HAND CLEANSING GEL- hand sanitizer llc gel Hand Sanitizer 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.

Ethyl Alcohol 70%

Antiseptic

Hand Cleanser to help reduce bacteria on the skin.

Flammable, Keep away from fire or flame.

For External use only

When using this product do not use in or near the eys. In case of contact, rinse eyes intensely with water.

Stop Use and ask a doctor if irritation or rash occurs and remains.

Keep out of reach of children. If swallowed get medical help or contact a Poison Control Center immediately.

Put enough product in your palm to cover hands well and rub hands together until dry. Children under 6 years old should be supervised when using.

Store under 110°F (43°C)

Water (Aqua), Hydroxypropyl Cellulose, Glycerin, Fragrance



Active Ingredients Purpose			
Ethyl Alcohol 70% v/v	Antiseptic		
Use - Hand Cleanser to help reduce	e bacteria on the skin		
Warnings Flammable, Keep away from fire o	r flame.		
For external use only			
When using this product do not use case of contact, rinse eyes intensely	e in or near the eyes. In with water.		
Stop use and ask a doctor if irritation or rash occurs and remains.			
Keep out of reach of children. If swallowed get medical help or contact a Poison Control Center immediately.			
Directions . Put enough product in hands well and rub hands together ur 6 years old should be supervised who	ntil dry . Children under		
Other information. Store under 1	10° F (43° C)		
Inactive ingredients Water (Aqua), Hydroxypropyl Cellulose, Glycerin, Fragrance			
Distributed by: Hand Cleansing G 850 Kaliste Saloom			

17 FL OZ (502mL)

www.HandSanitizerLLC.com Sales@HandSanitizerLLC.com

70% Alcohol Cleansing Gel

HAND CLEANSING GEL

hand sanitizer llc gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76701-700
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)		
GLYCERIN (UNII: PDC6A3C0OX)		
HYDRO XYPRO PYL CELLULO SE, UNSPECIFIED (UNII: 9 XZ8 H6 N6 O H)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76701-700- 17	17 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/16/2020	
2	NDC:76701-700- 02	2 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/16/2020	
3	NDC:76701-700- 08	8 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/16/2020	
4	NDC:76701-700- 32	32 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/16/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/16/2020	

Labeler - Hand Sanitizer LLC (117473019)

Establishment			
Name	Address	ID/FEI	Business Operations
Hand Sanitizer LLC		117473019	pack(76701-700), label(76701-700), manufacture(76701-700)

Revised: 4/2020 Hand Sanitizer LLC