SPECTRUM HEALTH - AMWAY HAND SANITIZER- alcohol gel
Access Business Group LLC

Spectrum Health - Amway Hand Sanitizer

Drug Facts

Active Ingredient

Ethyl alcohol 73% v/v

Purpose

Antiseptic

Uses

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

Warnings

For external use only. Flammable, keep away from heat or flame.

Do not use

- In children less than 2 years of age.
- On open skin woundsOn open skin wounds

When using this product

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

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 thoroughly into hands for at least 30 seconds. Allow to dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

Inactive ingredients

Water, Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Propylene Glycol, Triethanolamine, Panthenol, Aloe Barbadensis Leaf Juice, Maltodextrin, Sodium Benzoate, Potassium Sorbate

Questions?

USA 1.800.253.6500

Package Labeling:

SPECTRUM HEALTH - AMWAY HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10056-048
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	73 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	
PANTHENOL (UNII: WV9CM0O67Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10056-048-00	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/01/2021	

Labeler - Access Business Group LLC (839830713)

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