§ 520.930  Firocoxib paste.

(a) Specifications. Each milligram (mg) of paste contains 0.82 mg firocoxib.

(b) Sponsors. See No. 050604 in §510.600(c) of this chapter.

(c) Conditions of use in horses—(1) Amount. 0.1 mg per kilogram (0.045 mg per pound) body weight daily for up to 14 days.

(2) Indications for use. For the control of pain and inflammation associated with osteoarthritis.

(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.

§ 520.950  Flunixin oral dosage forms.

(a) Specifications. Each 10-gram packet contains flunixin meglumine equivalent to 250 milligrams of flunixin.

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

§ 520.955  Florfenicol.

(a) Specifications. Each milliliter (mL) contains 23 milligrams (mg) florfenicol.

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

(c) Related tolerances. See §556.283 of this chapter.

(d) Conditions of use in swine—(1) Amount. Administer in drinking water ad libitum at 400 mg per gallon (100 parts per million (ppm)) for 5 consecutive days.

(2) Indications for use. For the treatment of swine respiratory disease (SRD) associated with Actinobacillus pleuropneumoniae, Pasteurella multocida, Salmonella choleraesuis and Streptococcus suis.

(3) Limitations. Do not slaughter within 16 days of last treatment. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 520.970  Flunixin meglumine granules.

(a) Specifications. Each 10-gram packet contains flunixin meglumine equivalent to 250 milligrams of flunixin.

(b) Sponsor. No. 000061 in §510.600(c) of this chapter.
(c) Conditions of use—(1) Amount. 0.5 milligram of flunixin per pound of body weight (one packet per 500 pounds) per day.

(2) Indications for use. For alleviation of inflammation and pain associated with musculoskeletal disorders in the horse.

(3) Limitations. Administer daily dose for up to 5 days by sprinkling on small amount of feed. The effect of this drug on pregnancy has not been determined. Not for use in horses intended for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.


§ 520.970b Flunixin meglumine paste.

(a) Specifications. Each 30-gram syringe contains flunixin meglumine equivalent to 1,500 milligrams of flunixin.

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

(c) Conditions of use. Horses—(1) Amount. 0.5 milligram of flunixin per pound of body weight daily.

(2) Indications for use. For alleviation of inflammation and pain associated with musculoskeletal disorders.

(d) Limitations. For oral use only. Treatment should not exceed 5 consecutive days. The effect of this drug on pregnancy has not been determined. Not for use in horses intended for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.


§ 520.980 Fluoxetine.

(a) Specifications. Each chewable tablet contains 8, 16, 32, or 64 milligrams (mg) fluoxetine hydrochloride.

(b) Sponsor. See No. 00096 in § 510.600 of this chapter.

(c) Conditions of use in dogs—(1) Amount. 1 to 2 mg per kilogram body weight once daily.

(2) Indications for use. For the treatment of canine separation anxiety in conjunction with a behavior modification plan.

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

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§ 520.1010 Furosemide.

(a) Specifications. (1) Each tablet contains 12.5 or 50 milligrams (mg) furosemide.

(2) Each bolus contains 2 grams (g) furosemide.

(b) Sponsor. See sponsor numbers in § 510.600(c) of this chapter for use of dosage forms and strengths listed in paragraph (a) of this section for uses as in paragraph (d) of this section.

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

(d) Conditions of use. It is used as follows:

(1) Cattle—(i) Amount. 1 to 2 mg per pound (lb) body weight using powder, or one 2-g bolus per animal, per day.

(ii) Indications for use. For treatment of physiological parturient edema of the mammary gland and associated structures.

(iii) Limitations. Treatment not to exceed 48 hours post-parturition. Milk taken during treatment and for 48 hours after the last treatment must not be used for food. Cattle must not be slaughtered for food within 48 hours following last treatment.