#### HAND SANITIZER- alcohol gel ESKIOCHEM 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.

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# ALL-CLEAN -PACKAGE LABEL



74589-002-01 All-Clean Hand Sanitizer Kills 99.99% of all germs

### Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antimicrobrial

### 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

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

Carbormer, Tricolamine, water

# ALL-CLEAN



NDC:74589-002-01 ALL-CLEAN HAND SANITIZER KILLS 99.99% OF ALL GERMS

HAND SANITIZER alcohol gel						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:74589-002		
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ingredient Name			Basis of Strength	n Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	70 mL in 100 mL		

Inactive Ingredients							
Ingredient Name							
TROLAMINE (UNII: 903K93S3TK)							
WATER (UNII: 059QF0KO0R)							
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)							
Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
<b>1</b> NDC:74589-002-01	1000 mL in 1 BOTTLE; Type 0: Not a Combi	nation Product 04/08/2020					
Marketing Information							
Marketing Catego	y Application Number or Monograp	oh Citation Marketing Start Date	Marketing End Date				
OTC monograph not fin	al part333A	04/08/2020					

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

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

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
ESKBIOCHEM S.A. DE C.V.		951570433	manufacture(74589-002)				

Revised: 4/2020

ESKIOCHEM S.A. DE C.V.