the test and the basis for selecting participants.

(b) *NCAP testing.* For purposes of conducting an approved test program or procedure designed to evaluate planned components of the National Customs Automation Program (NCAP), as described in section 411(a)(2) of the Tariff Act of 1930 (19 U.S.C. 411), the Commissioner of Customs may impose requirements different from those specified in the Customs Regulations, but only to the extent that such different requirements do not affect the collection of the revenue, public health, safety, or law enforcement. In addition to the requirement of paragraph (a)(1) of this section, the imposition of any such different requirements shall be subject to the following conditions:

(1) *Prior publication requirement.* For tests affecting the NCAP, notice shall be published in the **Federal Register** not less than thirty days prior to implementing such test, followed by publication in the Customs Bulletin. The notice shall invite public comments concerning any aspect of the test program or procedure, and inform interested members of the public of the eligibility criteria for voluntary participation in the test and the basis for selecting participants; and,

(2) *Post publication requirement.* Within a reasonable time period following the completion of the test, a complete description of the results shall be published in both the **Federal Register** and the Customs Bulletin.

Approved: February 21, 1995.

George J. Weise,

Commissioner of Customs.

John P. Simpson,

Deputy Assistant Secretary of the Treasury.

[FR Doc. 95–6525 Filed 3–15–95; 8:45 am]

BILLING CODE 4820–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Social Security Administration

20 CFR Part 416

[Regulations No. 16]

RIN 0960–AC22

Supplemental Security Income for the Aged, Blind, and Disabled; Continuation of Benefits and Special Eligibility for Certain Severely Impaired Recipients Who Work; Appeal Rights Following Mass Change Resulting in Reduction, Suspension, or Termination of State Supplementary Payments; and Deemed Application Date Based on Misinformation; Correction

AGENCY: Social Security Administration, HHS.

ACTION: Correction to final rules.

SUMMARY: This document contains corrections to the final rules published in the **Federal Register** on Friday, August 12, 1994 (59 FR 41400), Monday, August 22, 1994 (59 FR 43035), and Wednesday, August 31, 1994 (59 FR 44918). We are correcting incorrect paragraph redesignations and related amendatory instructions, as well as making one technical correction to a paragraph in one regulatory section.

EFFECTIVE DATE: March 16, 1995.

FOR FURTHER INFORMATION CONTACT: Regarding this **Federal Register** document—Richard M. Bresnick, Legal Assistant, Office of Regulations, Social Security Administration, 6401 Security Boulevard, Baltimore, MD 21235, (410) 965–1758; regarding eligibility or filing for benefits—our national toll-free number, 1–800–772–1213.

SUPPLEMENTARY INFORMATION: In the final rules that appeared on page 41400 in the **Federal Register** issued of Friday, August 12, 1994, we had incorrect paragraph redesignations in amendatory item 15. We indicated that in § 416.1402 we were redesignating paragraphs (i) through (n) as paragraphs (h) through (m). However, paragraph (n) had not yet been published. Thus, paragraphs (i) through (m) should have been redesignated as paragraphs (h) through (l). The new paragraph (n) was to be contained in other final rules, ''Deemed Application Date Based on Misinformation,'' which were to be published before these rules but were not published until August 31, 1994 (59 FR 44918). We discovered this before ''Deemed Application Date Based on Misinformation'' was published, however, and changed the designation of that new paragraph in the later rules to paragraph (m) to reflect the proper redesignation which should have been made in the rules published on August 12, 1994.

In the interim, on August 22, 1994, we published other final rules, ''Appeal Rights Following Mass Change Resulting in Reduction, Suspension, or Termination of State Supplementary Payments'' (59 FR 43035), which contained a new paragraph designated paragraph (n) in § 416.1402 which would have been correct if the regulations had been published in the anticipated sequence. The amendatory item 3 in these final rules contained other incorrect paragraph designations and instructions for punctuating § 416.1402. Further, similar incorrect instructions for revising § 416.1402 were contained under part 416 in amendatory item 6 in the final rules published on August 31, 1994. Also, paragraph (m) as published in these rules was incorrectly punctuated and did not have the word ''and'' following it. Therefore, we are correcting all three amendatory items and paragraph (m) itself to reflect the correct paragraph designations, punctuation, and ending word ''and.'' With these corrections, all the paragraphs and amendatory instructions will be correct. Make the corrections as follows:

1. In the **Federal Register** issue of August 12, 1994, in the second column on page 41405, amendatory item 15 should read as follows:

15. Section 416.1402 is amended by revising paragraphs (a) and (b), removing paragraph (h), and redesignating paragraphs (i) through (m) as paragraphs (h) through (l), respectively to read as follows:

2. In the **Federal Register** issue of August 22, 1994, in the first column on page 43039, amendatory item 3 should read as follows:

3. Section 416.1402 is amended by removing the word ''and'' following the semicolon at the end of paragraph (k), replacing the period at the end of paragraph (l) with a semicolon and adding the word ''and'' after the semicolon, and adding a new paragraph (n) to read as follows:

3. In the **Federal Register** issue of August 31, 1994, in the third column on page 44927, amendatory item 6 under part 416 should read as follows:

6. Section 416.1402 is amended by removing the word ''and'' following the semicolon at the end of paragraph (l) and adding a new paragraph (m) to read as follows:

§ 416.1402 Administrative actions that are initial determinations.

