mg estradiol benzoate, and 1 pellet containing 29 mg tylosin tartrate) per implant dose.

(ii) Indications for use. For increased rate of weight gain and improved feed efficiency.

(iii) Limitations. For animals weighing 400 lb or more; for subcutaneous ear implantation, one dose per animal. Safety and effectiveness have not been established in veal calves. A withdrawal period has not been established for this product in preruminating calves. Do not use in calves to be processed for veal.

(3) Steers fed in confinement for slaughter—(i) Amount. Reimplant 200 mg progesterone and 20 mg estradiol benzoate on approximately day 70 following an initial implant of 100 mg progesterone and 10 mg estradiol benzoate or 200 mg progesterone and 20 mg estradiol benzoate.

(ii) Indications for use. For additional improvement in rate of weight gain.

(iii) Limitations. For subcutaneous ear implantation. Safety and effectiveness have not been established in veal calves. A withdrawal period has not been established for this product in preruminating calves. Do not use in calves to be processed for veal.

§ 522.2005 Propofol.

(a) Specifications. Each milliliter of emulsion contains 10 milligrams (mg) propofol.

(b) Sponsors. See sponsor numbers in §510.600(c) of this chapter.

(1) No. 000044 for use as in paragraphs (c)(1), (c)(2)(i), and (c)(3) of this section.

(2) No. 000044 for use as in paragraphs (c)(1), (c)(2)(i), and (c)(3) of this section.

(3) No. 054771 for use as in paragraphs (c)(1), (c)(2)(ii), and (c)(3) of this section.

§ 522.2002 Propiopromazine.

(a) Specifications. Each milliliter of solution contains 5 or 10 milligrams (mg) propiopromazine hydrochloride.

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

(c) Conditions of use in dogs and cats—(1) Amounts and indications for use. Administer 0.05 to 0.5 mg per pound of body weight by intravenous or intramuscular injection for tranquilization. Administer 0.25 mg per pound of body weight by intravenous injection as a preanesthetic.

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

§ 522.1962 Promazine.

(a) Specifications. Each milliliter of solution contains 50 milligrams (mg) promazine hydrochloride.

(b) Sponsors. See sponsors in §510.600(c) of this chapter for use as in paragraph (c) of this section:

(1) No. 054771 for use as in paragraphs (c)(1)(i)(A), (c)(1)(ii)(A), (c)(1)(iii), and (c)(2) of this section.

(2) No. 061623 for use as in paragraphs (c)(1)(i)(B), (c)(1)(ii)(B), and (c)(1)(iii) of this section.

(c) Conditions of use—(1) Horses—(A) Amount—(A) 0.2 to 0.5 milligrams per pounds (mg/lb) body weight intramuscularly or intravenously every 4 to 6 hours.

(B) 0.2 to 0.5 mg/lb body weight intravenously as required.

(ii) Indications for use—(A) For use as a tranquilizer, preanesthetic, or for minor operative procedures in conjunction with local anesthesia; and as adjunctive therapy for tetanus.

(B) For use as a tranquilizer and preanesthetic.

(iii) 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.

(2) Dogs and cats—(i) Amount. 1 to 2 mg/lb body weight intramuscularly or intravenously every 4 to 6 hours.

(ii) Indications for use. For use as a tranquilizer, preanesthetic, for minor operative procedures in conjunction with local anesthesia, as adjunctive therapy for tetanus, and as an antiemetic prior to worming; or to prevent motion sickness in dogs.

(iii) Limitations. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
§ 522.2012

(c) Conditions of use in dogs and cats—

(1) Amount. Administer by intravenous injection according to label directions. The use of preanesthetic medication reduces propofol dose requirements.

(2) Indications for use—

(i) As a single injection to provide general anesthesia for short procedures; for induction and maintenance of general anesthesia using incremental doses to effect; for induction of general anesthesia where maintenance is provided by inhalant anesthetics.

(ii) For the induction and maintenance of anesthesia and for induction of anesthesia followed by maintenance with an inhalant anesthetic.

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

§ 522.2063

(a) Specifications. Each milliliter of solution contains 20 milligrams (mg) pyrilamine maleate.

(b) Sponsors. See sponsor numbers in §510.600(c) of this chapter.

(c) Conditions of use—

(i) Horses, 40 to 60 mg per 100 pounds (lbs) body weight; foals, 20 mg/100 lbs body weight. Administer by intramuscular, subcutaneous, or intravenous injection. Dosage may be repeated every 6 to 12 hours whenever necessary.

(ii) Horses, 40 to 60 mg/100 lbs body weight; foals, 20 mg/100 lbs body weight. Administer by slow intravenous injection. Dosage may be repeated every 6 to 12 hours if necessary.

(2) Indications for use. It is intended for treating horses in conditions in which antihistaminic therapy may be expected to lead to alleviation of some signs of disease.

(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.

§ 522.2076

(a) Specifications. Each milliliter of solution contains 10 milligrams (mg) romifidine hydrochloride.

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

(c) Conditions of use in horses—

(i) Amount. 40 to 120 micrograms per kilogram of body weight (mcg/kg BW) intravenously for sedation and analgesia; 100 mcg/kg BW intravenously as a preanesthetic.

(2) Indications for use. For use as a sedative and analgesic to facilitate handling, clinical examinations, clinical procedures, and minor surgical procedures in adult horses; and for use as a preanesthetic prior to the induction of general anesthesia in adult horses.

(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.

§ 522.2100

(a)(1) Specifications. Each milliliter of emulsion contains 5.48 milligrams (mg) sodium selenite (equivalent to 2.5 mg selenium) and 50 mg of vitamin E (68 I.U.) (as d-alpha tocopheryl acetate).

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

(3) Conditions of use in horses—

(i) Amount. Administer 1 milliliter (mL) per (/) 100 pounds (lbs) of body weight by intravenous injection or by deep intramuscular injection in divided