SODIUM IODIDE- sodium iodide injection, solution
VetTek

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% Injection

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.

WARNING:

Not for use in lactating dairy cows.

DOSAGE AND ADMINISTRATION:

Using aseptic 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.

CAUTION:

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

RMS-92-1017 Iss. 09-11
18-819-25

Manufactured For
VETTEK
Blue Springs, MO 64014

NET CONTENTS: 250 mL

Manufactured by
Nova-Tech, Inc.
Grand Island, NE 68801

Lot No. Exp. Date

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 degrees C and 30 degrees C (59 degrees F and 86 degrees F).

TAKE TIME OBSERVE LABEL DIRECTIONS

SODIUM IODIDE

sodium iodide injection, solution

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:60270-863
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:60270-863-13	250 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/17/2011	

Labeler - VetTek (056387798)

Registrant - VetTek (056387798)

Establishment

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

Revised: 10/2011 VetTek