HAND SANITIZER- ethyl alcohol gel
Zhejiang Hongshiliang Health Technology Co., Ltd.

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.

75230-001 Hand Sanitizer

Active Ingredient

Ethyl Alcohol 65%

Purpose

Antiseptic

Use

Use To decrease bacteria on the skin

WARNINGS

Flammable. Keep away from fire or flame.
For external use only
When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water. Stop use and ask a doctor if irritation or rash appears and lasts.

keep out of reach of children

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

Directions

Directions●wet hands thoroughly with product and allow to dry without wiping●for children under 6, use only under adult supervision●not recommended for infants
Other information
●do not store above 110°F(43°C)
●may discolor some fabrics
●harmful to wood finishes and plastics

Inactive ingredients

Water, Carbomer,Triethanolamine, Aloe vera, Tocopheryl acetate,Glycerin, Fragrance

Drug Facts

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Distributed by PlasmaDent, Inc.
3809 Mojave Ct, Suite E, Columbia, MO 65202

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HAND SANITIZER
ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75230-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TOCOPHERYL RETINOATE (UNII: 0WN694NBMM)	
CARBOMER 934 (UNII: Z135WT9208)	

TRIETHANOLAMINE BENZOATE (UNII: M3EN4GC19W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75230-001-01	60 in 1 CARTON	03/21/2020	
1		1 in 1 BOX		
1		37 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:75230-001-02	120 in 1 CARTON	03/21/2020	
2		1 in 1 BOX		
2		37 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:75230-001-03	3 in 1 CARTON	03/21/2020	
3		60 in 1 CARTON		
3		1 in 1 BOX		
3		37 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:75230-001-04	18 in 1 CARTON	03/21/2020	
4		153 mL in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:75230-001-05	24 in 1 CARTON	03/21/2020	
5		153 mL in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:75230-001-06	30 in 1 CARTON	03/21/2020	
6		153 mL in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:75230-001-07	48 in 1 CARTON	03/21/2020	
7		153 mL in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:75230-001-08	18 in 1 CARTON	03/21/2020	
8		307 mL in 1 BOTTLE; Type 0: Not a Combination Product		
9	NDC:75230-001-09	24 in 1 CARTON	03/21/2020	
9		307 mL in 1 BOTTLE; Type 0: Not a Combination Product		
10	NDC:75230-001-10	54 in 1 CARTON	03/21/2020	
10		153 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/21/2020	

Labeler - Zhejiang Hongshiliang Health Technology Co., Ltd. (546641397)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Hongshiliang Health Technology Co., Ltd.		546641397	manufacture(75230-001)

Revised: 4/2020 Zhejiang Hongshiliang Health Technology Co., Ltd.