(c) Conditions of use—
(1) Amount—
(i) Preparation containing hydroxypropyl methylcellulose. Dogs: 1 to 2 drops per eye, every 6 hours.
(ii) Preparation without hydroxypropyl methylcellulose. Dogs and cats: 2 to 3 drops per eye, every 4 hours.
(2) Indications for use. Treatment of the inflammation, edema, and secondary bacterial infections associated with topical ophthalmological conditions of the eye such as corneal injuries, incipient pannus, superficial keratitis, conjunctivitis, acute nongranulomatous anterior uveitis, kerato-conjunctivitis, and blepharitis.
(3) Limitations. (i) In treating ophthalmological conditions associated with bacterial infections, the drug is contraindicated in those cases in which microorganisms are not susceptible to the antibiotics incorporated in the drug.
(ii) The drug is contraindicated in infectious tuberculous lesions of the eye, early acute stages of viral diseases of the cornea and conjunctiva, herpes simplex lesions of the eye, and fungal infections of the conjunctiva and eyelids.
(iii) The usual precautions and contraindications for corticosteroids and adrenocorticoids are applicable with this drug. Corticosteroids may inhibit essential inflammatory responses intrinsic to the fundamental healing mechanism. Adrenocorticoid compounds have been reported to cause an increase in intraocular pressure. Intraocular pressure should be checked frequently. Ocular reexaminations should be made at frequent intervals during long-term therapy.
(iv) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 524.981 Fluocinolone acetonide ophthalmic and topical dosage forms.

§ 524.981a Fluocinolone acetonide cream.

(a) Specifications. The drug contains 0.025 percent fluocinolone acetonide.
(b) Sponsor. See No. 099207 in § 510.600(c) of this chapter.

§ 524.981c Fluocinolone acetonide, neomycin sulfate cream.

(a) Specifications. The drug contains 0.025 percent fluocinolone acetonide and 0.5 percent neomycin sulfate (0.35 percent neomycin base).
(b) Sponsor. See No. 099207 in § 510.600(c) of this chapter.
(c) Conditions of use. (1) The drug is used in the treatment of superficial acute and chronic dermatoses in dogs. It is used in the treatment of allergic and acute moist dermatoses and burns.
(2) A small amount is applied to the affected area two or three times daily.
(3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.
[40 FR 13873, Mar. 27, 1975, as amended at 62 FR 40932, July 31, 1997]
in dogs. It is used in the treatment of wound infections in dogs and cats.
(2) A small amount is applied to the infected area two or three times daily.
(3) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 524.981d Fluocinolone acetonide, di- methyl sulfoxide solution.

(a) Specifications. Each milliliter of solution contains 0.01 percent fluocinolone acetonide and 20 percent dimethyl sulfoxide with propylene glycol and citric acid.
(b) Sponsor. See No. 000856 in §510.600(c) of this chapter.
(c) Conditions of use. (1) The drug is used in dogs for the relief of impaction commonly present in apparently normal anal sacs, for the reversal of inflammatory changes associated with abnormal anal sacs, and to counteract the offensive odor of anal sac secretions.
(2) It is administered by instillation of 1 to 2 milliliters into each anal sac following expression of anal sac contents. It may be necessary to repeat treatment at 60-day intervals to maintain an odor-free state. The total dosage used should not exceed 2 milliliters per anal sac per treatment.
(3) For use only by or on the order of a licensed veterinarian.

§ 524.1005 Furazolidone aerosol powder.

(a) Specifications. The product contains either 4 or 10 percent furazolidone in inert dispersing agent and propel- lant.
(b) Sponsors. (1) See No. 053501 in §510.600(c) of this chapter for use as in paragraphs (c)(1), (c)(2)(i), (c)(2)(ii), and (c)(3) of this section.
(2) See No. 017135 for use of the 4 percent product as in paragraph (c)(2)(iv) of this section.
(c) Conditions of use—(1) Amount. Hold container about 6 to 12 inches from the eye or affected area and apply only enough powder to impart a light yellow color.
(2) Indications of use—(i) Dogs. For treatment or prevention of bacterial infection of superficial wounds, abrasions, lacerations, and pyogenic dermatitis.
(ii) Horses. For treatment or prevention of bacterial infection of superficial wounds, abrasions, lacerations, and following firing (heat or electrocautery).
(iii) [Reserved]
(iv) Horses and ponies. For treatment or prevention of bacterial infection of superficial wounds, abrasions, and lacerations caused by Staphylococcus aureus, Streptococcus spp. and Proteus spp. sensitive to furazolidone.
(3) Limitations. For topical application in horses, ponies, and dogs: Clean affected area thoroughly, apply drug