

SANETTO- alcohol gel
CERGOMEX IMPORTS 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.

Hand Sanitaizer GEL

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Hand Sanitizer Sanetto



Drugs Facts	
Active Ingredients	Purpose
Organic Cane Alcohol 70% v/v	Antiseptic
Use for hand-washing to decrease bacteria on the skin only when water is not available	
Warnings For external use only	
When using this product *do not get into eyes *if contact occurs, rinse eyes thoroughly with water	
Stop use and ask a doctor if *irritations and redness develops	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions. *Press pump twice to deliver two squirts (about a quarter size) of product onto the palm of your hand. *rub hands together until dry. *wash hands with soap and water at earliest opportunity.	
Inactive Ingredients Water, glycerin, carbomer.	

 **Made in Mexico**
 Produced by:
 JE INDUSTRIAS
 QUÍMICAS S.A. DE C.V.
 Distributed by:
 CERGOMEX Imports LLC
 NDC 75107-250-01
 cergomeximports@gmail.com
 Tel. (956) 236-2621




Alcohol 80% 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.

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.

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

Hand Sanitizer Gel Sanetto



125 mL NDC: 81101-200-01

3780 mL NDC 81101-200-02

SANETTO			
alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81101-200
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	70 mL in 100 mL
Inactive Ingredients			
Ingredient Name		Strength	

GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81101-200-02	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:81101-200-03	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:81101-200-04	70 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - CERGOMEX IMPORTS LLC (117484660)

Registrant - CERGOMEX IMPORTS LLC (117484660)

Establishment

Name	Address	ID/FEI	Business Operations
CERGOMEX IMPORTS LLC		117484660	manufacture(81101-200)

Revised: 4/2023

CERGOMEX IMPORTS LLC