§ 520.62 Aminopentamide hydrogen sulphate tablets.

(a) Chemical name. 4-(Dimethylamino)-2,2-diphenylvaleramide hydrogen sulfate.

(b) Specifications. Each tablet contains 0.2 milligram of the drug.

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

(d) Conditions of use. (1) It is intended for use in dogs and cats only for the treatment of vomiting and/or diarrhea, nausea, acute abdominal visceral spasm, pylorospasm, or hypertrophic gastritis.

   Note: Not for use in animals with glaucoma because of the occurrence of mydriasis.

   (2) Dosage is administered by oral tablet every 8 to 12 hours, as follows:

<table>
<thead>
<tr>
<th>Weight of animal in pounds</th>
<th>Dosage in milligrams</th>
</tr>
</thead>
<tbody>
<tr>
<td>Up to 10</td>
<td>0.1</td>
</tr>
<tr>
<td>11 to 20</td>
<td>0.2</td>
</tr>
<tr>
<td>21 to 50</td>
<td>0.3</td>
</tr>
<tr>
<td>51 to 100</td>
<td>0.4</td>
</tr>
<tr>
<td>Over 100</td>
<td>0.5</td>
</tr>
</tbody>
</table>

   Dosage may be gradually increased up to a maximum of five times the suggested dosage. Oral administration of tablets may be preceded by subcutaneous or intramuscular use of the injectable form of the drug.

   (3) For use only by or on the order of a licensed veterinarian.

[40 FR 13838, Mar. 27, 1975, as amended at 53 FR 27651, July 25, 1988]

§ 520.82 Aminopropazine fumarate oral dosage forms.

§ 520.82a Aminopropazine fumarate tablets.

(a) Specifications. The drug is in tablet form. Each tablet contains both aminopropazine fumarate equivalent to 25 milligrams of aminopropazine base and neomycin sulfate equivalent to 50 milligrams of neomycin base.

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

(c) Conditions of use. (1) The drug is used in dogs to control bacterial diarrhea caused by organisms susceptible to neomycin and to reduce smooth muscle contractions.

   (2) It is administered at a dosage level of one to two tablets per 10 pounds of body weight twice daily for 3 days.

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


§ 520.88 Amoxicillin oral dosage forms.

§ 520.88a Amoxicillin trihydrate film-coated tablets.

(a) Specifications. Each tablet contains amoxicillin trihydrate equivalent to 50, 100, 150, 200, or 400 milligrams of amoxicillin.

(b) Sponsor. See No. 000069 in §510.600(c) of this chapter.

(c) Conditions of use. (1) The drug is used in dogs and cats for reducing excessive smooth muscle contractions, such as occur in urethral spasms associated with urolithiasis.

   (2) It is administered at a dosage level of 1 to 2 milligrams per pound of body weight. The dosage can be repeated every 12 hours, as indicated.

   (3) Not for use in animals intended for food purposes.

   (4) For use only by or on the order of a licensed veterinarian.

§ 520.88b

| 21 CFR Ch. 1 (4–1–10 Edition) |

Amoxicillin trihydrate for oral suspension.

(a) Specifications. When reconstituted, each milliliter contains amoxicillin trihydrate equivalent to 50 milligrams of amoxicillin.

(b) Sponsor. See No. 000069 in §510.600(c) of this chapter.

(1) Conditions of use—(i) Dogs—(A) Amount. 5 milligrams per pound of body weight twice daily.

(B) Indications for use. Treatment of infections caused by susceptible strains of organisms as follows: respiratory tract (tonsillitis, tracheobronchitis) caused by Staphylococcus aureus, Streptococcus spp., Escherichia coli, and Proteus mirabilis; and genitourinary tract (cystitis) caused by S. aureus, Streptococcus spp., E. coli, and P. mirabilis; gastrointestinal tract (bacterial gastroenteritis) caused by S. aureus, Streptococcus spp., E. coli, and P. mirabilis; bacterial dermatitis caused by S. aureus, Streptococcus spp., and P. mirabilis; and soft tissue (abscesses, lacerations, and wounds) caused by S. aureus, Streptococcus spp., E. coli, and P. mirabilis.

(ii) Indications for use. Treatment of infections caused by susceptible strains of organisms as follows: upper respiratory tract due to Staphylococcus spp., Streptococcus spp., Escherichia coli, Proteus mirabilis; gastrointestinal tract due to E. coli; skin and soft tissue (abscesses, lacerations, and wounds) due to S. aureus, Streptococcus spp., E. coli, and Pasteurella multocida.

(iii) Limitations. Administer for 5 to 7 days or 48 hours after all symptoms have subsided. If no improvement is seen in 5 days, review diagnosis and change therapy. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(B) Indications for use. Treatment of infections caused by susceptible strains of organisms as follows: upper respiratory tract due to Staphylococcus spp., Streptococcus spp., Hemophilus spp., E. coli, Pasteurella spp., and P. mirabilis; gastrointestinal tract (cystitis) due to S. aureus, Streptococcus spp., E. coli, P. mirabilis, and Corynebacterium spp.; gastrointestinal tract due to E. coli, Proteus spp., Staphylococcus spp., and Streptococcus spp.; skin and soft tissue (abscesses, lacerations, and wounds) due to Staphylococcus spp., Streptococcus spp., E. coli, and Pasteurella multocida.

(C) Limitations. Use for 5 to 7 days or 48 hours after all symptoms have subsided. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(ii) Cats—(A) Amount. 50 milligrams (5 to 10 milligrams per pound of body weight) once a day.

(B) Indications for use. Treatment of infections caused by susceptible organisms as follows: upper respiratory tract due to S. aureus, Streptococcus spp., and E. coli; genitourinary tract (cystitis) due to S. aureus, Streptococcus spp., E. coli, and P. mirabilis; gastrointestinal tract due to E. coli; and skin and soft tissue (abscesses, lacerations, and wounds) due to S. aureus, Streptococcus spp., E. coli, and Pasteurella multocida.

(C) Limitations. Use for 5 to 7 days or 48 hours after all symptoms have subsided. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(ii) Indications for use. Treatment of infections caused by susceptible strains of organisms as follows: respiratory tract due to S. aureus, Streptococcus spp., and P. mirabilis; bacterial dermatitis caused by S. aureus, Streptococcus spp., and P. mirabilis; and soft tissues (abscesses, lacerations, and wounds) caused by S. aureus, Streptococcus spp., E. coli, and P. mirabilis.

(C) Limitations. Use for 5 to 7 days or 48 hours after all symptoms have subsided. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(ii) Limitations. Use for 5 to 7 days or 48 hours after all symptoms have subsided. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(ii) Limitations. Use for 5 to 7 days or 48 hours after all symptoms have subsided. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(ii) Limitations. Use for 5 to 7 days or 48 hours after all symptoms have subsided. Federal law restricts this drug to use by or on the order of a licensed veterinarian.