(2) Dogs—(i) Amount. 1 to 2 mg/lb body weight, once or twice daily.

(ii) Indications for use—(A) For treatment of edema (pulmonary congestion, ascites) associated with cardiac insufficiency and acute noninflammatory tissue edema.

(B) For treatment of edema (pulmonary congestion, ascites) associated with cardiac insufficiency.

(3) Cats—(i) Amount. 1 to 2 mg/lb body weight, once or twice daily.

(ii) Indications for use. For treatment of edema (pulmonary congestion, ascites) associated with cardiac insufficiency and acute noninflammatory tissue edema.

§ 520.1044 Gentamicin sulfate oral dosage forms.

§ 520.1044a Gentamicin sulfate oral solution.

(a) Specifications. Each milliliter of aqueous solution contains gentamicin sulfate equivalent to 50 milligrams of gentamicin.

(b) Sponsor. See Nos. 000061 and 054925 in §510.600(c) of this chapter.

(c) Related tolerances. See §556.300 of this chapter.

(d) Conditions of use—(1) Amount. Colibacillosis: 1 milliliter per 2 gallons of drinking water for 3 consecutive days, to provide 0.5 milligram/pound/day; swine dysentery: 1 milliliter per 1 gallon of drinking water for 3 consecutive days, to provide 1.0 milligram/pound/day.

(2) Indications for use. In weanling swine for control and treatment of colibacillosis caused by strains of E. coli sensitive to gentamicin, and in swine for control and treatment of swine dysentery associated with Treponema hyodysenteriae.

(3) Limitations. For use in swine drinking water only. Do not store or offer medicated drinking water in rusty containers since the drug is quickly destroyed in such containers. Medicated drinking water should be prepared daily and be the sole source of drinking water for 3 consecutive days. Treatment may be repeated if dysentery recurs. Do not slaughter treated swine for food for at least 3 days following treatment.

§ 520.1044b Gentamicin sulfate pig pump oral solution.

(a) Specifications. Each milliliter of pig pump oral solution contains gentamicin sulfate equivalent to 4.35 milligrams of gentamicin.

(b) Sponsor. See Nos. 000061 and 059130 in §510.600(c) of this chapter.

(c) Related tolerances. See §556.300 of this chapter.

(d) Conditions of use—(1) Amount. Administer 1.15 milliliters of pig pump oral solution (5 milligrams of gentamicin) orally per pig one time.

(2) Indications for use. In neonatal swine 1 to 3 days of age for control and treatment of colibacillosis caused by strains of E. coli sensitive to gentamicin.

(3) Limitations. For use in neonatal swine only. Do not slaughter treated swine for food for at least 14 days following treatment.

§ 520.1044c Gentamicin sulfate soluble powder.

(a) Specifications. Each gram of gentamicin sulfate soluble powder contains gentamicin sulfate equivalent to 16.7, 66.7, or 333.3 milligrams of gentamicin.

(b) Sponsor. See Nos. 000061 and 057561 in §510.600(c) of this chapter.

(c) Related tolerances. See §556.300 of this chapter.

(d) Conditions of use—(1) Amount. Colibacillosis: gentamicin sulfate equivalent to 25 milligrams of gentamicin per gallon of drinking water for 3 consecutive days, to provide 0.5 milligram per pound per day; swine dysentery: gentamicin sulfate equivalent to 50 milligrams of gentamicin per gallon of drinking water for 3 consecutive days, to provide 1 milligram per pound of body weight per day.

(2) Indications for use. In weanling swine for control and treatment of
colibacillosis caused by strains of *E. coli* sensitive to gentamicin, and in swine for control and treatment of swine dysentery associated with *Treponema hyodysenteriae.*

(3) **Limitations.** For use in swine drinking water only. Do not store or offer medicated drinking water in rusty containers since the drug is quickly destroyed in such containers. Medicated drinking water should be prepared daily and be the sole source of drinking water for 3 consecutive days. Treatment may be repeated if dysentery recurs. Do not slaughter treated swine for food for at least 10 days following treatment.


§ 520.1100 **Griseofulvin.**

(a) **Specifications.**—(1) The powder complies with U.S.P. for griseofulvin, microsize.

(2) Each bolus contains 2.5 grams griseofulvin.

(3) Each tablet contains 125 or 500 milligrams griseofulvin.

(b) **Sponsors.** See sponsors in §510.600(c) of this chapter.

(1) No. 000061 for use of products described in paragraph (a) for use as in paragraph (d) of this section.

(2) No. 059130 for use of the powder described in paragraph (a)(1) for use as in paragraphs (d)(1)(i)(A) and (d)(1)(ii) 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.**—(1) **Horses.**—(i) **Amount and indications for use.**—(A) For equine ringworm infection caused by *Trichophyton equinum* or *Microsporum gypseum*, administer soluble powder described in paragraph (a)(1) of this section daily as a drench or as a top dressing on feed for not less than 10 days as follows: adults, 2.5 grams; yearlings, 1.25 to 2.5 grams; and foals, 1.25 grams.

(B) For treating ringworm infection caused by *T. equinum*, administer boluses described in paragraph (a)(2) of this section daily for not less than 10 days as follows: adults, 1 bolus; yearlings, one-half to 1 bolus; and foals, one-half bolus.

(ii) **Limitations.** Not for use in horses intended for food.

(2) **Dogs and cats:** (i) **Amount.** 125- and 500-milligram tablets administered orally as follows:

<table>
<thead>
<tr>
<th>Body weight (pounds)</th>
<th>Dosage (milligrams)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Up to 6</td>
<td></td>
</tr>
<tr>
<td>6 to 18</td>
<td>125</td>
</tr>
<tr>
<td>18 to 36</td>
<td>250</td>
</tr>
<tr>
<td>36 to 48</td>
<td>375</td>
</tr>
<tr>
<td>48 to 75</td>
<td>500</td>
</tr>
</tbody>
</table>

(B) **Weekly (single) dose:** If experience indicates that treatment is more effective for the drug given in large doses, administer at intervals of 7 to 10 days, a dose equal to 10 milligrams/pound of body weight × body weight × number of days between treatments. Dosage should be adjusted according to response. Administer additional dose after the animal is free of infection.

(ii) **Indications for use.** For treatment of fungal infections of the skin, hair, and claws caused by *Trichophyton mentagrophytes*, *T. rubrum*, *T. schoenleini*, *T. sulphurem*, *T. verrucosum*, *T. interdigitale*, *Epidermophyton floccosum*, *Microsporum gypseum*, *M. canis*, *M. audouini*.


§ 520.1120 **Haloxon oral dosage forms.**

§ 520.1120a **Haloxon drench.**

(a) **Chemical name.** 3-Chloro-7-hydroxy-4-methylcoumarin bis (2-chloroethyl) phosphate.

(b) **Specifications.** Haloxon assay of not less than 96 percent by infrared spectrum at 8.62 microns.

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

(d) **Special considerations.** Do not use any drug, insecticide, pesticide, or other chemical having cholinesterase-inhibiting activity either simultaneously or within a few days before or after treatment with haloxon.

(e) **Related tolerances.** See §556.310 of this chapter.

(f) **Conditions of use.** It is used as a drench as follows: