

TITLE 20—EMPLOYEES' BENEFITS**CHAPTER V—EMPLOYMENT AND TRAINING ADMINISTRATION, DEPARTMENT OF LABOR**

1. Part 655 of chapter V of title 20, Code of Federal Regulations, is amended as follows:

PART 655—TEMPORARY EMPLOYMENT OF ALIENS IN THE UNITED STATES

a. The authority citation for part 655 continues to read as follows:

Authority: Section 655.0 issued under 8 U.S.C. 1101(a)(15)(H) (i) and (ii), 1182 (m) and (n), 1184, 1188, and 1288(c); 29 U.S.C. 49 *et seq.*; sec. 3(c)(1), Pub. L. 101-238, 103 Stat. 2099, 2103 (8 U.S.C. 1182 note); sec. 221(a), Pub. L. 101-649, 104 Stat. 4978, 5027 (8 U.S.C. 1184 note); and 8 CFR 214.2(h)(4)(i).

Section 665.00 issued under 8 U.S.C. 1101(a)(15)(H)(ii), 1184, and 1188; 29 U.S.C. 49 *et seq.*; and 8 CFR 214.2(h)(4)(i).

Subparts A and C issued under 8 U.S.C. 1101(a)(15)(H)(ii)(b) and 1184; 29 U.S.C. 49 *et seq.*; and 8 CFR 214.2(h)(4)(i).

Subpart B issued under 8 U.S.C. 1101(a)(15)(H)(ii)(a), 1184, and 1188; and 29 U.S.C. 49 *et seq.*

Subparts D and E issued under 8 U.S.C. 1101(a)(15)(H)(i)(a), 1182(m), and 1184; 29 U.S.C. 49 *et seq.*; and sec. 3(c)(1), Pub. L. 101-238, 103 Stat. 2099, 2103 (8 U.S.C. 1182 note).

Subparts F and G issued under 8 U.S.C. 1184 and 1288(c); and 29 U.S.C. 49 *et seq.*

Subparts H and I issued under 8 U.S.C. 1101(a)(15) (H) (i)(b), 1182(n), and 1184; and 29 U.S.C. 49 *et seq.*

Subparts J and K issued under 29 U.S.C. 49 *et seq.*; and sec. 221(a), Pub. L. 101-649, 104 Stat. 4978, 5027 (8 U.S.C. 1184 note).

b. Part 655 is amended as set forth in the Joint Interim Final Rule, which appears at the end of the end of the common preamble.

TITLE 29—LABOR**CHAPTER V—WAGE AND HOUR DIVISION, DEPARTMENT OF LABOR**

2. Part 508 of chapter V of title 29, Code of Federal Regulations, is amended as follows:

PART 508—ATTESTATIONS FILED BY EMPLOYERS UTILIZING F-1 STUDENTS FOR OFF-CAMPUS WORK

a. The authority citation for part 508 continues to read as follows:

Authority: 29 U.S.C. 49 *et seq.*; and sec. 221(a), Pub. L. 101-649, 104 Stat. 4978, 5027 (8 U.S.C. 1184 note).

b. Part 508 is amended as set forth in the Joint Interim Final Rule, which appears at the end of the end of the common preamble.

Signed at Washington, DC, this 27th day of June, 1995.

Raymond Uhalde,

Deputy Assistant Secretary for Employment and Training.

Maria Echaveste,

Administrator, Wage and Hour Division, Employment Standards Administration.

[FR Doc. 95-16133 Filed 6-27-95; 12:52 pm]

BILLING CODE 4510-30-M

BILLING CODE 4510-27-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration****21 CFR Part 176**

[Docket No. 91F-0339]

Indirect Food Additives: Paper and Paperboard Components

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of 2-bromo-2-nitro-1,3-propanediol as an antimicrobial/preservative in fillers, pigment slurries, starch sizing solutions, and latex coatings used in the manufacture of paper and paperboard articles intended to contact food. This action responds to a food additive petition filed by Betz Laboratories, Inc.

DATES: Effective June 30, 1995; written objections and requests for a hearing by August 29, 1995.

ADDRESSES: Submit written objections to the Dockets Management Branch (HFA-305), Food and Drug Administration, rm. 1-23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Richard H. White, Center for Food Safety and Applied Nutrition (HFS-216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3094.

