PURIFIE- is opropyl alcohol liquid Brunner Indus trial Group

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 Santizier

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



946 ml NDC: 74271-542-32

HABLE RELAKE REL
HAND SANITIZER Isopropyl Alcohol Antisep 75% Topical Solution Hand Sanitizer
75% Topical Solution Hand Sanitizer
Non-sterile Solution
on Hano
Cleans Hands to Reduce Bacteria when soap &
water are not available
Brunner Industrial Group d.b.a. Smith Paint Product

118 ml NDC: 74271-542-14

roduct Type HUMAN OTC DRUG Item Code (Sour			nce) NDC:74271			
Route of Administration TOPICAL						
0.4x7						
Active Ingredient/Active Moiety						
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)			ISOPROPYL ALCOHOL			
Inactive Ingredients Ingredient Name						
GLYCERIN (UNII: PDC6A3C0OX)			1.45 mL in 100 mL			
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				0.125 mL in 100 mL		
	ety redient Name 416302) (ISOPROPYL ALCOHOL gredient Name	ety redient Name 416302) (ISOPROPYL ALCOHOL - gredient Name	ety redient Name Basis of Stree 416302) (ISOPROPYL ALCOHOL - ISOPROPYL ALCOHOL gredient Name 1.45 mL in 100	ety redient Name Basis of Strength 416302) (ISOPROPYL ALCOHOL - ISOPROPYL ALCOHOL gredient Name Streng 1.45 mL in 100 mL		

#	Item Code	Package Description	Marketing Start Date	Marketing End Date					
1	NDC:74271-542- 14	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020						
2	NDC:74271-542- 32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020						
N	Marketing Information								
	Marketing Cates	ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date					
0	TC monograph not	final part333A	03/30/2020						

Labeler - Brunner Industrial Group (003020062)

Registrant - Brunner Industrail Group (003020062)

Establishment

Name	Address	ID/FEI	Business Operations
Brunner Industrial Group		003020062	manufacture(74271-542)

Revised: 4/2020

Brunner Industrial Group