HAND SANITIZER- alcohol spray HAAN BRAND SL

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.

Hand Sanitizer Morning Glory

Uses

Uses

* To decrease bacterias on the hand. Recommended for repeated use.

Active Ingredient

Active Ingredient Purpose

• Ethyl Alcohol 65%.....Antiseptic

Warnings

- For external use only- hands
- **When using this product** keep out o **f** eyes. In case of contact, flush throughly with water.
- Do Not Inhale or ingest.
- Avoid contact with broken skin
- **Keep out of reach of children.** In case of incidental ingestion seek professional assistance or contact a Poison Control Center inmediately.
- **Stop use and ask Doctor:** if skin irritation develops.
- **Flammable:** Keep away from heat and flame.

Stop use and ask a doctor if

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* Irritation develops

Keep out of reach of children

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Directions

Directions: Wet hands throughly with product and allow to dry without wiping

 For Children under 6 years of age do not use without supervision of an adult. Not recomended for infants

Inactive Ingredients

Water, Water, C10-30 Alkyl Acrylate Crosspolymer, Parfum(Fragrance), Limonene, Tromethamine, Linalool, Hexyl Cinnamal, Potassium Sorbate, Citral, Sodium Benzoate



Questions?

QUESTIONS OR COMMENTS: Visit Haanready.com

Uses

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Hand Sanitizer Gel



Morning Glory



Hydrating Hand Sanitizer

Net Wt. 1 fl oz (30 ml)

HAND SANITIZER

alcohol spray

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79091-009

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
CARBOMER 934 (UNII: Z135WT9208)	
METHYL GLUCETH-10 (UNII: NOMWT4C7WH)	
WATER (UNII: 059QF0KO0R)	
LIMO NENE, (+)- (UNII: GFD7C86Q1W)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
LINALOOL , (+)- (UNII: F4VNO44C09)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TROMETHAMINE (UNII: 023C2WHX2V)	
BUTYL ACRYLATE/C16-C20 ALKYL METHACRYLATE/METHACRYLIC ACID/METHYL METHACRYLATE COPOLYMER (UNII: 7K68DGG29P)	

l	Packaging			
	# Item Code Package Description		Marketing Start Date	Marketing End Date
l	1 NDC:79091-009- 01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/18/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/18/2020	

Labeler - HAAN BRAND SL (468886727)

Registrant - Haan Brand SL (468886727)

Establishment			
Name	Address	ID/FEI	Business Operations

Laboratorios Magrina SL	460356033	manufacture(79091-009)	
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Establishment			
Name	Address	ID/FEI	Business Operations
HAAN BRAND SL		468886727	label(79091-009)

Revised: 8/2020 HAAN BRAND SL