RELIEF C HAND SANITIZER UNSCENTED FOR PROFESSIONALS- 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 8 oz rapidgel

Active ingredient

Ethyl Alcohol 70%

Purpose

Antimicrobial

Uses

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

Warnings

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.

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using hit produce to avoid swallowing.

- Store between 15-30 ^{o}C (59-86 ^{o}F)
- Avoid freezing and excessive heat 40 ^{o}C (104 ^{o}F)

Inactive ingredients

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

Relief C Hand Sanitizer Unscented for Professionals 8 oz rapidgel

RELIEF C HAND SANITIZER UNSCENTED FOR PROFESSIONALS

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73787-119
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**
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	

WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73787-119-01	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/11/2020	

Labeler - FGD, LLC (111927555)

Establishment

Name	Address	ID/FEI	Business Operations
Goodwin Co.		806987483	manufacture(73787-119)

Revised: 1/2021 FGD, LLC