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- (c) Related tolerances. See §556.625 of this chapter.
- (d) Conditions of use. It is used in the drinking water of broilers, breeder flocks, and replacement chickens as follows:
 - (1) Amount. 0.03 percent.
- (2) Indications for use. Treatment of coccidiosis.
- (3) Limitations. Administer in drinking water for 3 days as sole source of drinking water and sulfonamide medication; withdraw 4 days prior to slaughter; not to be administered to chickens producing eggs for human consumption.
- [40 FR 13838, Mar. 27, 1975, as amended at 50 FR 41489, Oct. 11, 1985; 54 FR 12188, Mar. 24, 1989; 55 FR 8460, Mar. 8, 1990; 64 FR 15684, Apr. 1, 1999; 67 FR 78355, Dec. 24, 2002]

§ 520.2200 Sulfachlorpyridazine.

- (a) Specifications.—(1) Sodium sulfachlorpyridazine powder.
- (2) Each bolus contains 2 grams sulfachlorpyridazine.
- (3) Each tablet contains 250 milligrams (mg) sulfachlorpyridazine.
- (b) *Sponsor*. See No. 000010 in §510.600(c) of this chapter.
- (c) Related tolerances. See §556.630 of this chapter.
- (d) Conditions of use. It is used as follows:
- (1) Calves—(i) Amount. Administer 30 to 45 mg sulfachlorpyridazine powder per pound (/lb) of body weight per day in milk or milk replacer, or in a bolus, in divided doses twice daily for 1 to 5 days.
- (ii) *Indications for use*. For the treatment of diarrhea caused or complicated by *Escherichia coli* (colibacillosis).
- (iii) Limitations. Treated ruminating calves must not be slaughtered for food during treatment or for 7 days after the last treatment. A withdrawal period has not been established for this product in preruminating calves. Do not use in calves to be processed for veal.
- (2) Swine—(i) Amount. Administer 20 to 35 mg/lb body weight per day, in divided doses twice daily for 1 to 5 days:
 - (A) In drinking water or
- (B) For individual treatment, in an oral suspension containing 50 mg per milliliter.

- (ii) *Indications for use*. For the treatment of diarrhea caused or complicated by *E. coli* (colibacillosis).
- (iii) Limitations. Treated swine must not be slaughtered for food during treatment or for 4 days after the last treatment.
- (3) Dogs—(i) Amount. Administer tablets orally at 500 mg per 10 to 15 lb of body weight daily, in two or three divided doses.
- (ii) Indications for use. As an aid in the treatment of infectious tracheobronchitis and infections caused by E. coli, and in the treatment of infections caused by other Grampositive and Gram-negative organisms that are susceptible to sulfonamide therapy.
- (iii) *Limitations*. Federal law restricts this drug to use only by or on the order of a licensed veterinarian.

[75 FR 10166, Mar. 5, 2010]

§ 520.2215 Sulfadiazine/pyrimethamine suspension.

- (a) Specifications. Each milliliter (mL) of suspension contains 250 milligrams (mg) sulfadiazine (as the sodium salt) and 12.5 mg pyrimethamine.
- (b) *Sponsor*. See No. 068718 in §510.600(c) of this chapter.
- (c) Conditions of use in horses—(1) Amount. Administer orally 20 mg sulfadiazine per kilogram (kg) body weight and 1 mg/kg pyrimethamine daily.
- (2) Indications for use. For the treatment of equine protozoal myeloencephalitis (EPM) caused by Sarcocystis neurona.
- (3) *Limitations*. Do not use in horses intended for human consumption. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- [69 FR 70054, Dec. 2, 2004, as amended at 73 FR 53686, Sept. 17, 2008]

§ 520.2218 Sulfamerazine, sulfamethazine, and sulfaquinoxaline powder.

- (a) Specifications. Each 195-gram (g) packet of powder contains 78 g sulfamerazine, 78 g sulfamethazine, and 39 g sulfaquinoxaline.
- (b) Sponsor. See No. 046573 in \$510.600(c) of this chapter.
- (c) Related tolerances. See §§ 556.670 and 556.685 of this chapter.