RELIEF C HAND SANITIZER UNSCENTED FOR PROFESSIONALS (2 OZ)- ethyl alcohol gel FGD, 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.

Relief C Hand Sanitizer Unscented for Professionals (2 oz) - rapidgel (alcohol)

Ethyl Alcohol 70% v/v

Antimicrobial

Hand sanitizer to help reduce bacteria that can potentially cause disease. For use when soap and water are not available

For external use only. Flammable. Keep away from heat or flame.

- on children less than 2 months of age
- on open skin wounds

keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

if irritation or rash occurs. These may be signs of a serious condition.

If swallowed, get medical help or contact a Poison Control Center right away.

- Wet hands thoroughly with product and allow to dry without wiping.
- Rub lightly until dry.
- Store between 15-30 °C (59-86 °F)
- Avoid freezing and excessive heat 40 °C (104 °F)

Glycerin, PEG-6 (and) AMP-Acrylate/vinyl-Isodecanoate Crosspolymer, Propylene-glycol, Purified Water

Relief C Hand Sanitizer Unscented for Professionals (2 oz) - rapidgel (alcohol)



Drug Facts

Active Ingredients Pu
Ethyl Alcohol 70% v/v.......Antir

Warnings

For external use only. Flammable. Keep away fro or flame.

Avoid contact with eyes. In case of contact, rinse ethoroughly with water immediately.

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

Stop and ask a doctor if irritation or rash occurs. To may be signs of a serious condition.

Directions

 Wet hands thoroughly with product and allow to dry without wiping.
 Rub lightly until dry.
 Do not rinse.

Other Information

Store below 110°F (43°C). Avoid freezing and exces

Inactive Ingredients Glycerin, PEG-6 (and)
AMP-Acrylates/Vinyl Isodecanoate Crosspolymer, P
Glycol, Purified Water.



RELIEF C HAND SANITIZER UNSCENTED FOR PROFESSIONALS (2 OZ)

ethyl alcohol gel

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73787-132		
Route of Administration	TOPICAL				

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

Inactive Ingredients				
Ingredient Name	Strength			
ACRYLATES/VINYL ISODECANO ATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8 MDB 79 NA)				
WATER (UNII: 059QF0KO0R)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
GLYCERIN (UNII: PDC6A3C0OX)				
POLYETHYLENE GLYCOL 300 (UNII: 5655G9 Y8 AQ)				

Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:73787-132-01	DC:73787-132-01 59 mL in 1 BOTTLE; Type 0: Not a Combination Product					
Marketing Information						
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not fi	nal part333E	08/03/2020				

Labeler - FGD, LLC (111927555)

Establishment						
Name	Address	ID/FEI	Business Operations			
Goodwin Co.		806987483	manufacture(73787-132)			

Revised: 1/2021 FGD, LLC