ALCOHOL ANTISEPTIC 80% TOPICAL SOLUTION- alcohol solution NCH Corporation

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 Sanitizer

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.

Active Ingredient(s)

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.

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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.

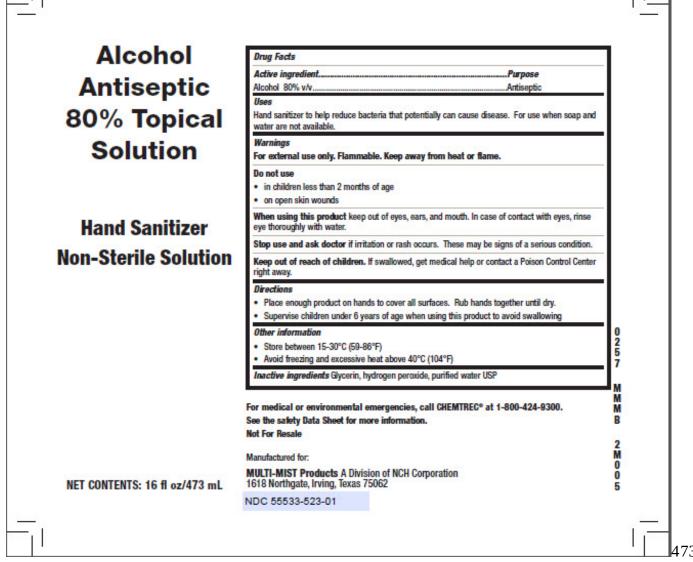
Other information

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

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel



NDC Code 55533-523-01

ALCOHOL ANTISEPTIC 80% TOPICAL SOLUTION

alcohol solution					
Product Inform	nation				
Product T ype	HUMAN OTC DRUG Item Cod			e (Source)	NDC:55533-523
Route of Administ	ration	TOPICAL			
Active Ingredie	ent/Active Moi	ety			
Ingredient Name				Basis of Strength	n Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)				ALCOHOL	80 mL in 100 mL
Inactive Ingred	lients				
Ingredient Name				Strength	
GLYCERIN (UNII: PDC6A3C0OX)				1.45 mL in 100 mL	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				0.125 mL in 100 mL	
WATER (UNII: 059Q	(FUKUUR)				
Packaging					
# Item Code		Package Description		Marketing Start Date	Marketing End Date
NDC:55533-523- 01 Product		'LE, PLASTIC; Type 0: Not a Con	ıbinatio n	05/15/2020	
Marketing Ir	nformation				
Marketing Categ	gory Applicat	ion Number or Monograph C	itation N	Marketing Start Date	Marketing End Date
OTC monograph not	final part333A		0	5/15/2020	

Labeler - NCH Corporation (126655562)

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NCH Corporation