

HUMCO IODIDES TINCTURE- ethyl alcohol liquid
Humco Holding Group, Inc.

Humco Iodides Tincture

Drug Facts

Active Ingredients

Alcohol 45 % denatured with ammonia, Ammonium and Potassium Iodides.

Purpose

Antiseptic

Uses

First aid to help prevent infection in minor cuts, scrapes and burns.

Warnings

For external use only.

Ask a doctor before use if you have

deep or puncture wounds, animal bites, serious burns.

Stop use and ask a doctor if:

The condition persists or gets worse, or if using this product for longer than 1 week.

When using this product:

Do not use in the eyes. If contact occurs, flush with large amounts of water while lifting upper and lower lids. Do not apply over large areas of the body.

Keep out of the reach of children.

In case of accidental ingestion give milk, then give a starch solution made by mixing two tablespoonfuls of cornstarch or flour to a pint of water. contact a poison Control Center immediately.

Directions

Clean the affected area. Apply a small amount to the affected area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let it dry first.

Inactive Ingredient:

Purified Water

Other Information:

Will stain skin and clothing

New label



HUMCO IODIDES TINCTURE			
ethyl alcohol liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0395-1207
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.45 mL in 1 mL	
Inactive Ingredients			
Ingredient Name	Strength		
AMMONIA (UNII: 5138Q19F1X)			
IODINE (UNII: 9679TC07X4)			
POTASSIUM IODIDE (UNII: 1C4QK22F9J)			
WATER (UNII: 059QF0KO0R)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0395-1207-92	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/30/2017	
2	NDC:0395-1207-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/30/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/25/1998	

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(0395-1207) , analysis(0395-1207) , pack(0395-1207) , label(0395-1207)

Revised: 12/2023

Humco Holding Group, Inc.