APPENDIX E TO PART 305—ROOM AIR CONDITIONERS

[Range information]

Manufacturer's rated cooling capacity in Btu's/yr	Range of energy efficiency ratios (EERs)	
	Low	High
Without Reverse Cycle and with Louvered Sides:		
Less than 6,000 Btu	8.0	10.0
6,000 to 7,999 Btu	8.5	10.3
8,000 to 13,999 Btu	9.0	12.0
14,000 to 19,999 Btu.	8.8	10.7
20,000 and more Btu	8.2	10.0
Without Reverse Cycle and without Louvered Sides:		
Less than 6,000 Btu	(*)	(*)
6,000 to 7,999 Btu	8.5	9.6
8,000 to 13,999 Btu	8.5	9.2
14,000 to 19,999 Btu.	(*)	(*)
20,000 and more Btu	(*)	(*)
With Reverse Cycle and with Louvered Sides	8.5	11.5
With Reverse Cycle, without Louvered Sides	8.0	9.0

* No data submitted for units meeting Federal Minimum Efficiency Standards effective January 1, 1990.

Cost Information for Appendix E

When the ranges of comparability in Appendix E are used on EnergyGuide labels for room air conditioners, the estimated annual operating cost disclosure appearing in the box at the bottom of the labels must be derived using the 1995 Representative Average Unit Costs for electricity (8.67¢ per kiloWatt-hour) and the text below the box must identify the costs as such.

Appendix H to part 305 [Amended]

■ 4. In section 2 of Appendix H of Part 305, the text is amended by removing the figure ''8.60¢'' wherever it appears and by adding, in its place, the figure ''9.06¢''. In addition, the text in this section is amended by removing the figure ''12.9¢'' wherever it appears and by adding, in its place, the figure ''13.6¢''.

■ 5. The formula in section 2 of Appendix H of part 305 is revised to read as follows in both places that it appears:

Appendix H to Part 305—Cooling Performance and Cost for Central Air Conditioners

* * * * *

$$\text{Your estimated cost} = \text{Listed average annual operating cost } * \times \frac{\text{Your cooling load hours } **}{1{,}000} \times \frac{\text{Your electrical rate in cents per KWH}}{9.06¢}$$

* * * * *

Appendix I to Part 305 [Amended]

■ 6. In section 2 of Appendix I of part 305, the text is amended by removing the figure ''8.60¢'' wherever it appears and by adding, in its place, the figure ''9.06¢''. In addition, the text and formulas are amended by removing the figure ''12.90¢'' wherever it appears and by adding, in its place, the figure ''13.6¢''.

■ 7. In section 2 of Appendix I of part 305, the formula is revised to read as follows in both places that it appears:

Appendix I to Part 305—Heating Performance and Cost for Central Air Conditioners

* * * * *

$$\text{Your estimated cost} = \text{Listed annual heating cost } * \times \frac{\text{Your electrical cost in cents per KWH}}{9.06¢}$$

* * * * *

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 05–11026 Filed 6–2–05; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

New Animal Drugs; Change of Sponsor's Name

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's name from Steris Laboratories, Inc., to Watson Laboratories, Inc.

DATES: This rule is effective June 3, 2005.

FOR FURTHER INFORMATION CONTACT: David R. Newkirk, Center for Veterinary

Medicine (HFV–100), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6967, e-mail: *david.newkirk@fda.gov*.

SUPPLEMENTARY INFORMATION: Steris Laboratories, Inc., 620 North 51st Ave., Phoenix, AZ 85043–4705, has informed FDA of a change of sponsor's name to Watson Laboratories, Inc. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c) to reflect the change.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

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 removing the entry for ''Steris Laboratories, Inc.'' and by alphabetically adding an entry for ''Watson Laboratories, Inc.''; and in the table in paragraph (c)(2) by revising the entry for ''000402'' 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
* * * * *	
Watson Laboratories, Inc., 620 North 51st Ave., Phoenix, AZ 85043–4705	000402
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
000402	Watson Laboratories, Inc., 620 North 51st Ave., Phoenix, AZ 85043–4705
* * * * *	

Dated: May 11, 2005.

Steven D. Vaughn,

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

[FR Doc. 05–11030 Filed 6–2–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 520, 522, and 558

New Animal Drugs; Change of Sponsor

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 for 16 approved new animal drug applications (NADAs) from Purina Mills, Inc., to Virbac AH, Inc.

DATES: This rule is effective June 3, 2005.

FOR FURTHER INFORMATION CONTACT: David R. Newkirk, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6967, e-mail: *david.newkirk@fda.gov*.

SUPPLEMENTARY INFORMATION: Purina Mills, Inc., P.O. Box 66812, St. Louis, MO 63166–6812, has informed FDA that it has transferred ownership of, and all rights and interest in, the following 16 approved NADAs to Virbac AH, Inc., 3200 Meacham Blvd., Ft. Worth, TX 76137:

TABLE 1.

NADA No.	21 CFR Section	Trade Name
011–779	522.940	Purina Pigemia 100
013–214	558.274	Purina Hygromix-Swine
013–663	520.100a	Purina Liquid Amprol
040–205	520.2380a	Purina Horse Wormer Medicated
042–116	558.185	Purina 6-Day Worm-Kill Feed
042–660	558.630	Purina Pork-Plus Medicated
043–387	558.625	Purina Hog Plus II
046–700	558.365	Statyl
049–729	520.2261a	Purina Sulfa
097–258	558.485	Purina Ban Worm For Pigs
099–767	558.630	Purina Tylan 40 Plus Sulfamethazine
113–748	520.1182	Purina Oral Pigemia
132–574	558.325	Purina Check-R-Ton LI
135–941	558.485	Check-E-Ton BM
136–116	520.905d	Purina Worm-A-Rest Litter Pack
140–869	520.1840	Purina Bloat Block; Purina Saf-T-Block BG

Accordingly, the agency is amending the regulations in parts 520, 522, and 558 (21 CFR parts 520, 522, and 558). Sections 520.100a, 520.905d, 520.1182, 520.1840, 520.2261a, 520.2380a, 522.940, 558.185, 558.274, 558.325, 558.365, 558.485, 558.625, and 558.630 will reflect the transfer of ownership and a current format. Sections 520.1182 and 522.940 are being revised to reflect a current format.

In addition, § 520.2380a is being revised to correct the citation for the approved indications for Virbac AH's thiabendazole dewormer approved under NADA 040–205. The citation was corrected in the **Federal Register** of March 3, 1976 (41 FR 9149), but an error was reintroduced in the 1978 printing of the Code of Federal Regulations. This action is being taken to improve the accuracy of the regulations.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects

21 CFR Parts 520 and 522

Animal drugs.

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 520, 522, and 558 are amended as follows: