HAND SANITIZER GEL- alcohol solution
Hill & Markes, 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.

Hand Sanitizer Gel

Moisturizing with Aloe

Effective at eliminating more than 99.9% of many common harmful germs & bacteria in as little as 15 seconds.

Active Ingredient(s)

Ethyl alcohol 70% v/v **Purpose** Antimicrobial

Purpose

Antimicrobial

Use

Hand sanitizer to help reduce bacteria on the skin that could cause disease Recommended for repeated use.

Warnings

Flammable. Keep away from fire or flame.

For external use only

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. 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.

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Directions

Place enough product in your palm to thoroughly cover your hands Rub hands together briskly until dry

No rinsing required

No towels needed

Other information

Do not store above 110°F (43°C) May discolor certain fabrics or surfaces.

Inactive ingredients

Water (Aqua), Aloe Vera, Propylene Glycol, Carboxethyl Acrylate Crosspolymer, Fragrance, FD&C Blue #1, FD&C Yellow #5

Package Label - Principal Display Panel

709.76 mL NDC: 78046-1124-1

HAND SANITIZER GEL

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78046-1124
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE (UNII: KBC00G95HI)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HEPTANAL (UNII: 92N104S3HF)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78046-1124-1	709.76 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/19/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/19/2020	

Labeler - Hill & Markes, Inc. (013616594)

Revised: 5/2020 Hill & Markes, Inc.