(b) *Sponsors*. See Nos. 000856, 059130, and 061623 in § 510.600(c) of this chapter.

(c) *Related tolerances*. See § 556.600 of this chapter.

(d) *Conditions of use in swine*—(1) *Amount*. Administer 5 mg per pound (lb) of body weight orally twice daily for 3 to 5 days.

(2) *Indications for use*. For the treatment and control of porcine enteric colibacillosis (scours) caused by *E. coli* susceptible to spectinomycin in pigs under 4 weeks of age.

(3) *Limitations*. Do not administer to pigs over 15 lb body weight or over 4 weeks of age. Do not administer within 21 days of slaughter.

[73 FR 6607, Feb. 5, 2008]

§ 520.2130 Spinosad.

(a) *Specifications*. Each chewable tablet contains 140, 270, 560, 810, or 1620 milligrams (mg) spinosad.

(b) *Sponsor*. See No. 000986 in § 510.600 of this chapter.

(c) *Conditions of use in dogs*—(1) *Amount*. Administer tablets once a month at a recommended minimum dosage of 13.5 mg per pound (30 mg per kilogram) of body weight.

(2) *Indications for use*. To kill fleas and for the prevention and treatment of flea infestations (*Ctenocephalides felis*) on dogs for 1 month.

(3) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[72 FR 60551, Oct. 25, 2007]

§ 520.2150 Stanozolol oral dosage forms.

§ 520.2150a Stanozolol tablets.

(a) *Specifications*. Each tablet contains 2 milligrams of stanozolol.

(b) *Sponsor*. No. 000009 in § 510.600(c) of this chapter.

(c) *Conditions of use*. (1) Used as an anabolic steroid treatment in dogs and cats.

(2) Administered orally to cats and small breeds of dogs, ½ to 1 tablet twice daily for several weeks; to large breeds of dogs, 1 to 2 tablets twice daily for several weeks. The tablets may be crushed and administered in feed.

(3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[40 FR 46101, Oct. 6, 1975, as amended at 42 FR 36995, July 19, 1977. Redesignated at 50 FR 38114, Sept. 20, 1985, and amended at 55 FR 23076, June 6, 1990]

§ 520.2150b Stanozolol chewable tablets.

(a) *Specifications*. Each chewable tablet contains 2 milligrams of stanozolol.

(b) *Sponsor*. No. 000009 in § 510.600(c) of this chapter.

(c) *Conditions of use*. (1) Used as an anabolic steroid treatment in dogs.

(2) Administered orally to small breeds of dogs, ½ to 1 tablet twice daily for several weeks; to large breeds of dogs, 1 to 2 tablets twice daily for several weeks.

(3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[50 FR 38114, Sept. 20, 1985, as amended at 55 FR 23076, June 6, 1990]

§ 520.2158 Streptomycin/dihydrostreptomycin oral dosage forms.

§ 520.2158a Streptomycin sulfate oral solution.

(a) *Specifications*. Solution containing 25 percent streptomycin sulfate.

(b) *Sponsor*. See Nos. 033008 and 055462 in § 510.600(c) of this chapter.

(c) *Related tolerances*. See § 556.610 of this chapter.

(d) *Conditions of use*. Use in drinking water as follows:

(1) *Calves and swine*—(i) *Amount*. 10 to 15 milligrams per pound (mg/pound) of body weight (1.0 to 1.5 grams per gallon).

(ii) *Indications for use*. Treatment of bacterial enteritis caused by *Escherichia coli* and *Salmonella* spp. susceptible to streptomycin.

(iii) *Limitations*. Calves: Do not administer for more than 5 days. Swine: Do not administer for more than 4 days. Prepare fresh solution daily. Calves: Withdraw 2 days before slaughter. As sole source of streptomycin. Warning: Certain strains of bacteria may develop a tolerance for streptomycin. Consult a veterinarian or animal pathologist for diagnosis.

(2) *Chickens*—(i) *Amount*. 10 to 15 mg/pound of body weight (0.6 to 0.9 grams per gallon).

(ii) *Indications for use*. Treatment of nonspecific infectious enteritis caused by organisms susceptible to streptomycin.

(iii) *Limitations*. Chickens: Do not administer for more than 5 days. Withdraw 4 days before slaughter. Do not administer to chickens producing eggs for human consumption. Prepare fresh solution daily. As sole source of streptomycin. Warning: Certain strains of bacteria may develop a tolerance for streptomycin. Consult a veterinarian or animal pathologist for diagnosis.

[57 FR 37327, Aug. 18, 1992, as amended at 58 FR 47211, Sept. 8, 1993; 63 FR 51821, Sept. 29, 1998]

§ 520.2158b Dihydrostreptomycin tablets.

(a) *Specifications*. Each tablet contains 37.5 milligrams dihydrostreptomycin (as the sulfate) with 375 milligrams chlorhexidine dihydrochloride.

(b) *Sponsor*. See No. 000856 in § 510.600(c) of this chapter.

(c) *Related tolerances*. See §§ 556.120 and 556.200 of this chapter.

(d) *Conditions of use*. *Calves*—(1) *Amount*. 150 milligrams of dihydrostreptomycin and 1.5 grams of chlorhexidine dihydrochloride per 100 pounds of body weight per day.

(2) *Indications for use*. Treatment of bacterial scours in calves.

(3) *Limitations*. Administer orally once a day for 5 days; withdraw 3 days before slaughter.

[57 FR 37327, Aug. 18, 1992; 57 FR 42623, Sept. 15, 1992]

§ 520.2158c Dihydrostreptomycin oral suspension.

(a) *Specifications*. Each milliliter contains 1.25 milligrams dihydrostreptomycin (as the sulfate) with 12.5 milligrams chlorhexidine dihydrochloride.

(b) *Sponsor*. See No. 000856 in § 510.600(c) of this chapter.

(c) *Related tolerances*. See §§ 556.120 and 556.200 of this chapter.

(d) *Conditions of use*. *Calves*—(1) *Amount*. 150 milligrams of dihydrostreptomycin and 1.5 grams of chlorhexidine dihydrochloride per 100 pounds of body weight per day.

(2) *Indications for use*. Treatment of bacterial scours in calves.

(3) *Limitations*. Administer orally once a day for 5 days; withdraw 3 days before slaughter.

[57 FR 37327, Aug. 18, 1992]

§ 520.2160 Styrylpyridinium, diethylcarbamazine oral dosage forms.

§ 520.2170 Sulfabromomethazine sodium boluses.

(a) *Specifications*. Each bolus contains 15 grams of sulfabromomethazine sodium.

(b) *Related tolerance*. See § 556.620 of this chapter.

(c) *Sponsor*. See No. 050604 in § 510.600(c) of this chapter.

(d) *NAS/NRC status*. These conditions of use are NAS/NRC reviewed and found effective. NADA's for these uses need not include effectiveness data as specified by § 514.111 of this chapter, but may require bioequivalency and safety information.

(e) *Conditions of use*. *Cattle*—(1) *Amount*. 90 milligrams per pound body weight.

(2) *Indications for use*. Treatment of necrotic pododermatitis (foot rot) and calf diphtheria caused by *Fusobacterium necrophorum;* colibacillosis (scours) caused by *Escherichia coli;* bacterial pneumonia and bovine respiratory disease complex (shipping fever complex) associated with *Pasteurella* spp.; acute metritis and acute mastitis caused by *Streptococcus* spp.

(3) *Limitations*. Administer orally; repeat in 48 hours if necessary; milk taken from animals within 96 hours (8 milkings) of latest treatment must not be used for food; do not administer within 18 days of slaughter; discontinue use if hematuria, crystalluria or severe depression are noticed; if signs persist after 2 or 3 days consult a veterinarian.

[47 FR 30243, July 13, 1982, as amended at 62 FR 63270, Nov. 28, 1997]

§ 520.2184 Sodium sulfachloropyrazine monohydrate.

(a) *Chemical name*. 2-Sulfamido-6-chloroxyrazine, sodium.

(b) *Sponsor*. See Nos. 053501 in § 510.600(c) of this chapter.