THIAMINE HYDROCHLORIDE- thiamine hydrochloride injection Covetrus North America

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.

Thiamine Hydrochloride 500mg/mL Injectable Sterile Solution

Recommended Use:

For use as a supplemental source of thiamine in dogs, cats, and horses.

Each mL Contains:

Thiamine Hydrochloride	500 mg
Disodium Edetate	0.01% w/v
Benzyle Alcohol (as preservative)	1.5% v/v
Water for Injection	q.s.
pH adjusted with Hydrochloric acid	or Sodium Hydroxide

Dosage:

For intravenous or intramuscular use as determined by the veterinarian.

Warning:

Anaphylactogenesis to parental thiamine has occurred. Administer slowly and with caution in doses over 0.10 mL (50mg)

Store at controlled temperatures between 15° and 30°C (59° - 86°F).

Reorder #002489

Questions?

(855) 724-346

Covetrus North America

400 Metro Place North

Dublin, OH 43017

covetrus.com

Made in the USA

AH-002489-02

V-0347-04

L6566-0220

ISS: 0220

REV: 0220

Principal Display Panel

THIAMINE HYDROCHLORIDE

500 mg/mL Injectable Sterile Solution

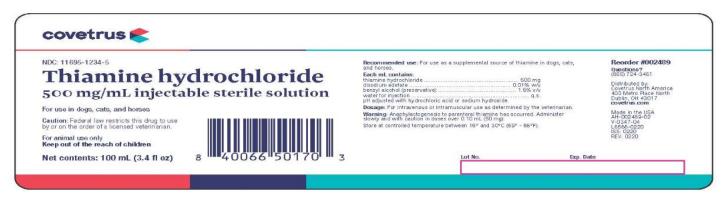
For use in dogs, cats and horses

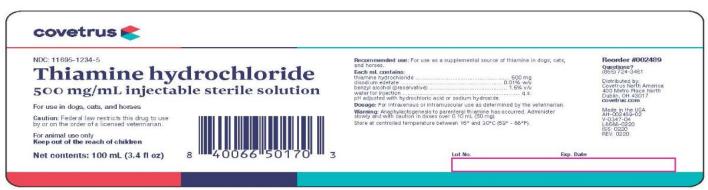
Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian

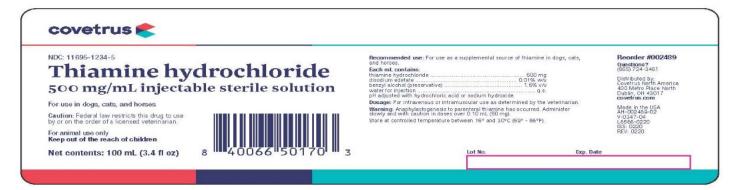
FOR ANIMAL USE ONLY

KEEP OUT OF THE REACH OF CHILDREN

100 mL (3.4 fl oz)







THIAMINE HYDROCHLORIDE

thiamine hydrochloride injection

Product Information

Product Type PRESCRIPTION ANIMAL DRUG Item Code (Source) NDC:11695-1234

Route of Administration INTRAVENOUS, INTRAMUSCULAR

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength

THIAMINE HYDROCHLORIDE (UNII: M572600E5P) (THIAMINE ION UNII:4ABT0J945J)

THIAMINE HYDROCHLORIDE

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Inactive Ingredients Ingredient Name Strength EDETATE DISODIUM (UNII: 7FLD91C86K) .1 mg in 1 mL BENZYL ALCOHOL (UNII: LKG8494WBH) .015 mL in 1 mL WATER (UNII: 059QF0K00R)

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:11695-1234-5	100 mL in 1 VIAL, MULTI-DOSE				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
unapproved drug other		06/01/2000				

Labeler - Covetrus North America (603750329)

Registrant - Sparhawk Laboratories, Inc. (147979082)

EstablishmentNameAddressID/FEIBusiness OperationsSparhawk Laboratories, Inc.147979082analysis, manufacture

Establishment						
Name	Address	ID/FEI	Business Operations			
Jiangxi Tianxin Pharmaceutical Co., Ltd.		527410270	api manufacture			

Revised: 12/2023 Covetrus North America