RIPCLEAR ANTISEPTIC HAND SANITIZER- isopropyl alcohol liquid
Rip Clear 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.

RIPCLEAR® ANTISEPTIC HAND SANITIZER

Drug Facts

Active ingredient

Isopropyl alcohol 75% v/v

Purpose

Antiseptic

Uses • hand sanitizer to decrease bacteria on the skin • recommended for repeated use • for use when soap and water are not available

Warnings

Flammable, keep away from fire/flame
For external use only

Do not use • in children less than 2 months of age • on open skin wounds

When using this product • do not get into eyes. In case of contact, rinse eyes thoroughly with water

Stop use and ask a doctor if • irritation and redness develop • condition persists for more than 72 hours

Keep out of reach of children. If swallowed get a medical help or contact a Poison Control Center right away.

Directions • wet hands thoroughly with product and allow to dry without wiping • supervise children under 6 years of age when using this product to avoid swallowing

Other information

• store between 59-86°F (15-30°C) • avoid freezing and excessive heat above 104°F (40°C)

Inactive ingredients

acrylates/c-10-30 alkyl acrylate crosspolymer, aloe vera leaf juice, aminomethylpropanol, FD&C Blue #1, FD&C Red #40, fragrance, glycerin, propylene glycol, propylparaben, water

***Questions?* +1-917-968-3098**

You may also report serious side effects to this phone number.

Mon-Fri 9:00 AM - 5:00 PM

Topical solution

Liquid, Non-Gel

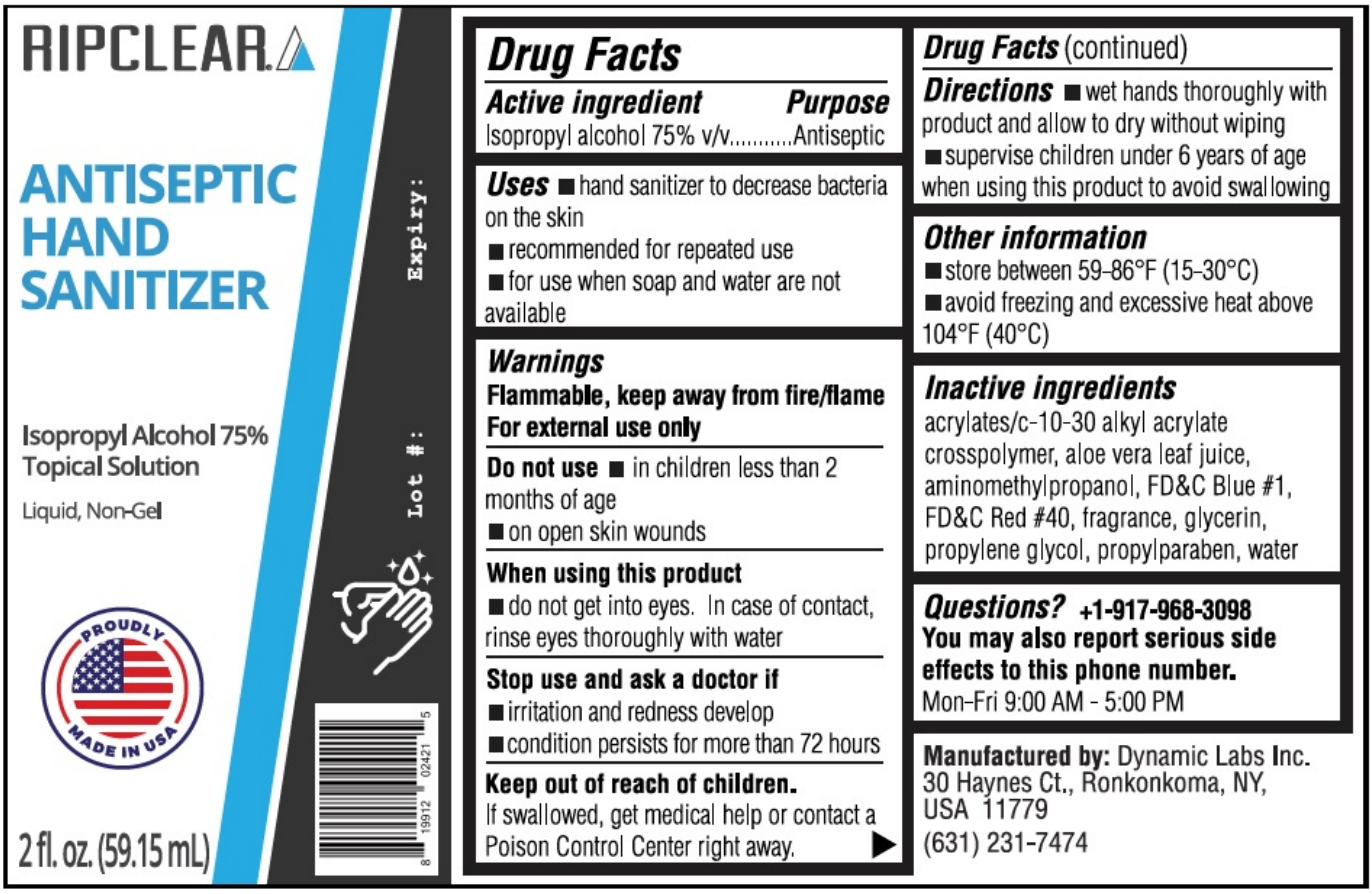
PROUDLY MADE IN USA

Manufactured by: Dynamic Labs Inc.
30 Haynes Ct., Ronkonkoma, NY, USA 11779

(631) 231-7474

Packaging

Package 59.15 mL



Package 236.6 mL

Package 3800 mL

RIPCLEAR ANTISEPTIC HAND SANITIZER

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80555-001

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
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	blue (Light blue)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80555-001-10	59.15 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/20/2020	
2	NDC:80555-001-20	236.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/20/2020	
3	NDC:80555-001-30	3800 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/20/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/20/2020	

Labeler - Rip Clear LLC (079813108)

Revised: 10/2020 Rip Clear LLC