

HAND SANITIZER 80ML- hand sanitizer gel, metered
NEW ZEALAND UNITED PHARMACEUTICALS LIMITED

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. Isopropyl Alcohol (75%, 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)

Isopropyl Alcohol 75% 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



500 ml NDC: 80400-001-02

HAND SANITIZER 80ML

hand sanitizer gel, metered

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80400-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	367.5000015 mL in 500 mL

Inactive Ingredients

Ingredient Name	Strength
TRIETHANOLAMINE CACODYLATE (UNII: M49TH27L7A)	1.15 mL in 500 mL
CENTELLA ASIATICA LEAF (UNII: 6810070TYD)	0.0000275 mL in 500 mL
CHRYSANTHELLUM INDICUM FLOWER OIL (UNII: HJ299SUZ05)	0.0000015 mL in 500 mL
ACRYLIC ACID (UNII: J94PBK7X8S)	1.5 mL in 500 mL
GLYCERIN (UNII: PDC6A3C0OX)	3 mL in 500 mL
WATER (UNII: 059QF0K00R)	126.8499865 mL in 500 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.0000275 mL in 500 mL
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	0.000005 mL in 500 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80400-001-02	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2020	

Labeler - NEW ZEALAND UNITED PHARMACEUTICALS LIMITED (594815782)

Establishment

Name	Address	ID/FEI	Business Operations
NEW ZEALAND UNITED PHARMACEUTICALS LIMITED		594815782	manufacture(80400-001) , pack(80400-001) , label(80400-001)

Revised: 9/2020

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