SODIUM IODIDE- sodium iodide injection, solution
Nova-Tech, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

SODIUM IODIDE 20%

INDICATIONS

For use as an aid in the treatment of actinomycosis (lumpy jaw), actinobacillosis (wooden tongue) and necrotic stomatitis in cattle.

CONTRAINDICATIONS

The use of sodium iodide is contraindicated in pregnancy and hyperthyroidism.

CAUTION

Animals vary in their susceptibility of iodides. Administer with caution until the animal's tolerance is determined. Discontinue treatment if adverse reactions occur.

DOSAGE AND ADMINISTRATION:

Using aspetic procedures, administer slowly by intravenous injection. Inject carefully to avoid deposition outside of the vein. The usual dose is 30 mg per pound of body weight (15 mL/100 lb). May be repeated at weekly intervals, if necessary.

WARNING

Not for use in lactating dairy cows.

For Animal Use Only

Keep Out of Reach Of Children

CAUTION:

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

COMPOSITION

Each 100 mL of sterile aqueous solution contains:
Sodium Iodide.....20 grams
Water For Injection.....q.s.

Store between 15°C-30°C (59°F-86°F).

TAKE TIME OBSERVE LABEL DIRECTIONS

Manufactured by:

Nova-Tech, Inc.

Grand Island, NE 68801 USA

18-819

RMS 92-366

NDC# 65207-819-25

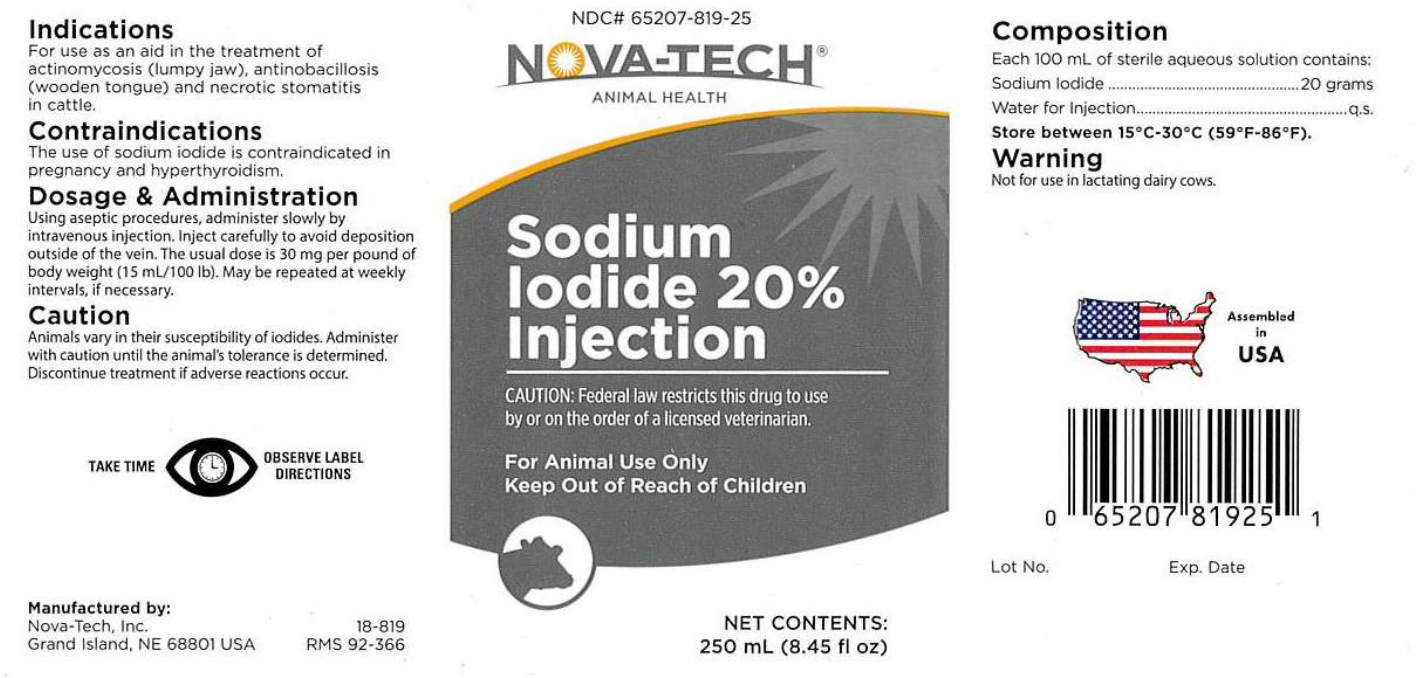
Net Contents:

250 mL (8.45 fl oz)

Assembled in USA

Lot No.

Exp. Date



SODIUM IODIDE

sodium iodide injection, solution

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:65207-819
Route of Administration	INTRAVENOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM IODIDE (UNII: F5WR8N145C) (IODIDE ION - UNII:09G4I6V86Q)	IODIDE ION	20 g in 100 mL

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:65207-819-25	250 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		08/22/2019	

Labeler - Nova-Tech, Inc. (196078976)

Registrant - Nova-Tech, Inc. (196078976)

Establishment

Name	Address	ID/FEI	Business Operations
Nova-Tech, Inc.		196078976	manufacture, api manufacture

Revised: 8/2019 Nova-Tech, Inc.