Food and Drug Administration, HHS

Respiratory infections (tonsillitis, tracheobronchitis) due to *Staphylococcus aureus*, *Streptococcus spp.*, *Escherichia coli*, and *Proteus mirabilis*; genitourinary infections (cystitis) due to *S. aureus*, *Streptococcus spp.*, *E. coli*, and *P. mirabilis*; gastrointestinal infections (bacterial gastroenteritis) due to *S. aureus*, *Streptococcus spp.*, *E. coli*, and *P. mirabilis*; bacterial dermatitis due to *S. aureus*, *Streptococcus spp.*, and *P. mirabilis*; soft tissue infections (abscesses, lacerations, and wounds), due to *S. aureus*, *Streptococcus spp.*, *E. coli*, and *P. mirabilis*.

(ii) Cats. Treatment of infections caused by susceptible strains of organisms as follows: Upper respiratory infections due to *S. aureus*, *Staphylococcus spp.*, *Streptococcus spp.*, *Hemophilus spp.*, *E. coli*, *Pasteurella spp.*, and *P. mirabilis*; genitourinary infections (cystitis) due to *S. aureus*, *Streptococcus spp.*, *E. coli*, *P. mirabilis*, and *Corynebacterium spp.*; gastrointestinal infections due to *E. coli*, *Proteus spp.*, *Staphylococcus spp.*, and *Streptococcus spp.*; skin and soft tissue infections (abscesses, lacerations, and wounds) due to *S. aureus*, *Staphylococcus spp.*, *Streptococcus spp.*, *E. coli*, and *Pasteurella multocida*.

(3) Limitations. For use in dogs and cats only. Administer once daily for up to 5 days by intramuscular or subcutaneous injection. Continue treatment for 48 to 72 hours after the animal has become afebrile or asymptomatic. Do not continue treatment beyond 5 days. Treated animals must not be slaughtered for food during treatment and for 25 days after the last treatment. With all antibiotics, appropriate in vitro culturing and susceptibility testing of samples taken before treatment should be conducted. Milk from treated cows must not be used for human consumption during treatment or for 96 hours (8 milkings) after last treatment. Maximum volume per injection should not exceed 30 milliliters. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

See §556.40 of this chapter.

[57 FR 37330, Aug. 18, 1992; 60 FR 55659, Nov. 2, 1995]

§ 522.90a Ampicillin trihydrate sterile injectible dosage forms.

(a) Specifications. Each milliliter contains ampicillin trihydrate equivalent to 200 milligrams of ampicillin.

(1) Sponsor. See No. 053501 in §510.900(c) of this chapter.

(2) Related tolerances. See §556.40 of this chapter.

(b) Indications for use. Treatment of bacterial enteritis caused by *Escherichia coli* and bacterial pneumonia caused by *Pasteurella spp*. susceptible to ampicillin.

(c) Limitations. Not for use in other animals raised for food production. Treated animals must not be slaughtered for food use during treatment or for 9 days after the last treatment. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
§ 522.90b  Ampicillin trihydrate for suspension.

(a) Specifications. When reconstituted, each milliliter contains ampicillin trihydrate equivalent to 50, 100, or 250 milligrams of ampicillin.

(b) Sponsor. See No. 000010 in §510.600(c) of this chapter for use of 50, 100, and 250 milligrams per milliliter ampicillin suspension.

(2) Related tolerances. See §556.40 of this chapter.

§ 522.90b  Ampicillin trihydrate for suspension.

(a) Specifications. When reconstituted, each milliliter contains ampicillin trihydrate equivalent to 50, 100, or 250 milligrams of ampicillin.

(b) Sponsor. See No. 000010 in §510.600(c) of this chapter for use of 50, 100, and 250 milligrams per milliliter ampicillin suspension.

(2) Related tolerances. See §556.40 of this chapter.