days, or for 2 to 3 days beyond the cessation of clinical signs, up to a maximum of 28 days.

(2) Indications for use. For the treatment of skin infections (wounds and abscesses) caused by susceptible strains of *Staphylococcus intermedius*, *S. aureus*, *Streptococcus canis* (group G, -hemolytic), *Escherichia coli*, *Pasteurella multocida*, and *Proteus mirabilis*.

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


§ 520.376 Cephalexin.

(a) Specifications. Each chewable tablet contains 75, 150, 300, or 600 milligrams (mg) cephalexin.

(b) Sponsor. See No. 051311 in §510.600(c) of this chapter.

(c) Conditions of use—(1) Dogs—(i) Amount. Administer 22 mg per kilogram of body weight twice daily for 28 days.

(ii) Indications for use. For the treatment of secondary superficial bacterial pyoderma in dogs caused by susceptible strains of *Staphylococcus pseudintermedius*.

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

(2) [Reserved]

(77 FR 47512, Aug. 9, 2012)

§ 520.390 Chloramphenicol oral dosage forms.

§ 520.390a Chloramphenicol tablets.

(a) Specifications. Each tablet contains 50, 100, 250, or 500 milligrams (mg); 1 or 2.5 grams (g) of chloramphenicol.

(b) Sponsors. See Nos. 000069 and 050057 in §510.600(c) of this chapter for use as in paragraph (d) of this section.

(c) Special considerations. Federal law prohibits the extralabel use of this product in food-producing animals.

(d) Conditions of use in dogs—(1) Amount. 25 mg per pound of body weight every 6 hours.

(2) Indications for use. For treatment of bacterial pulmonary infections, bacterial infections of the urinary tract, bacterial enteritis, and bacterial infections associated with canine distemper caused by susceptible organisms.

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


§ 520.390b Chloramphenicol capsules.

(a) Specifications. Each capsule contains 50, 100, 250, or 500 milligrams (mg) chloramphenicol.

(b) Sponsors. See Nos. 000069 and 050057 in §510.600(c) of this chapter for use as in paragraph (d) of this section.

(c) Special considerations. Federal law prohibits the extralabel use of this product in food-producing animals.

(d) Conditions of use in dogs—(1) Amount. 25 mg per pound of body weight every 6 hours.

(2) Indications for use. For treatment of bacterial pulmonary infections, bacterial infections of the urinary tract, bacterial enteritis, and bacterial infections associated with canine distemper caused by susceptible organisms.

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

§ 520.390c Chloramphenicol palmitate oral suspension.

(a) Specifications. Each milliliter contains chloramphenicol palmitate equivalent to 30 milligrams of chloramphenicol.

(b) Sponsor. See No. 000856 in §510.600(c) of this chapter.
§ 520.420 Chlorothiazide tablets and boluses.

(a)(1) Specifications. Each tablet contains 0.25 gram of chlorothiazide.

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

(3) Conditions of use—(i) Amount. Usual dosage is 5 to 10 milligrams per pound of body weight two or three times daily.

(ii) Indications for use. For use in dogs for treatment of congestive heart failure and renal edema.

(iii) Limitations. (a) Dosage must be adjusted to meet the changing needs of the individual animal. In mild and responsive cases, it is suggested that a dose of 5 milligrams per pound of body weight be administered two or three times daily. In moderately edematous and moderately responsive animals, a dose of 7.5 to 10 milligrams per pound of body weight may be administered three times daily. Severe conditions may require higher doses. Certain animals may respond adequately to intermittent therapy; in these cases, the drug may be administered either every other day or for 3 to 5 days each week.

(b) Animals should be regularly and carefully observed for early signs of fluid and electrolyte imbalance. Take appropriate countermeasures if this should occur. In some dogs, hypochloremic alkalosis may occur (that is, excretion of chloride in relation to sodium is excessive; the plasma bicarbonate level increases and alkalosis results). Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(b)(1) Specifications. Each bolus contains 2 grams of chlorothiazide.

(2) Sponsor. See No. 000006 in §510.600(c) of this chapter.

(3) Conditions of use—(i) Amount. 2 grams once or twice daily for 3 or 4 days.

(ii) Indications for use. For use in cattle as an aid in reduction of postparturient udder edema.

(iii) Limitations. Animals should be regularly and carefully observed for early signs of fluid and electrolyte imbalance. Take appropriate countermeasures if this should occur. Milk taken from dairy animals during treatment and for 72 hours (six milkings) after latest treatment must not be used for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

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