IODO ALBO IODIDE- alcohol liquid liquid
Menper Distributors Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Iodo Albo Iodide

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

Ethyl Alcohol 48%

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

For External Use Only.

When using this product:

Do not use in the eyes or apply over large areas of the body. In case of deep or puncture wounds, animal bites or serious burns consult a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately. Soap will deactivate the effects of this product.

- Ammonium Hydroxide
- Iodine
- Potassium Iodide
- Purified Water

First Aid Antiseptic

Iodo Albo Iodide

IODO ALBO IODIDE

alcohol liquid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53145-120

Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.48 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
IODINE (UNII: 9679TC07X4)	
POTASSIUM IODIDE (UNII: 1C4QK22F9J)	
AMMONIA (UNII: 5138Q19F1X)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:53145-120-91	30 mL in 1 CONTAINER; Type 0: Not a Combination Product	04/09/2019	

Marketing Information

Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph not final	part333A	04/09/2019	

Labeler - Menper Distributors Inc. (101947166)

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