

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.

75230-003 Hand Sanitizer

Active Ingredient

Ethyl Alcohol 75%

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

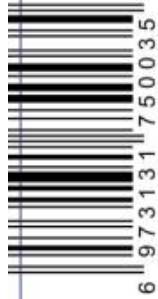
- do not store above 110°F(43°C)
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

Water, ,Glycerin,Carbomer,Triethanolamine



ShineX RINSE FREE
HAND SANITIZER, K-
EEPS YOUR HANDS
CLEAN AND HYDRA-
TED.



Distributed by

ADD.

NDC:

Drug Facts

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hand sanitizer

ShineX

RINSE-FREE
HAND
SANITIZER

8 fl oz (236.6ml)

HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75230-003
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: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER 934 (UNII: Z135WT9208)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75230-003-02	88.7 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
2	NDC:75230-003-03	177.4 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	

3	NDC:75230-003-07	150 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
4	NDC:75230-003-04	354.8 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
5	NDC:75230-003-06	75 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
6	NDC:75230-003-05	709.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
7	NDC:75230-003-01	44.3 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
8	NDC:75230-003-08	187.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
9	NDC:75230-003-09	375 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	
10	NDC:75230-003-10	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/09/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/08/2020	

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

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Hongshiliang Health Technology Co., Ltd.		546641397	manufacture(75230-003)

Revised: 5/2020

Zhejiang Hongshiliang Health Technology Co., Ltd.