

HAND SANITIZER- ethyl alcohol gel
QUANZHOU BESTHOPE HOUSEHOLD PRODUCTS 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.

50049-001 Hand Sanitizer

DRUG FACTS

Active ingredient

Ethyl Alcohol 75% v/v

Purpose

Antiseptic

Uses

For Hand Sanitizing

Warnings

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water
- Avoid contact with open skin
- Do not inhale or ingest
Stop use and ask a doctor
- If skin irritation or redness develops

Keep out of reach of children.

- Keep out of reach of children. Children under 6 years of age use only under adult supervision
- Not recommended for infants

Directions

- Wet hands thoroughly with product and allow to dry without wiping

Other information

- 1 Do not store above 105°F
- 1 May discolor some fabrics

Inactive ingredients

Alcohol, Water (Aqua), Glycerin, Butylene Glycol, Carbomer, Aminomethyl Propanol, Aloe Barbadensis Leaf Extract

Package Labeling



HAND SANITIZER

8 FL OZ (237ml)

Drug Facts

Active ingredients	Purpose
Ethyl Alcohol 75%(V/V).....	Antiseptic

Uses

- For Hand Sanitizing

Warnings

- For external use only
- Flammable. Keep away from fire or flame

When using this product

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water
- Avoid contact with open skin
- Do not inhale or ingest

Stop use and ask a doctor

- If skin irritation or redness develops

Directions

- Wet hands thoroughly with product and allow to dry without wiping
- Keep out of reach of children. Children under 6 years of age use only under adult supervision
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8 oz Item No: MED1103 ASI-90075

HiPatron™ Quanzhou besthope household products co.,ltd.

Best if used by: 7/2022

Batch Code: 04/2020

Made in China

HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50049-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)

ALOE VERA LEAF (UNII: ZY81Z83H0X)

BUTYLENE GLYCOL (UNII: 3XUS85K0RA)

CARBO MER 934 (UNII: Z135WT9208)

GLYCERIN (UNII: PDC6A3C0OX)

AMINO METHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50049-001-01	150 in 1 CARTON	04/04/2020	
1		45 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:50049-001-02	100 in 1 CARTON	04/04/2020	
2		75 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:50049-001-03	24 in 1 CARTON	04/04/2020	
3		187.5 mL in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:50049-001-04	20 in 1 CARTON	04/04/2020	
4		375 mL in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:50049-001-05	18 in 1 CARTON	04/04/2020	
5		750 mL in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:50049-001-06	200 in 1 CARTON	04/04/2020	
6		22.5 mL in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:50049-001-07	150 in 1 CARTON	04/04/2020	
7		37.5 mL in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:50049-001-08	24 in 1 CARTON	04/04/2020	
8		112.5 mL in 1 BOTTLE; Type 0: Not a Combination Product		
9	NDC:50049-001-09	24 in 1 CARTON	04/04/2020	
9		135 mL in 1 BOTTLE; Type 0: Not a Combination Product		
10	NDC:50049-001-10	24 in 1 CARTON	04/04/2020	
10		150 mL in 1 BOTTLE; Type 0: Not a Combination Product		
11	NDC:50049-001-11	24 in 1 CARTON	04/04/2020	
11		225 mL in 1 BOTTLE; Type 0: Not a Combination Product		
12	NDC:50049-001-12	24 in 1 CARTON	04/04/2020	
12		367.5 mL in 1 BOTTLE; Type 0: Not a Combination Product		
13	NDC:50049-001-13	12 in 1 CARTON	04/04/2020	
13		900 mL in 1 BOTTLE; Type 0: Not a Combination Product		
14	NDC:50049-001-14	12 in 1 CARTON	04/04/2020	
14		1500 mL in 1 BOTTLE; Type 0: Not a Combination Product		
15	NDC:50049-001-15	4 in 1 CARTON	04/04/2020	
15		2835 mL in 1 BOTTLE; Type 0: Not a Combination Product		
16	NDC:50049-001-16	4 in 1 CARTON	04/04/2020	
16		3750 mL in 1 BOTTLE; Type 0: Not a Combination Product		

17	NDC:50049-001-17	2 in 1 CARTON	04/04/2020	
17		7500 mL in 1 BOTTLE; Type 0: Not a Combination Product		
18	NDC:50049-001-18	1 in 1 CARTON	04/04/2020	
18		18750 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	04/04/2020	

Labeler - QUANZHOU BESTHOPE HOUSEHOLD PRODUCTS CO., LTD. (529121531)

Establishment

Name	Address	ID/FEI	Business Operations
QUANZHOU BESTHOPE HOUSEHOLD PRODUCTS CO.,LTD.		529121531	manufacture(50049-001)

Revised: 4/2020

QUANZHOU BESTHOPE HOUSEHOLD PRODUCTS CO., LTD.