HYPERTONIC SALINE- hypertonic saline injection, solution
Phoenix

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.

Hypertonic Saline Solution 7.2%

INDICATIONS:

For use in replacement therapy of sodium, chloride and water which may become depleted in many diseases.

CAUTION:

This product contains no preservatives. Use entire contents when first opened. Discard any unused solution.

DOSAGE AND ADMINISTRATION:

Warm to body temperature and administer slowly by intravenous or subcutaneous injection. The amount and rate of administration must be judged by the veterinarian in relation to the condition being treated and the clinical response of the animal, being careful to avoid overhydration.

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 Chloride ..... 7.2 g

Milliequivalents per liter

Cations
Sodium ..... 1232 mEq/L
Anions
Chloride ..... 1232 mEq/L

Total osmolarity is 2464 milliosmoles per liter.

Store between 15°C and 30°C (59°F and 86°F)

TAKE TIME OBSERVE LABEL DIRECTIONS

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

RMS 92-548

18-806-60

Rev. 06-10

NDC 57319-554-08

Sterile - Preservative Free

Net Contents: 1000 mL

Manufactured for:
Clipper Distributing Company, LLC
St. Joseph, MO 64507

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Clipper Distributing Company, LLC

Lot No.

Exp. Date

HYPERTONIC SALINE

hypertonic saline injection, solution

Product Information			
Product Type	PRESCRIPTION ANIMAL DRUG	**Item Code (Source)**	NDC:57319-554
Route of Administration	INTRAVENOUS, SUBCUTANEOUS		

Active Ingredient/Active Moiety		

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	7.2 g in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57319-554-08	1000 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/16/2018	

Labeler - Phoenix (150711039)

Registrant - Phoenix (150711039)

Establishment

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

Revised: 1/2018 Phoenix