QUICK-DRYING HAND SANITIZER- hand sanitizer liquid
Shandong Kehong Medical 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.

Active Ingredient(s)

Isopropyl Alcohol 75%,Chlorhexidine Gluconate 1.1%

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

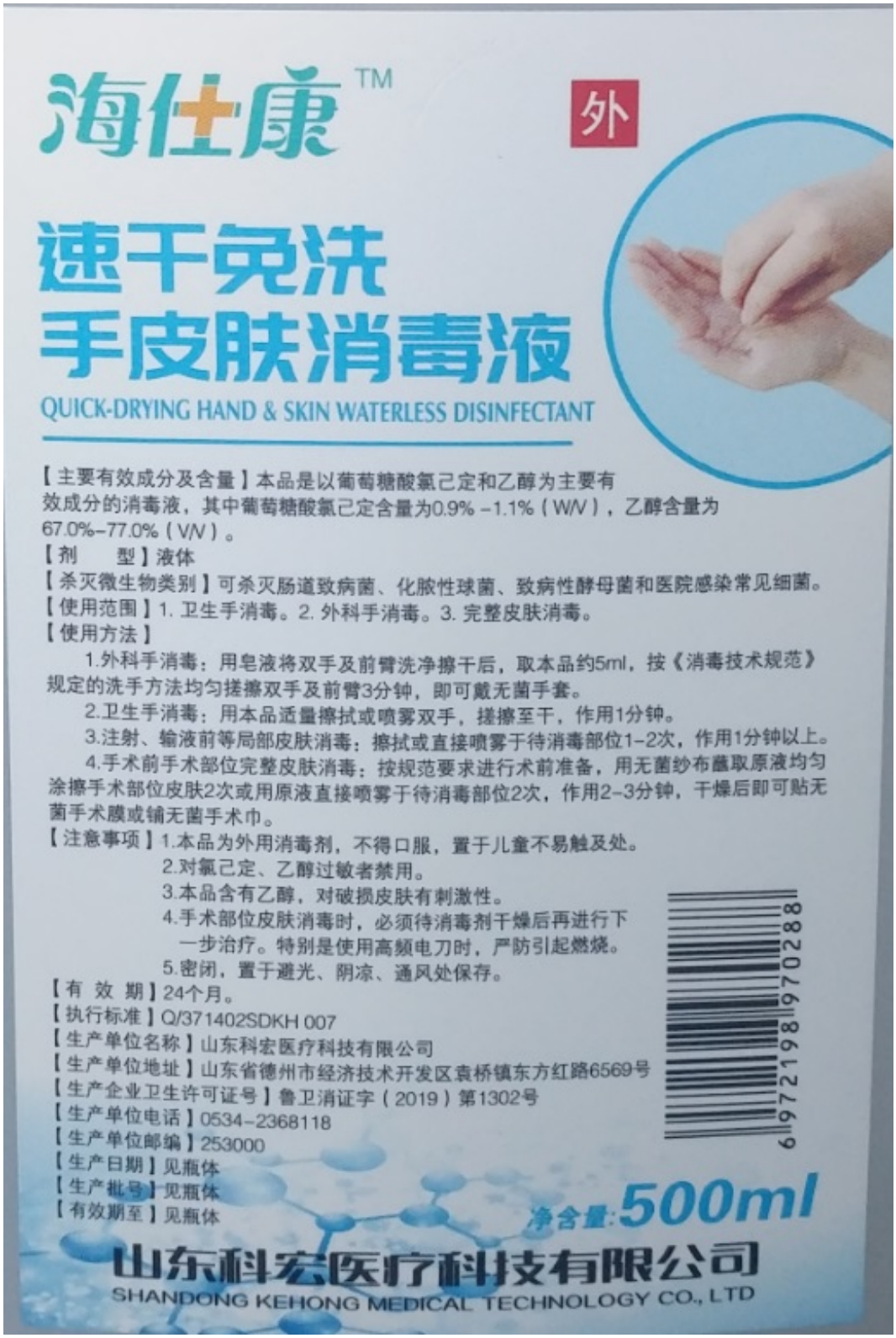
Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

water

Packaging Labels - Main Display Panel



QUICK-DRYING HAND SANITIZER

hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75244-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE GLUCONATE	1.1 g in 100 g
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	75 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75244-001-01	100 g in 1 BOTTLE; Type 0: Not a Combination Product	04/14/2020	
2	NDC:75244-001-02	500 g in 1 BOTTLE; Type 0: Not a Combination Product	04/14/2020	
3	NDC:75244-001-03	1000 g in 1 BOTTLE; Type 0: Not a Combination Product	04/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/14/2020	

Labeler - Shandong Kehong Medical Technology Co.,Ltd. (554527496)

Establishment

Name	Address	ID/FEI	Business Operations
Shandong Kehong Medical Technology Co.,Ltd		554527496	manufacture(75244-001)

Revised: 4/2020 Shandong Kehong Medical Technology Co.,Ltd.