PANITA HAND SANITIZER- alcohol 73% v/v gel
Empack Spraytech Inc.

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.

This is a hand sanitizer rub 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) (73%, 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. Glycerin
c. Carbomer
d. Triethanolamine
e. Purfied water

Active Ingredient(s)

Alcohol 73% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Rub

Use

Health care personnel hand rub to help reduce bacteria that potentially can cause disease.

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

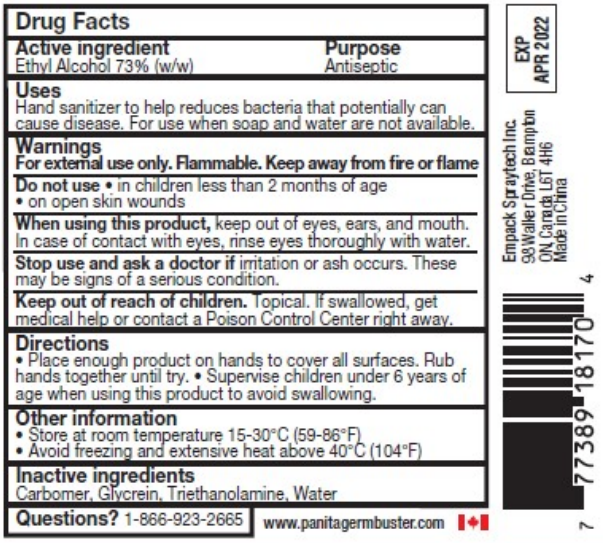
carbomer, glycerin, triethanolamine , purified water USP

Package Label - Principal Display Panel

3.4 fl.oz. (100 mL) NDC: 50021-010-01

8.1 fi.oz. (240 mL) NDC: 50021-010-02

16.9 fl.oz. (500 mL) NDC: 50021-010-03



PANITA HAND SANITIZER

alcohol 73% v/v gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50021-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	73 L in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	1 mL in 100 mL
CARBOMER 940 (UNII: 4Q93RCW27E)	0.45 in 100 mL

WATER (UNII: 059QF0KO0R)	25.3 mL in 100 mL
TRIETHANOLAMINE 2-CYCLOHEXYL-4,6-DINITROPHENOLATE (UNII: N2TK31JIAH)	0.25 in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50021-010-01	100 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/01/2020	
2	NDC:50021-010-02	240 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/01/2020	
3	NDC:50021-010-03	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

Labeler - Empack Spraytech Inc. (252047519)

Registrant - Empack Spraytech Inc. (252047519)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Rifeshow Cosmetics Co., Ltd.		560843477	manufacture(50021-010)

Revised: 4/2020 Empack Spraytech Inc.