respectively," is corrected to read, "; by removing paragraph (j), redesignating paragraphs (k), (l), and (m) as paragraphs (j), (k), and (l), respectively; and by revising the last sentence in newly redesignated paragraph (l)".

4. On page 52253, in the second column, in § 600.80, the last sentence of newly redesignated paragraph (l) is correctly revised to read as follows:

## § 600.80 Postmarketing reporting of adverse experiences.

\* \* \* \* \*

(l) \* \* \* For the purposes of this provision, this paragraph also includes any person reporting under paragraph (c)(1)(iii) of this section.

Dated: March 18, 1998.

#### William K. Hubbard,

Associate Commissioner for Policy Coordination.

 $[FR\ Doc.\ 98-7833\ Filed\ 3-25-98;\ 8:45\ am]$ 

BILLING CODE 4160-01-F

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **Food and Drug Administration**

#### 21 CFR Part 510

New Animal Drugs; Change of Sponsor 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 address for Koffolk, Inc.

FFECTIVE DATE: March 26, 1998.

FOR FURTHER INFORMATION CONTACT:
Thomas J. McKay, Center for Veterinary
Medicine (HFV–102), Food and Drug
Administration, 7500 Standish Pl.,
Rockville, MD 20855, 301–827–0213.

SUPPLEMENTARY INFORMATION: Koffolk,
Inc., One Parker Plaza, Fort Lee, NJ
07024, has informed FDA of a change of
sponsor address to P.O. Box 675935,
14735 Las Quintas, Rancho Santa Fe,
CA 92067. Accordingly, the agency is
amending the regulations in 21 CFR
510.600(c)(1) and (c)(2) to reflect the
change of sponsor 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:** 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 sponsor address for "Koffolk, Inc." and in the table in paragraph (c)(2) in the entry for "063271" by revising the sponsor address to read as follows:

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

(c) \* \* \*

(1) \* \* \*

(2) \* \* \*

| Drug labeler code |   |   | Firm name and addre | Firm name and address    |  |  |
|-------------------|---|---|---------------------|--------------------------|--|--|
| *                 | * | * | * *                 | * *                      |  |  |
| 063271            |   |   |                     | uintas, Rancho Santa Fe, |  |  |
| *                 | * | * | *                   | * *                      |  |  |

Dated: March 17, 1998.

#### Andrew J. Beaulieau,

Acting Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine. [FR Doc. 98–7835 Filed 3–25–98; 8:45 am]

BILLING CODE 4160-01-F

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Food and Drug Administration

21 CFR Part 558

New Animal Drugs For Use In Animal Feeds; Bacitracin Zinc; Correction

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule; correction.

**SUMMARY:** The Food and Drug Administration (FDA) is correcting a document that published in the Federal Register of October 23, 1997 (62 FR 55161). The document amended the animal drug regulations to reflect approval of abbreviated new animal drug application (ANADA) 200-223 filed by ALPHARMA Inc. In amending the animal drug regulations, the document did not reflect that Boehringer Ingelheim Animal Health, Inc.'s approval for use of bacitracin zinc Type A medicated articles is for chickens only. This document corrects that error.

**EFFECTIVE DATE:** October 23, 1997.

#### FOR FURTHER INFORMATION CONTACT:

David L. Gordon, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1739.

**SUPPLEMENTARY INFORMATION:** In FR Doc. 97–28015, appearing on page 55161 in the **Federal Register** of October 23, 1997, the following correction is made:

## § 558.78 [Amended]

1. On page 55162, in the second column, § 558.78 *Bacitracin zinc* is amended in paragraph (a)(3) by adding the phrase "for chickens" after the word "pound".

Dated: March 17, 1998.

### Andrew J. Beaulieau,

Acting Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine. [FR Doc. 98–7834 Filed 3–25–98; 8:45 am] BILLING CODE 4160–01–F

#### **DEPARTMENT OF THE TREASURY**

**Internal Revenue Service** 

26 CFR Parts 1 and 301

[TD 8767]

RIN 1545-AWO7

# Guidance Under Subpart F Relating to Partnerships and Branches

**AGENCY:** Internal Revenue Service (IRS), Treasury.

**ACTION:** Temporary and final regulations.

**SUMMARY:** This document contains regulations relating to the treatment under subpart F of certain payments involving branches of a controlled foreign corporation (CFC) that are treated as separate entities for foreign tax purposes or partnerships in which CFCs are partners. These regulations are necessary to provide guidance on transactions relating to such entities. These regulations will affect United States shareholders of controlled foreign corporations. The text of these temporary regulations also serves as the text of the proposed regulations published elsewhere in this issue of the Federal Register.

**DATES:** *Effective date:* These regulations are effective March 23, 1998.

Applicability date: For dates of applicability see §§ 1.904–5T(o), 1.954–1T(c)(1)(i)(E), 1.954–2T(a)(5)(iii) and (a)(6)(ii), 1.954–9T(d) and 301.7701–3T(f) of these regulations.

FOR FURTHER INFORMATION CONTACT: Valerie Mark, (202) 622–3840 (not a toll-free number).

#### SUPPLEMENTARY INFORMATION:

#### **Background**

I. In General

In these temporary regulations and in proposed regulations published elsewhere in this issue of the **Federal Register**, the Treasury and IRS set forth a framework for dealing with issues posed by the use of certain entities that are regarded as fiscally transparent for purposes of U.S. tax law, with regard to the application of subpart F of the Internal Revenue Code.

Subpart F was enacted by Congress to limit the deferral of U.S. taxation of certain income earned outside the United States by foreign corporations controlled by U.S. persons. Limited deferral was retained after the enactment of subpart F to protect the competitiveness of controlled foreign corporations (CFCs) doing business overseas. See S. Rep. No. 1881, 87th

Cong., 2d Sess. 78–80 (1962). This limited deferral furthers the objective of allowing a CFC engaged in an active business, and located in a foreign country for appropriate economic reasons, to compete in a similar tax environment with non-U.S.-owned corporations located in the same country.

Conversely, one of the purposes of subpart F is to prevent CFCs from converting active income that is not easily moveable and is earned in a jurisdiction in which a business is located for non-tax reasons, into passive, easily moveable income that is shifted to a lower tax jurisdiction primarily for tax avoidance. Moreover, when subpart F was first enacted it was realized that related person transactions can be easily manipulated to reduce both United States and foreign taxes. Consequently, in enacting subpart F, Congress provided that transactions of CFCs that involve related persons generally give rise to subpart F income with certain enumerated exceptions.

Hybrid branches, which, by definition, are not regarded as fiscally transparent under foreign law, are particularly well suited to the type of tax avoidance described above. In light of the recent proliferation of hybrid branches, Treasury and the IRS believe that it is appropriate to consider the issues related to transactions involving hybrid branches, or other hybrid entities, under subpart F.

The use of partnerships that are fiscally transparent for U.S. tax purposes raises additional issues in the context of subpart F that are similar to those raised in connection with hybrid branches. Such partnerships may or may not be fiscally transparent under foreign law. (Other fiscally-transparent entities, such as grantor trusts, will be the subject of guidance issued in conjunction with the finalization of regulations under section 672(f).)

The entity classification regulations of §§ 301.7701–1 through 301.7701–3 (the check-the-box regulations) make entity classification generally elective, in part so that taxpayers can choose a tax status that is consistent with their business objectives. This administrative provision was not intended to change substantive law. Particularly in the international area, the ability to more easily achieve fiscal transparency can lead to inappropriate results under certain substantive international provisions of the Code. Thus, the Treasury and the IRS believe that it is necessary to provide additional guidance regarding the use of hybrid entities in the international context. See