RUE21 ICON HAND SANITIZER- ethyl alcohol gel Pearl World 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.

rue 21_® ICON HAND SANITIZER

Drug Facts

Active ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

- To decrease bacteria on the skin that could cause disease.
- Recommended for repeated use.

Warnings

For external use only-hands. Flammable. Keep away from heat and flame. When using this product • Keep out of eyes. In case of contact with eyes, flush thoroughly with water. • Avoid contact with broken skin. • Do not inhale or ingest. Stop use and ask a doctor if skin irritation develops.

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

- Wet hands thoroughly with product and allow to dry without wiping.
- For children under 6, use only under adult supervision.
- Not recommended for infants.

Other information

• Do not store above 105°F. • May discolor some fabrics. • Harmful to wood finishes and plastics.

Inactive Ingredients

Water (Aqua), Glycerin, Propylene Glycol, Carbomer, Aloe Barbadensis Leaf Extract, Triethanolamine, Disodium EDTA, Tocopheryl Acetate (Vitamin E), Fragrance.

PEEL UP TO SEE ADDITIONAL DRUG FACTS

Manufactured exclusively for rue21.

Warrendale, PA 15086 Designed in the U.S.A.

Packaging









DRUG FACTS LABEL



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BATCH NO.: SA8228765 MFG: 20200805 EXP: 20220804

RUE21 ICON HAND SANITIZER

ethyl alcohol gel

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:69933-209 Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Basis of Strength	Strength			
ALCOHOL	70 mL in 100 mL			

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
TROLAMINE (UNII: 9O3K93S3TK)				
EDETATE DISO DIUM ANHYDRO US (UNII: 8 NLQ36 F6 MM)				
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)				

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:69933-209-30	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 21				
Marketing Information							

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	0 1/0 1/20 21	

Labeler - Pearl World Inc. (043130142)

Revised: 1/2021 Pearl World Inc.