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 80% 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

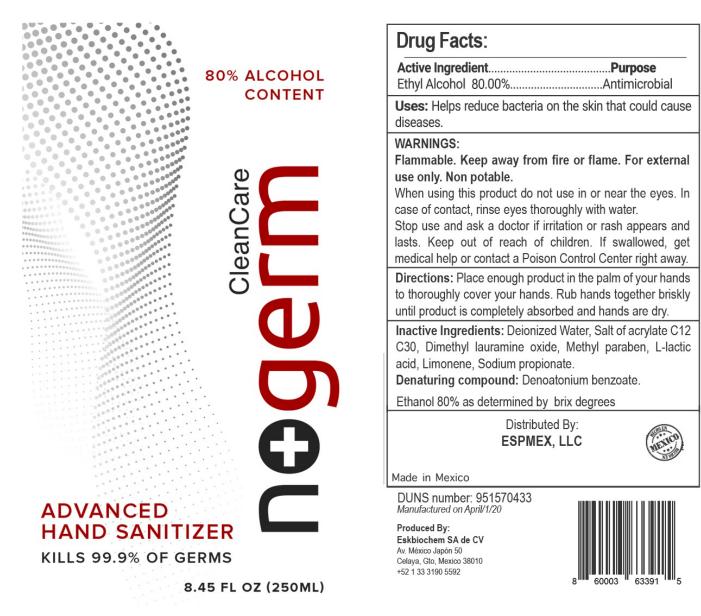
- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

Inactive ingredients

nogerm label 250ml



NDC:74589-005-03 NOGERM LABEL 250ML HAND SANITIZER KILLS 99.9% OF GERMS

HAND SANITIZER

alcohol gel

Product Information			
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:74589-005
Route of Administration	TOPICAL		

	t/Active Moiety				
	Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	80 mL in 250 mL		
		ALCOHOL			
Inactive Ingredie	nts				
	Ingredient Name		Strength		
CARBOMER INTERPO	DLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)				
TROLAMINE (UNII: 903K93S3TK)					
WATER (UNII: 059QF0KO0R)					
Packaging					
Packaging # Item Code	Package Description	Marketing Start Date	Marketing End Date		
# Item Code	Package Description 250 mL in 1 BOTTLE; Type 0: Not a Combination Product	•	Marketing End Date		
# Item Code	Ŭ Î	•	Marketing End Date		
# Item Code	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	•	Marketing End Date		
# Item Code 1 NDC:74589-005-03	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	•	Marketing End Date Marketing End Date		
 # Item Code 1 NDC:74589-005-03 Marketing Inference 	250 mL in 1 BOTTLE; Type 0: Not a Combination Product Ormation ry Application Number or Monograph Citation	04/17/2020			

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(74589-005)

Revised: 4/2020

ESKBIOCHEM S.A. DE C.V.