

§ 520.23

- 520.2345d Tetracycline hydrochloride soluble powder.
- 520.2345e Tetracycline oral liquid.
- 520.2345f Tetracycline phosphate complex and sodium novobiocin capsules.
- 520.2345g Tetracycline hydrochloride and sodium novobiocin tablets.
- 520.2345h Tetracycline hydrochloride, sodium novobiocin, and prednisolone tablets.
- 520.2362 Thienium cloylate tablets.
- 520.2380 Thiabendazole oral dosage forms.
- 520.2380a Thiabendazole top dressing and mineral protein feed block.
- 520.2380b Thiabendazole drench or oral paste.
- 520.2380c Thiabendazole bolus.
- 520.2380d Thiabendazole, piperazine citrate suspension.
- 520.2380e Thiabendazole with trichlorfon.
- 520.2380f Thiabendazole, piperazine phosphate powder.
- 520.2455 Tiamulin soluble powder.
- 520.2456 Tiamulin liquid concentrate.
- 520.2473 Tioxidazole oral dosage forms.
- 520.2473a Tioxidazole granules.
- 520.2473b Tioxidazole paste.
- 520.2481 Triamcinolone acetone tablets.
- 520.2482 Triamcinolone acetone oral powder.
- 520.2520 Trichlorfon oral dosage forms.
- 520.2520b Trichlorfon and atropine.
- 520.2520e Trichlorfon boluses.
- 520.2520f Trichlorfon granules.
- 520.2520g Trichlorfon, phenothiazine, and piperazine dihydrochloride powder.
- 520.2582 Triflupromazine hydrochloride tablets.
- 520.2604 Trimeprazine tartrate and prednisolone tablets.
- 520.2605 Trimeprazine tartrate and prednisolone capsules.
- 520.2610 Trimethoprim and sulfadiazine tablets.
- 520.2611 Trimethoprim and sulfadiazine oral paste.
- 520.2612 Trimethoprim and sulfadiazine oral suspension.
- 520.2613 Trimethoprim and sulfadiazine powder.
- 520.2640 Tylosin.

AUTHORITY: 21 U.S.C. 360b.

SOURCE: 40 FR 13838, Mar. 27, 1975, unless otherwise noted.

§ 520.23 Acepromazine maleate tablets.

(a) *Sponsors.* See drug labeler codes in § 510.600(c) of this chapter for identification of sponsors as follows:

(1) For No. 000856, use of 5-, 10-, or 25-milligram tablets as in paragraph (b) of this section.

(2) For No. 000010, use of 10- or 25-milligram tablets as in paragraph (c) of this section.

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(b) *Conditions of use.* It is used in dogs and cats as follows:

(1) *Indications for use.* It is used in dogs and cats as a tranquilizer.

(2) *Amount.* Dogs: 0.25 to 1.0 milligram per pound of body weight; Cats: 0.5 to 1.0 milligram per pound of body weight.

(3) *Limitations.* The drug is administered orally. Dosage may be repeated as required. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(c) *Conditions of use.* It is used in dogs as follows:

(1) *Indications for use.* It is used in dogs as an aid in tranquilization and as a preanesthetic agent.

(2) *Amount.* Dogs: 0.25 to 1.0 milligram per pound of body weight.

(3) *Limitations.* The drug is administered orally. Dosage may be repeated as required. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[46 FR 44443, Sept. 4, 1981, as amended at 49 FR 49091, Dec. 18, 1984; 52 FR 666, Jan. 8, 1987; 53 FR 40727, Oct. 18, 1988; 56 FR 37473, Aug. 7, 1991; 62 FR 35075, June 30, 1997]

§ 520.44 Acetazolamide sodium soluble powder.

(a) *Specifications.* The drug is in a powder form containing acetazolamide sodium, USP equivalent to 25 percent acetazolamide activity.

(b) *Sponsor.* See No. 053501 in § 510.600(c) of this chapter.

(c) *Conditions of use.* (1) It is used in dogs as an aid in the treatment of mild congestive heart failure and for rapid reduction of intraocular pressure.¹

(2) It is administered orally at a dosage level of 5 to 15 milligrams per pound of body weight daily.¹

(3) For use only by or on the order of a licensed veterinarian.¹

[40 FR 13838, Mar. 27, 1975, as amended at 67 FR 78355, Dec. 24, 2002]

¹These conditions are NAS/NRC reviewed and deemed effective. Applications for these uses need not include effectiveness data as specified by § 514.111 of this chapter, but may require bioequivalency and safety information.