infectious enteritis) caused by organisms susceptible to erythromycin.

(iii) Limitations. Administer for 7 days; do not use in turkeys producing eggs for human consumption; to assure effectiveness, treated birds must consume enough medicated water to provide a therapeutic dosage; solutions older than 3 days should not be used; withdraw 1 day before slaughter.


§ 520.863 Ethylisobutrazine hydrochloride tablets.

(a) Specifications. Each tablet contains either 10 milligrams or 50 milligrams of ethylisobutrazine hydrochloride.
(b) Sponsor. See No. 000061 in § 510.600(c) of this chapter.
(c) Conditions of use. (1) It is administered orally to dogs as a tranquillizer.1
(2) It is administered once daily at a dosage level of 2 to 5 milligrams of ethylisobutrazine hydrochloride per pound of body weight.1
(3) It is not to be used in conjunction with organophosphates and/or procaine hydrochloride because phenothiazine may potentiate the toxicity of organophosphates and the activity of procaine hydrochloride.1
(4) Federal law restricts this drug to use by or on the order of a licensed veterinarian.1


§ 520.870 Etodolac.

(a) Specifications. Each tablet contains 150, 300, or 500 milligrams (mg) of etodolac.
(b) Sponsor. See No. 000010 in § 510.600(c) of this chapter.
(c) [Reserved]
(d) Conditions of use—(1) Amount. Administer 10 to 15 mg per kilogram (4.5 to 6.8 mg per pound) of body weight per day orally.
(ii) Indications for use. For the management of pain and inflammation associated with osteoarthritis in dogs.
(iii) Limitations. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
(2) [Reserved]


§ 520.903 Febantel oral dosage forms.

§ 520.903a Febantel paste.

(a) Chemical name. Dimethyl [2-[(methoxyacetyl)amino]-4-(phenylthio)phenyl] carbonimidoyl]bis [carbamate].
(b) Specifications. The drug is a paste containing 45.5 percent febantel.
(c) Sponsor. See No. 000859 in § 510.600(c) of this chapter.
(d) Conditions of use—(1) Amount. Six milligrams per kilogram (2.73 milligrams per pound) of body weight in horses.
(2) Indications for use. For removal of large strongyles (Strongylus vulgaris, S. edentatus, S. equinus); ascarids (Parascaris equorum—sexually mature and immature); pinworms (Oxyuris equi—adult and 4th stage larva); and the various small strongyles in horses, foals, and ponies.
(3) Limitations. (i) The paste may be administered on the base of the tongue or well mixed into a portion of the normal grain ration.
(ii) [Reserved]
(iii) For animals maintained on premises where reinfection is likely to occur, retreatment may be necessary. For most effective results, retreat in 6 to 8 weeks.
(iv) Not for use in horses intended for food.
(v) Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism.


§ 520.903b Febantel suspension.

(a) Specifications. The suspension contains 9.3 percent (2.75 grams per ounce) febantel.
(b) Sponsor. See 000859 in § 510.600(c) of this chapter.