NEW WAVE HAND SANITIZER GEL- alcohol gel BRENNTAG MID-SOUTH, 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(s)



Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria on the skin.

Warnings

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

Do not use

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

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

Keep out of reach of children. 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 this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Inactive Ingredients: Water, Acrylates/Vinyl Isodecanoate Crosspolymer (10000 MPA.S Neutralized at 0.5%), Aloe Vera Leaf Extract, Isopropyl Myristate, Propylene Glycol, Fragrance, Vitamin E Acetate, Yellow 10, Yellow 5, Blue 1

Product Name



NEW WAVE HAND SANITIZER GEL

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65353-8442
Route of Administration	TOPICAL		

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

Inactive Ingredients		
Ingredient Name	Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ALOE (UNII: V5VD430YW9)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
WATER (UNII: 059QF0KO0R)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65353- 8442-1	1032 L in 1 TANK; Type 0: Not a Combination Product	10/19/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	10/19/2020	01/31/2025	

Labeler - BRENNTAG MID-SOUTH, INC. (122625064)

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