2003–14–04 Transport Category Airplanes: Amendment 39–13223. Docket 2003–NM–152–AD.

Applicability: The airplanes listed in Table 1 of this AD, certificated in any category. Table 1 of this AD follows:

TABLE 1.—AFFECTED AIRPLANE MODELS

Airplane manufacturer	Airplane model	As listed in C & D Aerospace Service Bulletin
Boeing	737–200, –300, –400, –500, –600, –700, –800, and –900 series.	B221001–52–03, Revision 3, dated March 25, 2003.
Boeing	757–200 and –300 series	B231001–52–02, Revision 4, dated March 19, 2003.
McDonnell Douglas	DC–10–10F, DC–10–30, DC–10–30F, DC–10–40, MD–10–30F, MD–11, and MD–11F.	B211200–52–02, Revision 1, dated June 3, 2003.

Compliance: Required as indicated, unless accomplished previously.

To prevent inadvertent release of the decompression latch and consequent opening of the decompression panel in the reinforced flight deck door, which could result in the decompression panel hitting and injuring an airplane crewmember, if the crewmember is in close proximity to the flight deck door when the decompression panel opens, accomplish the following:

Note 1: Where there are differences between this AD and the referenced service bulletins, this AD prevails.

Modification

(a) Within 90 days after the effective date of this AD, modify the reinforced flight deck door according to paragraph (a)(1), (a)(2), or (a)(3) of this AD, as applicable.

(1) For Boeing Model 737–200, –300, –400, –500, –600, –700, –800, and –900 series airplanes: Modify the upper and lower pressure relief latch assemblies on the flight deck door by doing all actions specified in and according to paragraphs 3.A., 3.B., and 3.C. of the Accomplishment Instructions of C & D Aerospace Service Bulletin B221001–52–03, Revision 3, dated March 25, 2003. One latch strap should be installed at the bottom of the upper pressure relief assembly, and a second latch strap should be installed at the top of the lower pressure relief assembly. When properly installed, the strap should cover a portion of the latch hook.

(2) For Boeing Model 757–200 and –300 series airplanes: Modify the upper and lower pressure relief latch assemblies on the flight deck door by doing all actions specified in and according to paragraphs 3.A., 3.B., and 3.C. of the Accomplishment Instructions of C & D Aerospace Service Bulletin B231001–52–02, Revision 4, dated March 19, 2003. One latch strap should be installed at the bottom of the upper pressure relief assembly, and a second latch strap should be installed at the top of the lower pressure relief assembly. When properly installed, the strap should cover a portion of the latch hook.

(3) For McDonnell Douglas DC–10–10F, DC–10–30, DC–10–30F, DC–10–40, MD–10–30F, MD–11, and MD–11F airplanes: Install spacers in the upper and lower pressure relief latch assemblies of the flight deck door, by doing all actions specified and according to paragraphs 3.A., 3.C., and 3.D. of C & D Aerospace Service Bulletin B211200–52–02, Revision 1, dated June 3, 2003. One latch strap should be installed at the bottom of the upper pressure relief assembly, and a second latch strap should be installed at the top of the lower pressure relief assembly. When properly installed, the strap should cover a portion of the latch hook.

Modifications Accomplished Per Previous Issues of Service Bulletin

(b) Modifications accomplished before the effective date of this AD per a service bulletin listed in paragraph (b)(1), (b)(2), or (b)(3) of this AD; as applicable; are considered acceptable for compliance with the corresponding action specified in paragraph (a) of this AD.

(1) For Boeing Model 737–200, –300, –400, –500, –600, –700, –800, and –900 series airplanes: C & D Aerospace Service Bulletin B221001–52–03, dated December 6, 2002; Revision 1, dated January 2, 2003; or Revision 2, dated February 20, 2003.

(2) For Boeing Model 757–200 and –300 series airplanes: C & D Aerospace Service Bulletin B231001–52–02, dated December 6, 2002; Revision 1, dated January 2, 2003; Revision 2, dated February 20, 2003; or Revision 3, dated March 7, 2003.

(3) For McDonnell Douglas DC–10–10F, DC–10–30, DC–10–30F, DC–10–40, MD–10–30F, MD–11, and MD–11F airplanes: C & D Aerospace Service Bulletin B211200–52–02, dated April 30, 2003.

Parts Installation

(c) As of the effective date of this AD, no person may install, on any airplane, a reinforced flight deck door having any part number listed in the paragraph 1.A. of C & D Aerospace Service Bulletin B221001–52–03, Revision 3, dated March 25, 2003; B231001–52–02, Revision 4, dated March 19, 2003; or B211200–52–02, Revision 1, dated June 3, 2003; as applicable; unless the door has been modified as required by paragraph (a) of this AD.

