HAND SANITIZER- isopropyl alcohol gel
Kennecott Utah Copper LLC

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

RioTinto

Isopropyl Alcohol Antiseptic 75%

Topical Solution

Hand Sanitizer
Non-sterile Solution

Do Not Ingest
Temporary Use Only

10,000 mL (338.14 oz)

Manufactured by
Kennecott Utah Copper LLC
4700 Daybreak Parkway
South Jordan, UT 84009

Americas Emergency Numbers
Toll Free (24 Hr)
+1 866 928 0789
Non-Toll Free (24 Hr)
+1 215 207 0061

Drug Facts

Active ingredient[s] ..... Isopropyl Alchol 75% v/v
Purpose ..... Antiseptic

Use[s]
Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

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Inactive ingredients glycerin, hydrogen peroxide, purified water USP

10,000 ml NDC: 74270-801-32

RioTinto

Isopropyl Alcohol Antiseptic 75%

Topical Solution

Hand Sanitizer
Non-sterile Solution

Do Not Ingest
Temporary Use Only

946.353 mL (32 oz)

Manufactured by
Kennecott Utah Copper LLC
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RioTinto

Isopropyl Alcohol Antiseptic 75%

Topical Solution

Hand Sanitizer
Non-sterile Solution

Do Not Ingest
Temporary Use Only

828.058 mL (28 oz)

Manufactured by
Kennecott Utah Copper LLC
4700 Daybreak Parkway
South Jordan, UT 84009

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+1 866 928 0789
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+1 215 207 0061

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Inactive ingredients glycerin, hydrogen peroxide, purified water USP

236 ml NDC: 74270-801-08

RioTinto

Isopropyl Alcohol Antiseptic 75%

Topical Solution

Hand Sanitizer
Non-sterile Solution

Do Not Ingest
Temporary Use Only

236.588 mL (8 oz)

Manufactured by
Kennecott Utah Copper LLC
4700 Daybreak Parkway
South Jordan, UT 84009

Americas Emergency Numbers
Toll Free (24 Hr)
+1 866 928 0789
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RioTinto

Isopropyl Alcohol Antiseptic 75%

Topical Solution

Hand Sanitizer
Non-sterile Solution

Do Not Ingest
Temporary Use Only

118.294 mL (4 oz)

Manufactured by
Kennecott Utah Copper LLC
4700 Daybreak Parkway
South Jordan, UT 84009

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Drug Facts

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HAND SANITIZER

isopropyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74270-801

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74270-801-04	118.294 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74270-801-08	236.588 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Kennecott Utah Copper LLC (619968779)

Revised: 6/2020 Kennecott Utah Copper LLC