

RUE21 BLACK 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.

BLACK rue21 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.

Manufactured exclusively for rue21®

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

Made in China. www.rue21.com

Packaging



DRUG FACTS LABEL

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\$2.99 4 00276 83796 4	

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RUE21 BLACK HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69933-206
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C00X)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69933-206-30	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/17/2020	

Labeler - Pearl World Inc. (043130142)

Revised: 8/2020

Pearl World Inc.