HYPERTONIC SALINE SOLUTION 7.2% - hypertonic saline solution 7.2% injection, solution Vedco

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.

EQUI-PHAR EQUINE 7 HSS

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.

CAUTION:

Federal law (U.S.A.) restricts this drug to use by or on the order of a licensed veterinarian.

FOR VETERINARY USE ONLY

COMPOSITION:

Milliequivalents per literCations Sodium....... 1232 mEq/LAnions Chloride......... 1232 mEq/L

Total osmolarity is 2464 milliosmoles per liter.

Store between 15°C-30°C (59°F-86°F).

TAKE TIME OBSERVE LABEL DIRECTIONS

For Animal Use Only

KEEP OUT OF REACH OF CHILDREN

MADE IN U.S.A.

Iss. 03-01

18-806-60

Distributed By VEDCO, INC. St. Joseph, MO 64507

NDC 50989-638-17

STERILE HYPERTONIC

7.2% SALINE SOLUTION

NET CONTENTS: 1000 mL

Lot No.

Exp. Date

FOR VETERINARY USE ONLY

Milliequivalents per liter:

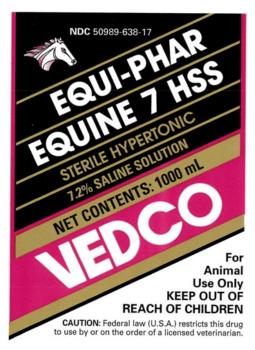
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Product Information						
Product Type	PRESCRIPTION ANIMAL DRUG	Item Code (Source)	NDC:50989-638			
Route of Administration	INTRAVENOUS, SUBCUTANEOUS					

Active Ingredient/Active Moiety								
	Ba	sis of Streng	gth Strengt	h				
Sodium Chloride (UNII: 451W47IQ8X) (Sodium Cation - UNII:LYR4M0NH37)			Sodi	um Chloride	7.2 g in 100 i	mL		
Packaging								
# Item Code	Package Description	Marketing Start Date		e Ma	Marketing End Date			
1 NDC:50989-638-17	1000 mL in 1 BOTTLE							
Marketing Information								
Marketing Category	Application Number or Monograph Citation Ma		Marketing Start Date		Marketing End D)at		
unapproved drug other			08/03/2017					

Labeler - Vedco (021634266)

Registrant - Vedco (021634266)

Establishment

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

Revised: 3/2018

Vedco