7A994 Other navigation direction finding equipment, airborne communication equipment, all aircraft inertial navigation systems not controlled under 7A003 or 7A103, and other avionic equipment, including parts and components, n.e.s.

* * * * *

List of Items Controlled

Unit: * * *
Related Controls: N/A.
Related Definitions: * * *
Items: * * *

* * * * *

■ 9. Supplement No. 1 to Part 774 (the Commerce Control List), Category 9—Propulsion Systems, Space Vehicles, and Related Equipment, ECCN 9A106, List of Items Controlled Section is amended by revising the ''Items'' paragraph to read as follows:

9A106 Systems or components, other than those controlled by 9A006, usable in ''missiles'', as follows (see List of Items Controlled), and specially designed for liquid rocket propulsion systems.

* * * * *

List of Items Controlled

Unit: * * *
Related Controls: * * *
Related Definitions: * * *
Items:

a. Ablative liners for thrust or combustion chambers;

b. Rocket nozzles;

c. Thrust vector control sub-systems;

Technical Note: Examples of methods of achieving thrust vector control controlled by 9A106.c includes:

1. Flexible nozzle;
2. Fluid or secondary gas injection;
3. Movable engine or nozzle;
4. Deflection of exhaust gas steam (jet vanes or probes); or
5. Thrust tabs.

d. Liquid and slurry propellant (including oxidizers) control systems, and specially designed components therefor, designed or modified to operate in vibration environments of more than 10 g rms between 20 Hz and 2000 Hz.

Note: The only servo valves and pumps controlled by 9A106.d, are the following:

a. Servo valves designed for flow rates of 24 liters per minute or greater, at an absolute pressure of 7 Mpa or greater, that have an actuator response time of less than 100 ms;

b. Pumps, for liquid propellants, with shaft speeds equal to or greater than 8,000 rpm or with discharge pressures equal to or greater than 7 Mpa.

e. Flight control servo valves designed or modified for use in ''missiles'' and designed or modified to operate in a vibration environment of more than 10g RMS over the entire range between 20Hz and 2KHz.

Dated: September 12, 2003.

Matthew Borman,

Acting Assistant Secretary for Export Administration.

[FR Doc. 03–23888 Filed 9–17–03; 8:45 am]

BILLING CODE 3510–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form 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 two approved new animal drug applications (NADAs) from Teva Pharmaceuticals USA to Delmarva Laboratories, Inc.

DATES: This rule is effective September 18, 2003.

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: *dnewkirk@cvm.fda.gov*.

SUPPLEMENTARY INFORMATION: Teva Pharmaceuticals USA, 650 Cathill Rd., Sellersville, PA 18960, has informed FDA that it has transferred ownership of, and all rights and interest in, the following two approved NADAs to Delmarva Laboratories, Inc., 1500 Huguenot Rd., suite 106, Midlothian, VA 23113:

NADA No.	Trade Name
65–492	ROBAMOX V (amoxicillin trihydrate) Tablets
65–495	ROBAMOX V (amoxicillin trihydrate)

Accordingly, the agency is amending the regulations in 21 CFR 520.88b and 520.88f to reflect the transfer of ownership.

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 520

Animal drugs.

■ 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 520 is amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 520 continues to read as follows:

Authority: 21 U.S.C. 360b.

§ 520.88b [Amended]

■ 2. Section 520.88b *Amoxicillin trihydrate for oral suspension* is amended in paragraph (c) by removing ''*Sponsor*. See Nos. 000093 and 000856'' and by adding in its place ''*Sponsors*. See Nos. 000856 and 059079''.

§ 520.88f [Amended]

■ 3. Section 520.88f *Amoxicillin trihydrate tablets* is amended in paragraph (b) by removing ''*Sponsor*. See Nos. 000093 and 000856'' and by adding in its place ''*Sponsors*. See Nos. 000856 and 059079''.

Dated: August 28, 2003.

Steven D. Vaughn,

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

[FR Doc. 03–23779 Filed 9–17–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 556 and 558

New Animal Drugs; Ractopamine

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 new animal drug application (NADA) filed by Elanco Animal Health. The NADA provides for use of ractopamine hydrochloride Type A medicated articles to make Type B and Type C medicated feeds used for increased rate of weight gain, improved feed efficiency, and increased carcass leanness in cattle fed in confinement for slaughter.

DATES: This rule is effective September 18, 2003.

FOR FURTHER INFORMATION CONTACT: Eric S. Dubbin, Center for Veterinary