HAND SANITIZER- alcohol liquid The no nasties holdings pty ltd

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.

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

575 mL NDC 75740-001-00



VTISEPTIC 80 % • TOPICAL SOLUTION SANITIZER NON-STERILE SOLUTION





MATE IN AUSTRAL

60mL

DRUG FACTS

Active Ingredient(s)

Alcohol 80% v/v

Purpose

Antiseptic

Use(s)

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.

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 15–30°C (59–86°F) Avoid freezing and excessive heat about 40°C (104°F)

INACTIVE INGREDIENTS

Glycerin, Hydrogen Peroxide, Purifed Water USP

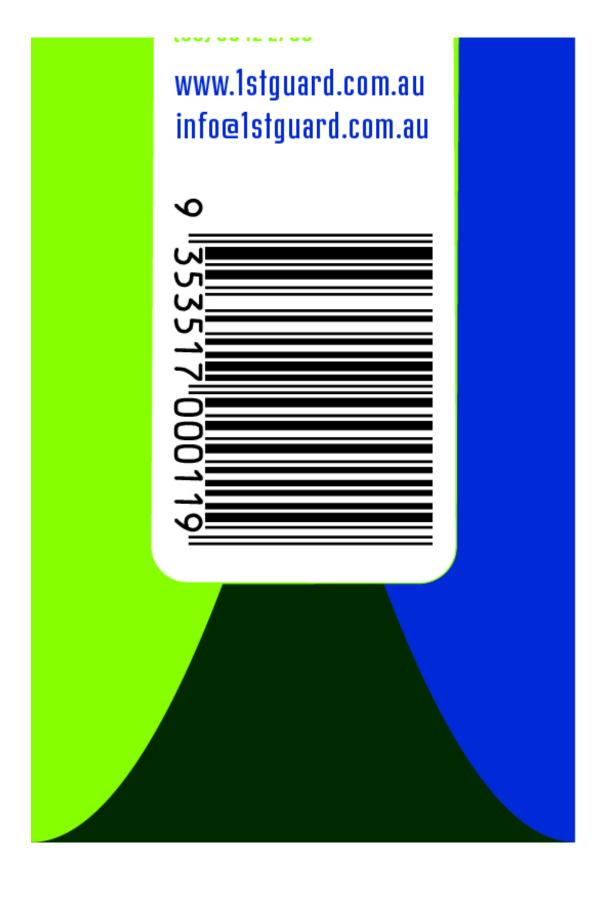


99.9%



DISTRIBUTED BY Royal Emerald Pharmaceuticals

Made in Australia by Inspi Beverages Pty Ltd 2/75a Chapel st Windsor 3181 (03) 9042 2765





HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75740-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL		

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6 A3C0 OX)	1.45 mL in 100 mL		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)			

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:75740-001-00	575 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			
Marketing Inf	ormation				
Marketing Inf		Marketing Start Date	Marketing End Date		
	ry Application Number or Monograph Citation	Marketing Start Date 03/30/2020	Marketing End Date		

Labeler - The no nasties holdings pty ltd (746897666)

Establishment					
Name	Address	ID/FEI	Business Operations		
The no nasties holdings pty ltd		746897666	manufacture(75740-001)		

Revised: 4/2020 The no nasties holdings pty ltd