time that provides an acceptable level of safety may be used if approved by the Manager, International Branch, ANM–116, Transport Airplane Directorate, FAA. Operators shall submit their requests through an appropriate FAA Principal Maintenance Inspector, who may add comments and then send it to the Manager, International Branch, ANM–116

Note 3: Information concerning the existence of approved alternative methods of compliance with this AD, if any, may be obtained from the International Branch, ANM–116.

Special Flight Permits

(d) Special flight permits may be issued in accordance with sections 21.197 and 21.199 of the Federal Aviation Regulations (14 CFR 21.197 and 21.199) to operate the airplane to a location where the requirements of this AD can be accomplished.

Incorporation by Reference

(e) The actions shall be done in accordance with Airbus Service Bulletin A320–35–1022, dated June 27, 2001. This incorporation by reference was approved by the Director of the Federal Register in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Copies may be obtained from Airbus Industrie 1 Rond Point, Maurice Bellonte, 31707 Blagnac Cedex, France. Copies may be inspected at the FAA, Transport Airplane Directorate, 1601 Lind Avenue, SW., Renton, Washington; or at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC.

Note 4: The subject of this AD is addressed in French airworthiness directive 2001–363(B), dated August 8, 2001.

Effective Date

(f) This amendment becomes effective on January 11, 2002

Issued in Renton, Washington, on December 17, 2001.

Kalene C. Yanamura,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 01–31549 Filed 12–26–01; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Airspace Docket No. 01–ACE–7]

Amendment to Class E Airspace; Ankeny, IA

AGENCY: Federal Aviation Administration, DOT.

ACTION: Direct final rule; confirmation of effective date.

SUMMARY: This document confirms the effective date of a direct final rule which revises Class E airspace at Ankeny, IA.

EFFECTIVE DATE: 0901 UTC, December 27, 2001.

FOR FURTHER INFORMATION CONTACT: Kathy Randolph, Air Traffic Division, Airspace Branch, ACE–52OC, DOT Regional Headquarters Building, Federal Aviation Administration, 901 Locust, Kansas City, MO 64106; telephone: (816) 329–2525.

SUPPLEMENTARY INFORMATION: The FAA published this direct final rule with a request for comments in **Federal Register** on September 24, 2001 (66 FR 48794). The FAA uses the direct final rulemaking procedure for a non-controversial rule where the FAA believes that there will be no adverse public comment. This direct final rule advised the public that no adverse comments were anticipated, and that unless a written adverse comment, or a written notice of intent to submit such an adverse comment, were received within the comment period, the regulation would become effective on December 27, 2001. Adverse comments were received, and thus this document confirms that this direct final rule will become effective on that date.

Issued in Kansas City, MO on December 18, 2001.

Herman J. Lyons, Jr.,

Manager, Air Traffic Division, Central Region.

[FR Doc. 01–31727 Filed 12–26–01; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Chapter I

Change of Address; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending its regulations to remove references to certain room numbers that no longer are valid because the address of the Center for Food Safety and Applied Nutrition (CFSAN) changed to 5100 Paint Branch Pkwy., College Park, MD, on December 14, 2001. FDA also is amending its regulations to remove a reference to an alternate site for submissions of documents to the Docket Management Branch. This alternate site is no longer available effective December 14, 2001. This action is editorial in nature and is intended to improve the accuracy and clarity of the agency's regulations.

EFFECTIVE DATE: December 14, 2001.

FOR FURTHER INFORMATION CONTACT: Louis B. Brock, Office of Regulations and Policy, Center for Food Safety and Applied Nutrition (HFS–24), 5100 Paint Branch Pkwy., College Park, MD, 20740–3835, 301–436–2378.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of November 6, 2001 (66 FR 56034), FDA amended its regulations to reflect that on December 14, 2001, CFSAN's address was changed to 5100 Paint Branch Pkwy., College Park, MD 20740. This change of address will make room numbers cited in certain regulations invalid. Therefore, FDA is amending its regulations in 21 CFR parts 73, 101, 172, 173, 177, 178, and 184 to remove the phrase ''rm. 3321'' wherever it appears. FDA also is amending 21 CFR 10.20(f) to add a period after ''20857'' and to remove the following phrase: '', except that a submission which is required to be received by the Branch by a specified date may be delivered in person to the FDA building in Washington (Room 6819, 200 C Street, SW., Washington, DC 20204) and will be considered as received by the Branch on the date on which it is delivered.'' This alternate site, originally provided for the convenience of the public, is no longer available because CFSAN's address has changed to College Park.

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedure are unnecessary because FDA is merely correcting nonsubstantive errors.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR chapter I is amended as follows:

1. Section 10.20(f) is amended by adding a period after ''20857'' and by removing the rest of the sentence.

2. Parts 73, 101, 172, 173, 177, 178, and 184 are amended by removing the words ''rm. 3321'' wherever they appear.

Dated: December 19, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–31714 Filed 12–26–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

New Animal Drugs; Change of Sponsor's Address

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor's address for Phibro Animal Health.

DATES: This rule is effective December 27, 2001.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209.

SUPPLEMENTARY INFORMATION: Phibro Animal Health, One Parker Plaza, Fort Lee, NJ 07024, has informed FDA of a change of sponsor's address to 710 Rt. 46 East, suite 401, Fairfield, NJ 07004. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to reflect the change.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects in 21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 510 is amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

2. Section 510.600 is amended in the table in paragraph (c)(1) by revising the entry for ''Phibro Animal Health'' and in the table in paragraph (c)(2) by revising the entry for ''066104'' to read as follows:

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

* * * * *

(c) * * *

(1) * * *

Firm name and address	Drug labeler code
* * * * * * *	
Phibro Animal Health, 710 Rt. 46 East, suite 401, Fairfield, NJ 07004 ..	066104
* * * * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * * * *	
066104	Phibro Animal Health, 710 Rt. 46 East, suite 401, Fairfield, NJ 07004