(v) Look through the sight gage opening while using a flashlight pointed into the filler vent hole to verify the gears are meshed properly. Gears are properly meshed when the ''X'' marked on the pinion gear of the output cartridge is between the two ''X's'' marked on the gear of the input cartridge (see figure 3). Do not torque the MS20074–04–06 bolts until both cartridges are installed on the case and the gears are properly meshed. Torque the output cartridge bolts to 60 in.-lbs. first, then torque the input cartridge bolts to 60 in.-lbs. Safety wire with 0.032-inch stainless steel safety wire.

(vi) Reinstall sight gage with MS35769–11 or AN900–10 gasket. Oil threads to prevent threads from locking up. Torque to 200 in.-lbs.

(vii) Reinstall the chip detector with a MS35769–8 or AN900–9 gasket after lubricating the threads with oil. Torque the chip detector to 150 in.-lbs. Safety wire the sight gage to the chip detector using 0.032-inch stainless steel safety wire.

(viii) Fill the T/R gearbox with oil to the level indicated on the T/R sight glass decal. Reinstall the filler vent plug, P/N A610–1, with a MS35769–9 or AN900–8 gasket, after lubricating the threads with oil.

(ix) Inspect the T/R gearbox assembly to ensure that the shafts and gears rotate freely.

(7) Reinstall the T/R gearbox onto the helicopter in accordance with the applicable maintenance manual. Verify that the oil level of the T/R gearbox is at the recommended mark on the sight glass.

(d) An alternative method of compliance or adjustment of the compliance time that provides an acceptable level of safety may be used when approved by the Manager, Los Angeles Aircraft Certification Office, FAA. Operators shall submit their requests through an FAA Principal Maintenance Inspector, who may concur or comment and then send it to the Manager, Los Angeles Aircraft Certification Office.

Note 2: Information concerning the existence of approved alternative methods of compliance with this AD, if any, may be obtained from the Los Angeles Aircraft Certification Office.

(e) Special flight permits may be issued in accordance with sections 21.197 and 21.199 of the Federal Aviation Regulations (14 CFR 21.197 and 21.199) to operate the helicopter to a location where the requirements of this AD can be accomplished.

(f) This amendment becomes effective on December 27, 1995.

Issued in Fort Worth, Texas, on November 2, 1995.

Eric Bries,

Acting Manager, Rotorcraft Directorate, Aircraft Certification Service.

[FR Doc. 95–28537 Filed 11–21–95; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 520 and 522

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 name for three new animal drug applications (NADA's) from Vet-A-Mix, Inc., to Lloyd, Inc.

EFFECTIVE DATE: November 22, 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: Vet-A-Mix, Inc., 604 West Thomas Ave., P.O. Box A, Shenandoah, IA 51601, has informed FDA of a change of sponsor name for approved NADA's 92–836 (diethylcarbamazine citrate), 140–866 (yohimbine hydrochloride injectable), and 140–921 (prednisolone tablets) to Lloyd, Inc., 604 West Thomas Ave., Shenandoah, IA 51601. Accordingly, FDA is amending the regulations in 21 CFR 520.622c, 520.1880, and 522.2670 to reflect the change of sponsor name.

List of Subjects in 21 CFR Parts 520 and 522

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 parts 520 and 522 are 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: Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

§ 520.622c [Amended]

2. Section 520.622c *Diethylcarbamazine citrate chewable tablets* is amended in paragraph (b)(3) by removing ''011789'' and adding in its place ''061690''.

§ 520.1880 [Amended]

3. Section 520.1880 *Prednisolone tablets* is amended in paragraph (b) by removing ''011789'' and adding in its place ''061690''.

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

4. The authority citation of 21 CFR part 522 continues to read as follows:

Authority: Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

§ 522.2670 [Amended]

5. Section 522.2670 *Yohimbine injectable* is amended in paragraph (b) by removing ''032998'' and adding in its place ''061690''.

Dated: November 13, 1995.

Robert C. Livingston,

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

[FR Doc. 95–28542 Filed 11–21–95; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 522

Implantation or Injectable Dosage Form New Animal Drugs; Selenium/Vitamin E Injection

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 an abbreviated new animal drug application (ANADA) filed by Fort Dodge Laboratories. The ANADA provides for subcutaneous or intramuscular use of a selenium/vitamin E injection for prevention and treatment of selenium/tocopherol deficiency syndrome in weanling calves and breeding beef cattle.

EFFECTIVE DATE: November 22, 1995.

FOR FURTHER INFORMATION CONTACT: Melanie R. Berson, Center for Veterinary Medicine (HFV–135), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1643.

SUPPLEMENTARY INFORMATION: Fort Dodge Laboratories, 800 Fifth St. NW., P.O. Box 518, Fort Dodge, IA 50501, filed ANADA 200–109, which provides for subcutaneous or intramuscular use of Velenium™ (selenium, vitamin E) Injection for prevention and treatment of selenium/tocopherol deficiency syndrome in weanling calves and breeding beef cattle. The drug is limited to use by or on the order of a licensed veterinarian.

Approval of ANADA 200–109 for Fort Dodge's selenium/vitamin E injection is as a generic copy of Schering-Plough's Mu-Se® (selenium/vitamin E) Injection