* * * * *

(m) A claim for benefits under § 416.351 based on alleged misinformation; and

* * * * *

Dated: March 7, 1995.

Neil J. Stillman,

Deputy Assistant Secretary for Information Resources Management.

[FR Doc. 95–6502 Filed 3–15–95; 8:45 am]

BILLING CODE 4190–29–M

Food and Drug Administration

21 CFR Part 510

Animal Drugs, Feeds, and Related Products; 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 for a new animal drug application (NADA) from Zoecon Industries, Inc., to Sandoz Agro, Inc.

EFFECTIVE DATE: March 16, 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: Due to a merger with Zoecon Industries, Inc., 12200 Denton Dr., Dallas, TX 75234, and Sandoz Agro, Inc., 1300 East Touhy Ave., Des Plaines, IL 60018, the firms have requested that FDA publish a notice of a change of sponsor name and address for their new animal drug application NADA 98–895, Starbar GX–118 (N-(mercaptomethyl) phthalimide S-(O,O dimethylphosphorodithioate) emulsifiable liquid). 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. The drug labeler code ''011536'' for Zoecon Industries, Inc., is being retained for the new sponsor.

List of Subjects in 21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and record keeping 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 ''Zoecon Industries, Inc.,'' and alphabetically adding a new entry for ''Sandoz Agro, Inc.,'' and in the table in paragraph (c)(2) in the entry for ''011536'' 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
* * * *	*
Sandoz Agro, Inc., 1300 East Touhy Ave., Des Plaines, IL 60018	011536
* * * *	*

(2) * * *

Drug labeler code	Firm name and address
*	* * * *
011536	Sandoz Agro, Inc., 1300 East Touhy Ave., Des Plaines, IL 60018
*	* * * *

Dated: March 8, 1995.

Robert C. Livingston,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 95–6528 Filed 3–15–95; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Lincomycin Hydrochloride Soluble Powder

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 a supplemental new animal drug application (NADA) filed by The Upjohn Co. The NADA provides for the use of lincomycin hydrochloride soluble powder to make medicated swine and broiler chicken drinking water. The supplement provides for use of a packet containing the equivalent of 32 grams (g) of lincomycin in addition to the currently approved packet containing the equivalent of 16 g of lincomycin.

EFFECTIVE DATE: March 16, 1995.

FOR FURTHER INFORMATION CONTACT: David R. Newkirk, Center for Veterinary Medicine (HFV–142), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–2701.

SUPPLEMENTARY INFORMATION: The Upjohn Co., Kalamazoo, MI 49001, filed a supplement to NADA 111–636 for Lincomix (lincomycin hydrochloride) soluble powder. The supplemental NADA provides for the use of an 80-g packet containing the equivalent of 32 g of lincomycin in addition to the approved 40-g packet containing the equivalent of 16 g of lincomycin. Both packets are used to make a swine drinking water containing 250 milligrams (mg) of lincomycin per gallon used for the treatment of swine dysentery and broiler chicken drinking water containing 64 mg of lincomycin per gallon for the control of necrotic enteritis.

This supplemental NADA is approved as of February 9, 1995, and the regulations in § 520.1263c(a) (21 CFR 520.126c(a)) are amended to reflect the approval.

This is a manufacturing supplement to an approved NADA. The approval does not require a summary of safety, effectiveness data, or information. Therefore, a freedom of information summary as provided in part 20 (21 CFR part 20) and § 514.11(e)(2)(ii) (21 CFR 514.11(e)(2)(ii)) is not required.

Under section 512(c)(2)(F)(iii) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(c)(2)(F)(iii)), this approval for food-producing animals does not qualify for marketing exclusivity because the supplemental application does not contain new clinical or field investigations (other than bioequivalence or residue studies) and new human food safety studies (other than bioequivalence or residue studies) essential to the approval and conducted or sponsored by the applicant.

The agency has determined under 21 CFR 25.24(d)(1)(iii) 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.

List of Subjects in 21 CFR Part 520

Animal drugs.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under