PREMIER PURE PROTECTION- alcohol gel RPP Products, 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.

Active Ingredient

Ethyl Alcohol 72% v/v.

Purpose

Antimicrobial

Use

Hand sanitizer to help reduce bacteria on the skin.

Warnings

Flammable. Keep away from fire or flame

For external use only.

When using this product do not use in or near the eyes. In case of contact rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

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

Directions

- Put enough product in your palm to cover hands and rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using PREMIER PURE.

Other information

- Store below 110° F (43° C).
- May discolor certain fabrics or surfaces.
- This product does not contain any chemicals known to the State of California to cause cancer, birth defects, or any other reproductive harm.

Inactive ingredients

Water, Acrylates Copolymer, Glycerin, Hydrolyzed Jojoba Esters, Aminomethyl Propanol, Fragrance

Product Label

Drug Facts

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GEL Hand Sanitizer

72% ETHYL ALCOHOL

- MAY KILL 99.9% OF GERMS
- REFRESHING SCENT
- HYDRATING MOISTURIZERS
 3 FL OZ (89 mL)



PREMIER PURE PROTECTION

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73945-005

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)
ALCOHOL (UNII: 3K9958 V90M) ALCOHOL 72 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ACRYLATES/VINYL ISODECANO ATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8 MDB79 NA)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROLYZED JOJOBA ESTERS (POTASSIUM SALTS) (UNII: CH428W5O62)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

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Item Code Package Description Marketing Start Date Date

NDC-730 4E OOE EO mI in 1 DOTTI E DI ACTIC: Type O. Not a Combination

1	NDC:/3945-005-	59 III. III 1 BO I I LE, PLASTIC; Type 0: Not a Combination Product	06/08/2020
2	NDC:73945-005- 02	89 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2020
3	NDC:73945-005- 03	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2020
4	NDC:73945-005- 04	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2020
5	NDC:73945-005- 05	1893 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2020
6	NDC:73945-005- 06	$3785\ mL$ in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2020

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	06/08/2020				

Labeler - RPP Products, Inc. (623623852)

Revised: 6/2020 RPP Products, Inc.