AD No.	Amendment No.	Federal Register citation	Date of publication
93–17–07	39–8678	58 FR 45827.	Aug. 31, 1993.
93–03–14	39–8518	58 FR 14513.	Mar. 18, 1993.
93–24–51	39–8439	57 FR 60118.	Dec. 18, 1992.

(d) 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, Seattle Aircraft Certification Office (ACO), FAA, Transport Airplane Directorate. Operators shall submit their requests through an appropriate FAA Principal Maintenance Inspector, who may add comments and then send it to the Manager, Seattle ACO.

Note 2: Information concerning the existence of approved alternative methods of compliance with this AD, if any, may be obtained from the Seattle ACO.

(e) Special flight permits may be issued in accordance with §§ 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.

(f) The modification, inspections, checks, and correction of discrepancies shall be done in accordance with Boeing Alert Service Bulletin 747–54A2156, dated December 15, 1994. 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. 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.

(g) This amendment becomes effective on July 28, 1995.

Issued in Renton, Washington, on June 16, 1995.

Darrell M. Pederson,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 95–15300 Filed 6–27–95; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 558

Animal Drugs, Feeds, and Related Products; Nicarbazin Type A Medicated Article

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 Planalquimica Industrial Ltda. The ANADA provides for the use of a nicarbazin-containing Type A medicated article in making Type C medicated chicken feeds for the prevention of coccidiosis.

EFFECTIVE DATE: June 28, 1995.

FOR FURTHER INFORMATION CONTACT: Melanie R. Berson, Center for Veterinary Medicine (HFV–135), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1643.

SUPPLEMENTARY INFORMATION: Planalquimica Industrial Ltda., Rua das Magnolias nr. 2405, Jardim das Bandeiras, CEP 13053–120, Campinas, Sao Paulo, Brazil, has filed ANADA 200–027, which provides for the use of a nicarbazin-containing Type A medicated article in making Type C medicated chicken feeds as an aid in preventing outbreaks of cecal and intestinal coccidiosis.

Planalquimica's ANADA 200–027 for a 113.5-gram-per-pound nicarbazin Type A medicated article (Nicarmix) is approved as a generic copy of Merck Research Laboratories' NADA 9–476 for Nicarb®. The ANADA is approved as of June 28, 1995, and the regulations are amended in 21 CFR 510.600(c) and 558.366(a) to reflect the approval. The basis for approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of part 20 (21 CFR part 20) and § 514.11(e)(2)(ii) (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, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

The agency has carefully considered the potential environmental effects of this action. FDA has concluded that the action will not have a significant impact on the human environment, and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects

21 CFR Part 510

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

21 CFR Part 558

Animal drugs, Animal feeds.

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 558 are amended as follows:

PART 510—NEW ANIMAL DRUGS

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

Authority: Secs. 201, 301, 501, 502, 503, 512, 701, 721 of the Federal Food, Drug, and Cosmetic Act (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 ''Planalquimica Industrial Ltda.'' and in the table in paragraph (c)(2) by numerically adding a new entry for ''060728'' 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
* * * * * * *	
Planalquimica Industrial Ltda., Rua das Magnolias nr. 2405, Jardim das Bandeiras, CEP 13053–120, Campinas, Sao Paulo, Brazil	060728
* * * * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * * * *	
060728	Planalquimica Industrial Ltda., Rua das Magnolias nr. 2405, Jardim das Bandeiras, CEP 13053–120, Campinas, Sao Paulo, Brazil
* * * * * * *	

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

3. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.366 [Amended]

4. Section 558.366 *Nicarbazin* is amended in paragraph (a) by removing the phrase ''000006 and 000986'' and adding in its place ''000006, 000986, and 060728''.

Dated: June 21, 1995.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 95–15768 Filed 6–27–95; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

29 CFR Parts 1910, 1915, and 1926

RIN: 1218–AB25

Occupational Exposure to Asbestos

AGENCY: Occupational Safety and Health Administration, Department of Labor.

ACTION: Final rule; extension of start-up dates for compliance.

SUMMARY: OSHA is extending the start-up dates for some provisions of the asbestos standards until October 1, 1995 to give the public more time to implement compliance.

DATES: These amendments take effect on June 28, 1995. For Part 1910—General Industry, for § 1910.1001, the start-up dates for compliance for paragraph (d)(2)—initial monitoring, for paragraph (e)—regulated area, for paragraph (f)—methods of compliance, for paragraph (g)—respiratory protection, for paragraph (i)—hygiene facilities, for paragraph (j)—communication of hazards, and for paragraph (l)—medical surveillance are extended to October 1, 1995. For Part 1915—Shipyards, for § 1915.1001, the start-up dates for compliance for paragraph (g)—methods of compliance, for paragraph (h)—respiratory protection, for paragraph (j)—hygiene facilities, for paragraph (k)—communication of hazards, for paragraph (l)—housekeeping, for paragraph (m)—medical surveillance and for paragraph (o)—qualified person are extended to October 1, 1995. For Part 1926—Construction, for § 1926.1101, the start-up dates for compliance for paragraph (g)—methods of compliance, for paragraph (h)—respiratory protection, for paragraph (j)—hygiene facilities, for paragraph (k)—communication of hazards, for paragraph (l)—housekeeping, for paragraph (m)—medical surveillance and for paragraph (o)—competent person are extended to October 1, 1995.

FOR FURTHER INFORMATION CONTACT: Mr. Richard Liblong, Director of Information and Consumer Affairs, Occupational Safety and Health Administration, U.S. Department of Labor, Room N3647, 200 Constitution Avenue NW., Washington, DC 20210, telephone (202) 219–8151.

SUPPLEMENTARY INFORMATION: OSHA issued improved asbestos standards for general industry, construction and shipyards on August 10, 1994 at 59 FR 40964 to better protect workers from lung cancer, asbestosis and other diseases caused by asbestos exposure. The new standards took effect on October 11, 1994 and that date was the start-up date for some provisions such as the new lower exposure limit of 0.1 f/cc. However, various other provisions such as the new medical surveillance, respiratory protection and training provisions and the engineering control requirements had start-up dates from between January 9, 1995 and April 10, 1995.

Various members of the public have requested that OSHA grant more time for the public to comply with some provisions. OSHA is publishing a correction and clarification notice and various compliance and training materials to assist in the understanding of the new standard.

OSHA has concluded that it is appropriate to give the public additional time to implement some of the provisions of the new asbestos standards which may require more time for implementation. Other provisions such as the new exposure limit had a start-up date of October 11, 1994 and again, OSHA is not extending the start-up date of those provisions. In the interim, the provisions of the preexisting asbestos standards remain in effect for those provisions of the new standards whose start-up dates have been extended.

The provisions extended are listed above in the **DATES** section of this preamble. OSHA has concluded that October 1, 1995 is a reasonable time for employers to fully come into compliance with the provisions for