PUMPKIN HAND SANITIZER- alcohol gel Reaction Retail, 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.

Pumpkin Hand Sanitizer

Drug Facts

Active Ingredient:

Ethyl Alcohol 75%

Antiseptic

Uses:

Hand sanitizer to help decrease bacteria on the skin.

Warnings:

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

Stop use and ask doctor if

irritation or rash appears and lasts.

Keep out of reach of children.

If swallowed, get medical help or contact a doctor right away.

Directions:

Squirt as needed into your palms and thoroughly spread on both hands. Rub into skin until dry.

Other Information:

Store below 118 F

Inactive Ingredients:

Aqua (Water), Glycerin, Propylene Glycol, Aloe Barbadensis Leaf Juice, Parfum (Fragrance), Acrylates/C10-30 Alkyl Acrylate Crosspolymer,

Triethanolamine, Citric Acid, Potassium Sorbate, Sodium Benzoate, Sucrose, Zea Mays (Corn) Starch, Hydroxypropyl Methylcellulose, Polyvinyl Alcohol, Benzyl Benzoate, Eugenol, Coumarin, Cinnamyl Alcohol, Amyl Cinnamal, Cinnamal, Benzyl Alcohol, CI 77267 (D&C Black No. 3), CI 19140 (Yellow 5), CI 14700 (Red 4)

Package Labeling:30ml





Made in TURKEY Distributed by Reaction Retail 1010 Westmore Ave, Rockville, MD 20850

www.reactionretail.com





Drug Facts
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Other Information: Store below 118°F

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Iriethanolamine, Citric Acid, Potassium Sorbate, Sodium Benzoate, Sucrose, Zea Mays (Corn) Starch, Hydroxypropyl Methylcellulose, Polyvinyl Alcohol, Benzyl Benzoate, Eugenol, Coumarin, Cinnamyl Alcohol, Amyl Cinnamal, Cinnamal, Benzyl Alcohol, CI 77267 (D&C Black No. 3), CI 19140 (Yellow 5), CI 14700 (Red 4)



Package Labeling:35ml



Drug Facts

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\$2.00





PUMPKIN HAND SANITIZER

alcohol gel

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:80026-007

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.75 mL in 1 mL

mactive ingredients					
Ingredient Name	Strength				
WATER (UNII: 059QF0KO0R)					

GLYCERIN (UNII: PDC6A3C0OX)

Inactive Inquedient

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)
TROLAMINE (UNII: 9O3K93S3TK)
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)
SODIUM BENZOATE (UNII: OJ245FE5EU)
SUCROSE (UNII: C151H8M554)
CORN (UNII: 0N8672707O)
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)
BENZYL BENZOATE (UNII: N863NB338G)
EUGENOL (UNII: 3T8H1794QW)
COUMARIN (UNII: A4VZ22K1WT)
CINNAMYL ALCOHOL (UNII: SS8 YOP444F)
.ALPHAAMYLCINNAMALDEHYDE (UNII: WC51CA3418)
CINNAMALDEHYDE (UNII: SR60A3XG0F)
BENZYL ALCOHOL (UNII: LKG8494WBH)
D&C BLACK NO. 2 (UNII: 4XYU5U00C4)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)

I	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:80026-007-30	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020		
2	NDC:80026-007-35	35 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/15/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	09/15/2020		

Labeler - Reaction Retail, LLC (968085212)

Revised: 8/2020 Reaction Retail, LLC