EMHS1 HAND SANITIZER- alcohol gel Sayida Group 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.

EMHS1 Hand Sanitizer (gel - 1 Gal) bottle, pump NDC: 77858-0753-1 EMHS1 Hand Sanitizer (gel - 1 Gal) bottle NDC: 77858-0753-2

Active Ingredient(s)

Ethyl Alcohol 70% v/v.

Purpose

Antiseptic

Use

To help reduce bacteria that can potentially cause disease.

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

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.

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

Do not store above 43C (110F)

Inactive ingredients

Carbomer, Isopropyl alcohol, Quaternary ammonium, Water

Package Label - Principal Display Panel

DISTRIBUTED BY: Sayida Group

6538 Collins Ave #401

Miami Beach, FL 33141

EMHS1 Hand Sanitizer (bottle, pump - 1Gal) NDC: 77858-0753-1

EMHS1 Hand Sanitizer (bottle - 1Gal) NDC: 77858-0753-2



Drug Facts

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Distributed by:

Sayida Group 6538 Collins Ave #401 Miami Beach, FL 33141 www.emhs1.com Made in Mexico

EMHS1 HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77858-0753

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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	Ingredient Name	Basis of Streng	th Strength
ALCOHOL (UNII: 3K9958V	90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 L in 100 L

Inactive Ingredients			
Ingredient Name	Strength		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM30 7FC)			
ISOPROPYL ALCOHOL (UNII: ND2M416302)			
WATER (UNII: 059QF0KO0R)			

F	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:77858-0753- 1	0.26 L in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/16/2020			
2	NDC:77858-0753- 2	0.26 L in 1 BOTTLE; Type 0: Not a Combination Product	06/16/2020			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	06/11/2020				

Labeler - Sayida Group LLC (117520711)

Revised: 6/2020 Sayida Group LLC