§ 524.2350

rash or irritation develops, discontinue use and consult a veterinarian.


§ 524.2350 Tolnaftate cream.

(a) Specifications. The drug contains 1 percent tolnaftate (2-naphthyl-N-methyl-N-(3-tolyl) thionocarbamate) in an anhydrous cream base.

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

(c) Conditions of use. (1) The drug is indicated for treatment of ringworm lesions due to Microsporum canis and Microsporum gypseum in dogs and cats.

(2) A small amount of the cream is applied to the affected areas once or twice a day for 2 to 4 weeks. The areas to be treated are first cleared of exudate and the hair clipped if the areas are not already denuded. The cream is massaged into each lesion and immediate surrounding area until the cream is no longer visible.

(3) If no response is seen after 2 weeks of treatment with the drug the diagnosis should be reviewed.

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


§ 524.2482 Triamcinolone spray.

(a) Specifications. Each milliliter of solution contains 0.15 milligrams triamcinolone acetonide.

(b) Sponsor. See No. 067292 in § 510.600(c) of this chapter.

(c) Conditions of use in dogs—(1) Amount. Rub into affected areas two to four times daily for 4 to 10 days.

(2) Indications for use. For topical treatment of allergic dermatitis and summer eczema.

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

[68 FR 4916, Jan. 31, 2003]

PART 526—INTRAMAMMARY DOSAGE FORM NEW ANIMAL DRUGS

§ 524.2483 Triamcinolone cream.

(a) Specifications. The vanishing cream contains 0.1 percent triamcinolone acetonide.

(b) Sponsor. See Nos. 000010, 015914, and 054925 in § 510.600(c) of this chapter.

(c) Conditions of use in dogs—(1) Amount. Rub into affected areas two to four times daily for 4 to 10 days.

(2) Indications for use. For topical treatment of allergic dermatitis and summer eczema.

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


§ 524.2620 Liquid crystalline trypsin, Peru balsam, castor oil.

(a)(1) Specifications. The drug is a liquid for direct application or an aerosol preparation formulated so that each gram delivered to the wound site contains 0.12 milligram of crystalline trypsin, 87.0 milligrams of Peru balsam, and 788.0 milligrams of castor oil.

(b)(1) Specifications. The drug is a liquid for direct application or an aerosol preparation formulated so that each gram delivered to the wound site contains 0.1 milligram of crystalline trypsin, 72.5 milligrams of Peru balsam, and 800 milligrams of castor oil.

(2) Sponsor. See No. 051079 in § 510.600(c) of this chapter.

(3) Conditions of use. The drug is used as an aid in the treatment of external wounds and assists healing by facilitating the removal of necrotic tissue, exudate and organic debris.