SUPPLEMENTARY INFORMATION: In a notice published in the *Federal Register* of September 19, 1991 (56 FR 47478), FDA announced that a food additive petition (FAP 1B4279) had been filed by Betz Laboratories, Inc., 4636 Somerton Rd., Trevese, PA 19053. The petition proposed to amend the food additive regulations in § 176.170 *Components of paper and paperboard in contact with aqueous and fatty foods* (21 CFR 176.170) to provide for the safe use of 2-bromo-2-nitro-1,3-propanediol as an antimicrobial/preservative in fillers,

binders, pigment slurries, sizings, and coatings used in the manufacture of paper and paperboard articles intended for food-contact use. The petitioner later limited and clarified the requested use of the additive to provide for its safe use as an antimicrobial/preservative in fillers, pigment slurries, starch sizing solutions, and latex coatings used as components of paper and paperboard in contact with aqueous and fatty foods.

FDA has evaluated data in the petition and other relevant material. The agency concludes that the additive is safe and effective for the proposed use and that 21 CFR 176.170 of the food additive regulations should be amended as set forth below.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition by appointment with the information contact person listed above. As provided in 21 CFR 171.1(h), the agency will delete from the documents any materials that are not available for public disclosure before making the documents available for inspection.

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.

Any person who will be adversely affected by this regulation may at any time on or before July 31, 1995, file with the Dockets Management Branch (address above) written objections thereto. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any

particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this document. Any objections received in response to the regulation may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 176

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director of the Center for Food Safety and Applied Nutrition, 21 CFR Part 176 is amended as follows:

PART 176—INDIRECT FOOD ADDITIVES: PAPER AND PAPERBOARD COMPONENTS

1. The authority citation for 21 CFR part 176 continues to read as follows:

Authority: Secs. 201, 402, 406, 409, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 342, 346, 348, 379e).

2. Section 176.170 is amended in the table in paragraph (a)(5) by alphabetically adding a new entry under the headings "List of Substances" and "Limitations" to read as follows:

§ 176.170 Components of paper and paperboard in contact with aqueous and fatty foods.

* * * * *
(a) * * *
(5) * * *

List of Substances	Limitations
* * * * *	* * * * *
2-Bromo-2-nitro-1,3-propanediol (CAS Reg. No. 52-51-7)	For use only as an antimicrobial/preservative in fillers, pigment slurries, starch sizing solutions, and latex coatings at levels not to exceed 0.01 percent by weight of those components.
* * * * *	* * * * *

* * * * *

Dated: June 21, 1995.

Fred R. Shank,
Director, Center for Food Safety and Applied Nutrition.
[FR Doc. 95-16092 Filed 6-29-95; 8:45 am]
BILLING CODE 4160-01-F

21 CFR Part 510

New Animal Drugs; Change of Sponsor Name and 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 name and address from DDI Pharmaceuticals, Inc., to OXIS International, Inc.

EFFECTIVE DATE: June 30, 1995.

FOR FURTHER INFORMATION CONTACT: Benjamin A. Puyot, Center for Veterinary Medicine (HFV-130), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301-594-1646.

SUPPLEMENTARY INFORMATION: DDI Pharmaceuticals, Inc., 518 Logue Ave., Mountain View, CA 94043, has informed FDA of a change of sponsor name and address to OXIS International, Inc., 6040 N. Cutter Circle, suite 317, Portland, OR 97217-3935. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to

reflect the change of sponsor name and address.

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: 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 removing the entry for "DDI Pharmaceuticals, Inc.," and by alphabetically adding a new entry for "OXIS International, Inc.," and in the table in paragraph (c)(2) in the entry "024991" by revising the sponsor name and address 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
* * * * *	* * * * *
OXIS International, Inc., 6040 N. Cutter Circle, suite 317, Portland, OR 97217-3935.	024991
* * * * *	* * * * *

(2) * * *

Drug labeler code	Firm name and address
* * * * *	* * * * *
024991	OXIS International, Inc., 6040 N. Cutter Circle, suite 317, Portland, OR 97217-3935
* * * * *	* * * * *

Dated: June 23, 1995.

Andrew J. Beaulieu,
Acting Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.
[FR Doc. 95-16093 Filed 6-29-95; 8:45 am]
BILLING CODE 4160-01-F