Food and Drug Administration, HHS

§ 522.88 Sterile amoxicillin trihydrate for suspension.

(a)(1) Specifications. Each vial contains 3 grams of amoxicillin as the trihydrate. The powder is reconstituted with sterile water for injection USP to a concentration of 100 or 250 milligrams per milliliter for use as in paragraph (d) of this section.

1These conditions are NAS/NRC reviewed and deemed effective. Applications for these uses need not include effectiveness data as specified by §514.111 of this chapter, but may require bioequivalency and safety information.

Dosage can be repeated every 12 hours, as indicated.1

1Not for use in animals intended for food purposes.1

1For use only by or on the order of a licensed veterinarian.1

[40 FR 13858, Mar. 27, 1975, as amended at 63 FR 44382, Aug. 19, 1998]

§ 522.84 Beta-aminopropionitrile fumarate.

(a) Specifications. Each vial contains 7.0 milligrams of beta-aminopropionitrile fumarate sterile lyophilized powder which is reconstituted for injection with 10 milliliters of sterile physiologic saline, USP.

(b) Sponsor. See No. 064146 in §510.600(c) of this chapter.

(c) [Reserved]

(d) Conditions of use—(1) Horses—(i) Amount. 7 milligrams (10 milliliters) intralesionally every other day for 5 treatments beginning about 30 days after initial injury.

(2) [Reserved]

[63 FR 44382, Aug. 19, 1998]

§ 522.82 Aminopropazine fumarate sterile solution injection.

(a) Specifications. Each milliliter of aminopropazine fumarate sterile aqueous solution, veterinary, contains aminopropazine fumarate equivalent to 25 milligrams of aminopropazine base.

(b) Sponsor. See No. 000061 in §510.600(c) of this chapter.

(c) Conditions of use. (1) The drug is used for reducing excessive smooth muscle contractions, such as occur in urethral spasms associated with urolithiasis in cats and dogs and in colic spasms in horses.1

(2) It is administered intramuscularly or intravenously to dogs and cats at a level of 1 to 2 milligrams per pound of body weight. It is administered intramuscularly or intravenously to horses at a level of 0.25 milligrams per pound of body weight.

Dosage may be gradually increased up to a maximum of five times the suggested dosage. Following parenteral use dosage may be continued by oral administration of tablets.

(2) For use only by or on the order of a licensed veterinarian.