

Food and Drug Administration, HHS

§ 520.45a

- 520.2345a Tetracycline hydrochloride capsules.
- 520.2345b Tetracycline tablets.
- 520.2345c Tetracycline boluses.
- 520.2345d Tetracycline 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 Thienem cloylate tablets.
- 520.2380 Thiabendazole oral dosage forms.
- 520.2380a Thiabendazole top dressing and mineral protein 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.
- 520.2473 Tioxidazole oral dosage forms.
- 520.2473a Tioxidazole granules.
- 520.2473b Tioxidazole paste.
- 520.2475 Toceranib.
- 520.2481 Triamcinolone acetonide tablets.
- 520.2482 Triamcinolone acetonide oral powder.
- 520.2483 Triamcinolone.
- 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.
- 520.2598 Trilostane.
- 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 paste.
- 520.2612 Trimethoprim and sulfadiazine oral suspension.
- 520.2613 Trimethoprim and sulfadiazine powder.
- 520.2640 Tylosin.
- 520.2645 Tylvalosin.

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

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

§ 520.23 Acepromazine.

(a) *Specifications.* Each tablet contains 5, 10, or 25 milligrams (mg) acepromazine maleate.

(b) *Sponsors.* See No. 000010 in § 510.600(c) of this chapter.

(c) *Conditions of use*—(1) *Dogs*—(i) *Amount.* 0.25 to 1.0 mg per pound (lb) body weight orally.

(ii) *Indications for use.* As an aid in tranquilization and as a preanesthetic agent.

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

(2) *Cats*—(i) *Amount.* 0.5 to 1.0 mg/lb body weight orally.

(ii) *Indications for use.* As a tranquilizer.

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

[75 FR 10165, Mar. 5, 2010]

§ 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]

§ 520.45 Albendazole oral dosage forms.

§ 520.45a Albendazole suspension.

(a) *Specifications.* Each milliliter of suspension contains 45.5 milligrams (mg) (4.55 percent) or 113.6 mg (11.36 percent) albendazole.

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

(c) *Related tolerances.* See § 556.34 of this chapter.

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.