IV. References

The following information has been placed on display in the Dockets Management Branch (address above) and may be seen by interested persons between 9 a.m. and 4 p.m., Monday through Friday.


16. Letter from William A. Craig, Professor of Medicine and Pharmaceutics, University of Wisconsin, to Dockets Management Branch (HFA–305), Food and Drug Administration, dated July 31, 1996.


18. “Report from the Danish Veterinary Laboratory: The Effect of Avoparcin Used as a Feed Additive on the Occurrence of Vancomycin Resistant Enterococcus Faecium in Pig and Poultry Production,” Danish Veterinary Laboratory, Copenhagen, July 1995.

V. Request for Comments

Interested persons may, on or before July 21, 1997, submit to the Dockets Management Branch (address above) written comments regarding this document. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office between 9 a.m. and 4 p.m., Monday through Friday.

VI. Order of Prohibition

Therefore, under the Federal Food, Drug, and Cosmetic Act, and under the authority delegated to the Commissioner of Food and Drugs, I hereby issue the following order under section 2(a)(4)(D) of the AMDUCA, Pub. L. 1–3–396 (21 U.S.C. 331, 351, 352, 353, 355, 357, 360b, 371, 379e).

1. The authority citation for 21 CFR part 530 continues to read as follows:

PART 530—EXTRALABEL DRUG USE IN ANIMALS

1. The following drugs, families of drugs, and substances are prohibited for extralabel use in animals.

(a) The following drugs, families of drugs, and substances are prohibited for extralabel animal and human use in food-producing animals.

1. Chloramphenicol;

2. Clobenoterol;

3. Dextran stilbestrol (DES);

4. Dimetridazole;

5. Ipronidazole;

6. Other nitroimidazoles;

7. Furazolidone (except for approved topical use);

8. Nitrofurazone (except for approved topical use);

9. Sulfonamide drugs in lactating dairy cattle (except approved use of sulfadimethoxine, sulfamerazine, sulfadex, and sulfadimidine);

10. Fluoroquinolones; and


(b) The following drugs, families of drugs, and substances are prohibited for extralabel animal and human use in nonfood-producing animals:

1. Sulfonamide drugs in lactating dairy cattle (except approved use of sulfadimethoxine, sulfamerazine, sulfadex, and sulfadimidine).

2. Fluoroquinolones; and

3. Glycopeptides.

VII. List of Subjects in 21 CFR Part 530

A. Administrative procedure and procedure, Advertising, Animal drugs, Animal feeds, Drugs, Labeling, Prescription drugs, Reporting and recordkeeping requirements.

B. 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 530 is amended to read as follows:

PART 530—EXTRALABEL DRUG USE IN ANIMALS

1. The authority citation for 21 CFR part 530 continues to read as follows:


2. Section 530.41 is revised to read as follows:

§ 530.41 Drugs prohibited for extralabel use in animals.

(a) The following drugs, families of drugs, and substances are prohibited for extralabel use in animals.

1. Chloramphenicol;

2. Clobenoterol;

3. Dextran stilbestrol (DES);

4. Dimetridazole;

5. Ipronidazole;

6. Other nitroimidazoles;

7. Furazolidone (except for approved topical use);

8. Nitrofurazone (except for approved topical use);

9. Sulfonamide drugs in lactating dairy cattle (except approved use of sulfadimethoxine, sulfamerazine, sulfadex, and sulfadimidine).

10. Fluoroquinolones; and


(b) The following drugs, families of drugs, and substances are prohibited for extralabel use in food-producing animals:

1. Chloramphenicol;

2. Clobenoterol;

3. Dextran stilbestrol (DES);

4. Dimetridazole;

5. Ipronidazole;

6. Other nitroimidazoles;

7. Furazolidone (except for approved topical use);

8. Nitrofurazone (except for approved topical use);

9. Sulfonamide drugs in lactating dairy cattle (except approved use of sulfadimethoxine, sulfamerazine, sulfadex, and sulfadimidine).
(ACDA) is revoking its existing superseded employee responsibility and conduct regulations at 22 CFR part 606, and, in their stead, inserting cross-references to the executive branch-wide Standards, as well as to executive branch financial disclosure regulations.

**EFFECTIVE DATE:** These regulations are effective May 22, 1997.


**SUPPLEMENTARY INFORMATION:**

I. Background

On August 7, 1992, the Office of Government Ethics published the Standards of Ethical Conduct for Employees of the Executive Branch. See 57 FR 35066-35067, as corrected at 57 FR 48557 and 57 FR 52583, with additional extensions for certain existing provisions at 59 FR 4779-4780 and 60 FR 6390-6391. The executive branch-wide Standards are now codified at 5 CFR part 2635. Effective February 3, 1993, they established uniform ethical conduct standards applicable to all executive branch personnel.

ACDA is revoking the provisions of its existing standards of conduct regulations that have already been superseded or that are superseded upon issuance of this regulation and replacing them with a new section that provides a cross reference to 5 CFR parts 2634 and 2635.

II. Revocation of ACDA’s Responsibilities and Conduct Regulations

This final rule revokes ACDA’s employee responsibility and conduct regulations at 22 CFR part 606, now superseded. Some of those regulations were superseded when the confidential financial disclosure provisions of the executive branch-wide financial disclosure regulations at 5 CFR part 2634 took effect on October 5, 1992, and many others were superseded when the Standards of Ethical Conduct for Employees of the Executive Branch at 5 CFR part 2635 became effective on February 3, 1993. Others were retained in ACDA’s internal regulations since they dealt with other aspects of employee conduct such as indebtedness and political activity.

The ACDA residual standards rule replaces ACDA’s revoked ethics regulations with a cross-reference at new 22 CFR part 606 to OGE’s rules at 5 CFR parts 2634 and 2635.

### III. Matters of Regulatory Procedure

**Executive Order 12866**

In issuing this rule, ACDA has adhered to the regulatory philosophy and the applicable principles of regulation as set forth in Section 1 of Executive Order 12866, Regulatory Planning and Review. This regulation has not been reviewed by the Office of Management and Budget under that Executive Order, as it deals with agency organization, management, and personnel matters and is not, in any event, deemed “significant” thereunder.

**Paperwork Reduction Act**

ACDA has determined that the Paperwork Reduction Act (44 U.S.C. chapter 35) does not apply because the proposed regulation does not contain any information collection requirements that require the approval of the Office of Management and Budget.

**Administrative Procedure Act**

This rulemaking is related solely to ACDA’s organization, procedure, and practice. Consequently, ACDA has found that good cause exists under 5 U.S.C. 553(b)(2); 5 CFR 2634.

**Final rule.**

This rule amends the surety bond provisions of Minerals Management Service (MMS) regulations to establish December 8, 1997, as the deadline for Outer Continental Shelf (OCS) oil and gas and sulphur lessees to comply with new levels of bond coverage established in 1993. It also makes other changes that reduce the risk of default by an underfunded entity who operates a lease or holds a pipeline or other operating rights.

### § 606.1 Cross-reference to employee ethical conduct standards and financial disclosure regulations.

Employees of the United States Arms Control and Disarmament Agency (ACDA) should refer to the Standards of Ethical Conduct for Employees of the Executive Branch at 5 CFR part 2635 and the Executive Branch financial disclosure regulations at 5 CFR part 2634.

[FR Doc. 97-13390 Filed 5-21-97; 8:45 am]
BILLING CODE 6820-32-P

### DEPARTMENT OF THE INTERIOR

**Minerals Management Service**

30 CFR Parts 250, 251, 256, 281, and 282

RIN 1010-AB92

**Surety Bonds for Outer Continental Shelf Leases**

**AGENCY:** Minerals Management Service, Interior.

**ACTION:** Final rule.

**SUMMARY:** This rule amends the surety bond provisions of Minerals Management Service (MMS) regulations to establish December 8, 1997, as the deadline for Outer Continental Shelf (OCS) oil and gas and sulphur lessees to comply with new levels of bond coverage established in 1993. It also makes other changes that reduce the risk of default by an underfunded entity who operates a lease or holds a pipeline right-of-way or geological and geophysical (G&G) exploration permit to drill a deep stratigraphic test well.

**EFFECTIVE DATE:** August 20, 1997.

**FOR FURTHER INFORMATION CONTACT:** John V. Mirabella, Engineering and Operations Division, (703) 787-1607.

**SUPPLEMENTARY INFORMATION:** This rule: (1) Establishes December 8, 1997, as the deadline for every lessee to comply with the bond coverage requirements established in the rule published August 27, 1993 (58 FR 45255). (2) Clarifies our position that co-lessees and operating rights owners are...