HAND SANITIZER- alcohol gel ESKBIOCHEM S.A. DE C.V.

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.

Active Ingredient(s)

Alcohol 75% v/v.

Purpose

Antimicrobial, 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
- do not use in or near eyes

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

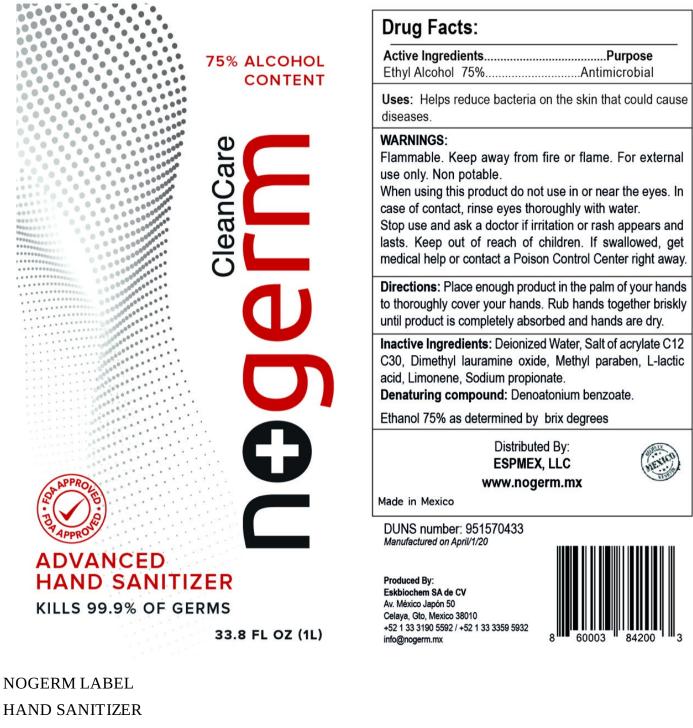
- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

Inactive ingredients

nogerm label 75%



KILLS 99.9% OF GERMS

ndc74589-009-01

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74589-009
Route of Administration	TOPICAL		
Active Ingredient/Ac	ctive Moiety		
	Ingredient Name	Basis of Streng	th Strength
LCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL		75 mL in 100 mL	
Inactive Ingredients			
	Ingredient Name		Strength
WATER (UNII: 059QF0KO	0 R)		
TROLAMINE (UNII: 903KS	93S3TK)		
CARBOMER INTERPOL V	MER TYPE A (55000 CPS) (UNII: 59TL3W	<i>1</i> G5CO)	
Child O ment in the OE I	MER I IPE A (55000 CPS) (UNII. 591L5W	(4566)	
	MER 1 1PE A (55000 CPS) (UNIL 59 115W		
Packaging	MER 1 1 PE A (55000 CPS) (UNII. 59 115W		
Packaging	Package Description		nte Marketing End Date
Packaging # Item Code		Marketing Start Da	nte Marketing End Date
Packaging # Item Code	Package Description	Marketing Start Da	nte Marketing End Dat
Packaging # Item Code	Package Description	Marketing Start Da	nte Marketing End Dat
Packaging # Item Code 1 NDC:74589-009-01 100	Package Description 0 mL in 1 BOTTLE; Type 0: Not a Combina	Marketing Start Da	nte Marketing End Dat
Packaging # Item Code	Package Description 0 mL in 1 BOTTLE; Type 0: Not a Combina	Marketing Start Da tion Product 04/15/2020	nte Marketing End Date

Labeler - ESKBIOCHEM S.A. DE C.V. (951570433)

Registrant - ESKBIOCHEM S.A. DE C.V. (951570433)

Establishment

Name	Address	ID/FEI	Business Operations
ESKBIOCHEM S.A. DE C.V.		951570433	manufacture(74589-009)

Revised: 4/2020

ESKBIOCHEM S.A. DE C.V.