(c) **Conditions of use in horses**—(1) **Dosage.** Five milligrams per kilogram of body weight intravenously followed by maintenance oral therapy of 10 milligrams per kilogram of body weight twice daily for up to 14 consecutive days.

(2) **Indications for use.** For the relief of inflammation and associated pain and lameness exhibited with arthritis, as well as myositis and other soft tissue diseases of the musculoskeletal system of the horse.

(3) **Limitations.** Not for use in horses intended for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 522.1484 Neomycin.

(a) **Specifications.** Each milliliter of solution contains 50 milligrams (mg) of neomycin sulfate (equivalent to 35 mg of neomycin base).

(b) **Sponsor.** See No. 054771 in § 510.600(c) of this chapter.

(c) **Conditions of use in dogs and cats**—(1) **Amount.** Administer 5 mg per pound of body weight daily by intramuscular or intravenous injection, divided into portions administered every 6 to 8 hours for 3 to 5 days.

(2) **Indications for use.** For the treatment of acute and chronic bacterial infections due to organisms susceptible to neomycin.

(3) **Limitations.** Not for parenteral use in food-producing animals because of prolonged residues in edible tissues. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 522.1503 Neostigmine.

(a) **Specifications.** Each milliliter of solution contains 2 milligrams (mg) neostigmine methylsulfate.

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

(c) **Conditions of use**—(1) **Amount.** Administer to cattle and horses at a dosage level of 1 mg per (/) 100 pounds (lbs) of body weight subcutaneously. Administer to sheep at a dosage level of 1 to 1½ mg/100 lbs body weight subcutaneously. Administer to swine at a dosage level of 2 to 3 mg/100 lbs body weight intramuscularly. These doses may be repeated as indicated.

(2) **Indications for use.** For treating rumen atony; initiating peristalsis which causes evacuation of the bowel; emptying the urinary bladder; and stimulating skeletal muscle contractions.

(3) **Limitations.** Not for use in animals producing milk, since this use will result in contamination of the milk. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 522.1610 Oleate sodium.

(a) **Specifications.** Each milliliter of solution contains 50 milligrams (mg) of sodium oleate.

(b) **Sponsor.** See No. 037990 in § 510.600(c) of this chapter.

(c) **Conditions of use in horses**—(1) **Amount.** Administer by parenteral injection depending on the area of response desired. An injection of 1 milliliter (mL) will produce a response of approximately 15 square centimeters. Do not inject more than 2 mL per injection site. Regardless of the number of injection sites, the total volume used should not exceed 10 mL.

(2) **Indications for use.** It is used in horses to stimulate infiltration of cellular blood components that subsequently differentiate into fibrous and/or fibrocartilagenous tissue.

(3) **Limitations.** Do not use in horses intended for human consumption. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

§ 522.1620 Orgotein for injection.

(a) **Specifications.** Orgotein for injection is packaged in a vial containing 5 milligrams of orgotein and 10 milligrams of sucrose as lyophilized sterile nonpyrogenic powder with directions for dissolving the contents of the vial in 2 milliliters of diluent which is sodium chloride injection, U.S.P.

(b) **Sponsor.** See No. 024991 in § 510.600(c) of this chapter.