CLEACE 75% ALCOHOL HAND SANITIZER- alcohol gel Warehouse Agency Llc

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.

CLEACE 75% Alcohol hand sanitizer

Active Ingredient

Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Uses

- hand sanitizer to help decrease bacteria on the skin, when water, soap & towel are not available.
- recommended for repeated use.

Warnings

For external use only.

Flammable. Keep away from fire or flame.

Do not apply around eyes.

Do not use in ears & mouth.

When using this product

avoid contact with eyes. In case of contact flush eyes with water.

Stop use and ask a doctor if

redness or irritation develop and persist for more than 72 hours

Keep out of reach of children.

Children must be supervised in use of this product.

Directions

• place enough product into your palms and thoroughly spread on both hands.

• rub into skin until dry.

Other information

- store below 110F (43C)
- may discolor certain fabrics or surfaces.

Inactive ingredients

Carbomer, Triethanolamine, Water

Package Label - Principal Display Panel

500 mL NDC: 80125-001-01

A CLEACE HOME

IS A CLEAN HOME

CLEACE

75% INSTANT

HAND

SANITIZER

Kills 99% of germs

Contain 75% Ethyl Alcohol

16.9 FL OZ

Drug Facts

500mL

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Drug Facts

Active ingredient Purpose
Ethyl Alcohol 75% v/v......Antiseptic

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CLEACE 75% ALCOHOL HAND SANITIZER

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80125-001(NDC:74621-002)	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	7.5 mg in 1 mL		

Inactive Ingredients		
Ingredient Name	Strength	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
WATER (UNII: 059QF0KO0R)		
TROLAMINE (UNII: 903K93S3TK)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80125-001- 01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/13/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	08/13/2020		

Labeler - Warehouse Agency Llc (113134877)

Revised: 2/2023 Warehouse Agency Llc