PRACTICAL CARE HAND SANITIZER- ethyl alcohol liquid Lindie Nzie Trading and Business Services CC

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.

Hand Sanitizer

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - PDP and Information Panel (Front and Back of Bottle) 250 mL NDC: 77682-001-01



Antiseptic Hand Sanitizer Ethyl Alcohol 70% Non-sterile Solution

Manufactured by - Linzie Nzie Trading, South Africa Imported and Distributed by BJM Products Inc. 6 Cedarhill Park Drive #13 Plymouth MA 02360 • 800-685-1859

Purpose

.Antiseptic

PRACTICAL CARE HAN	ND SANITIZER				
ethyl alcohol liquid					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:77682-001	
Route of Administration	TOPICAL				
Active Ingradient/Active Mei	a t v				
Active Ingredient/Active Moi					
Ingred	lient Name		Basis of Strength		Strength
ALCOHOL (UNII: 3K9958V90M) (ALC	COHOL - UNII:3K9958V90M)		ALCOHOL		70 mL in 100 mL
Inactive Ingredients					
U U	Ingredient Name				Strength
GLYCERIN (UNII: PDC6A3C0OX)					
HYDROGEN PEROXIDE (UNII: BBX06	0 AN9 V)				
WATER (UNII: 059QF0KO0R)					

Packaging					
# I	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC	C:77682-001-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020		
Mar	keting Info	ormation			
	keting Inf o		Marketing Start Date	Marketing End Date	
Mar	Ŭ	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	

Labeler - Lindie Nzie Trading and Business Services CC (538995764)

Revised: 7/2020

Lindie Nzie Trading and Business Services CC