Alternative Methods of Compliance

(d) In accordance with 14 CFR 39.19, the Manager, Los Angeles Aircraft Certification Office (ACO), FAA, is authorized to approve alternative methods of compliance for this AD.

Incorporation by Reference

(e) Unless otherwise specified in this AD, the actions shall be done in accordance with C & D Aerospace Service Bulletin B211200–52–02, Revision 1, dated June 3, 2003; C & D Aerospace Service Bulletin B221001–52–03, Revision 3, dated March 25, 2003; or C & D Aerospace Service Bulletin B231001–52–02, Revision 4, dated March 19, 2003; as applicable. 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 Boeing Commercial Airplane Group, P.O. Box 3707, Seattle, Washington 98124–2207; or C & D Aerospace, 5701 Bolsa Avenue, Huntington Beach, California 92647–2063. Copies may be inspected at the FAA, Transport Airplane Directorate, 1601 Lind Avenue, SW., Renton, Washington; at the FAA, Los Angeles Aircraft Certification Office, 3960 Paramount Boulevard, Lakewood, California; or at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC.

Effective Date

(f) This amendment becomes effective on July 25, 2003.

Issued in Renton, Washington, on July 2, 2003.

Vi L. Lipski,

Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 03–17311 Filed 7–9–03; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 520

Oral Dosage Form New Animal Drugs; Phenylbutazone Tablets and Boluses

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of an abbreviated new animal drug application (ANADA) filed by West-Ward Pharmaceutical Corp. The ANADA provides for oral use of phenylbutazone tablets in horses for relief of inflammatory conditions

associated with the musculoskeletal system.

DATES: This rule is effective July 10, 2003.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–8549, e-mail: *lluther@cvm.fda.gov.*

SUPPLEMENTARY INFORMATION: West-Ward Pharmaceutical Corp., 465 Industrial Way West, Eatontown, NJ 07724, filed ANADA 200–323 for the oral use of Phenylbutazone Tablets in horses for relief of inflammatory conditions associated with the musculoskeletal system. West-Ward Pharmaceutical's Phenylbutazone Tablets are approved as a generic copy of Boehringer Ingelheim Vetmedica's BIZOLIN (phenylbutazone) Tablets, approved under NADA 99–618. The ANADA is approved as of March 28, 2003, and the regulations are amended in 21 CFR 520.1720a to reflect the approval and current format. The basis of approval is discussed in the freedom of information summary.

In addition, West-Ward Pharmaceutical Corp., has not been previously listed in the animal drug regulations as a sponsor of an approved application. At this time, 21 CFR 510.600(c) is being amended to add entries for the firm.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR part 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

FDA has determined under 21 CFR 25.33(a)(1) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

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 21 CFR Part 510

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

21 CFR Part 520

Animal drugs.

■ 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 parts 510 and 520 are 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 alphabetically adding a new entry for ''West-Ward Pharmaceutical Corp.'' and in the table in paragraph (c)(2) by numerically adding a new entry for ''000143'' 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
* * *	* *
West-Ward Pharmaceutical Corp., 465 Industrial Way West, Eatontown, NJ 07724.	000143
* * *	* *

(2) * * *

Drug labeler code	Firm name and address
* *	* * *
000143	West-Ward Pharmaceutical Corp., 465 Industrial Way West, Eatontown, NJ 07724
* *	* * *

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 3. The authority citation for 21 CFR part 520 continues to read as follows:

Authority: 21 U.S.C. 360b.

■ 4. Section 520.1720a is amended by adding paragraph (b)(5) to read as follows:

§ 520.1720a Phenylbutazone tablets and boluses.

* * * * *

(b) * * *

(5) No. 000143 for use of 1-gram tablets in horses.

* * * * *

Dated: June 26, 2003.

Andrew J. Beaulieu,

Acting Director, Center for Veterinary Medicine.

[FR Doc. 03–17439 Filed 7–9–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Salinomycin, Chlortetracycline, and Roxarsone

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of an abbreviated new animal drug application (ANADA) filed by Pennfield Oil Co. The ANADA provides for the use of single-ingredient Type A medicated articles containing salinomycin, chlortetracycline, and roxarsone to make three-way combination drug Type C medicated feeds for broiler chickens.

DATES: This rule is effective July 10, 2003.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–8549, e-mail: *lluther@cvm.fda.gov.*

SUPPLEMENTARY INFORMATION: Pennfield Oil Co., 14040 Industrial Rd., Omaha, NE 68144, filed ANADA 200–355 for use of PENNCHLOR (chlortetracycline), salinomycin, and roxarsone Type A medicated articles to make three-way combination drug Type C medicated feeds for broiler chickens. Pennfield Oil Co.'s ANADA 200–355 is approved as a generic copy of Alpharma, Inc.'s NADA 140–867. The ANADA is approved as of March 31, 2003, and the regulations are amended in 21 CFR 558.550 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9