(iii) **Limitations.** For intraarticular injection in horses only. Treatment may be repeated after 1 or more weeks but not to exceed 2 injections per week for a total of 4 weeks. Not for use in horses intended for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

(d)(1) **Specifications.** Each milliliter of sterile aqueous solution contains 10 milligrams of hyaluronate sodium.

(2) **Sponsor.** See 000061 in §510.600(c) of this chapter.

(e)(1) **Specifications.** Each milliliter of solution contains:

(i) 10 milligrams (mg) hyaluronate sodium; or

(ii) 10 mg hyaluronate sodium with benzyl alcohol as a preservative.

(2) **Sponsor.** See 060865 in §510.600(c).

(3) **Conditions of use**—(i) **Amount.** 5 to 10 milliliters (125 to 250 milligrams) intravenously or intramuscularly once or twice a day. After onset of diuresis, treatment may be continued with an orally administered maintenance dose.

(2) **Indications for use.** For use in cattle as an aid in the treatment of postparturient udder edema.**

(3) **Limitations.** Animals should be regularly and carefully observed for early signs of fluid and electrolyte imbalance. Take appropriate countermeasures if this should occur. Milk taken from dairy animals during treatment and for 72 hours (6 milkings) after the latest treatment must not be used for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.**

§ 522.1150 **Hydrochlorothiazide injection.**

(a) **Specifications.** Each milliliter contains 25 milligrams of hydrochlorothiazide.

(b) **Sponsor.** See No. 050604 in §510.600(c) of this chapter.

(c) **Conditions of use**—(1) **Amount.** 20 mg of the product described in paragraph (e)(1)(i) of this section by intra-articular injection into the carpus or fetlock; or 40 mg of the product described in paragraph (e)(1)(i) or (e)(1)(ii) of this section by slow intravenous injection into the jugular vein. Treatment may be repeated at weekly intervals for a total of three treatments.

(ii) **Indications for use.** For treatment of carpal or fetlock joint dysfunction due to noninfectious synovitis associated with equine osteoarthritis.

(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.

§ 522.1150 **Hydrochlorothiazide injection.**

(a) **Specifications.** Each milliliter contains 25 milligrams of hydrochlorothiazide.

(b) **Sponsor.** See No. 050604 in §510.600(c) of this chapter.

(c) **Conditions of use**—(1) **Amount.** 5 to 10 milliliters (125 to 250 milligrams) intravenously or intramuscularly once or twice a day. After onset of diuresis, treatment may be continued with an orally administered maintenance dose.

(ii) **Indications for use.** For use in cattle as an aid in the treatment of postparturient udder edema.**

(3) **Limitations.** Animals should be regularly and carefully observed for early signs of fluid and electrolyte imbalance. Take appropriate countermeasures if this should occur. Milk taken from dairy animals during treatment and for 72 hours (6 milkings) after the latest treatment must not be used for food. Federal law restricts this drug to use by or on the order of a licensed veterinarian.**


§ 522.1150 **Hydrochlorothiazide injection.**

(a) **Specifications.** Each milliliter contains 25 milligrams of hydrochlorothiazide.

(b) **Sponsor.** See No. 050604 in §510.600(c).

(c) **Conditions of use**—(1) **Amount.** Small and medium-size joints (carpal, fetlock)—22 milligrams; larger joint (hock)—44 milligrams.

(ii) **Indications for use.** Treatment of joint dysfunction in horses due to noninfectious synovitis associated with equine osteoarthritis.

(iii) **Limitations.** For intra-articular injection in horses only. Treatment may be repeated at weekly intervals for a total of three treatments. 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.