KEKE HAND SANITIZER 750ML 01- alcohol liquid Shenzhen Lantern Scicence 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.

KEKE hand sanitizer 750ml

Drug Facts



Active Ingredient

Active Ingredient Purpose Ethyl Alcohol 75% [V/V] Antiseptic

Use

Hand sanitizer helps to reduce bacteria that potentially can cause disease.

Warning

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

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Flammable, keep away from heat and flame.

Discontinue if skin becomes irritated and ask a doctor .

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a poson control center immediately.

When using this product

do not get into eyes, If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor

If iritation or rash appears and lasts.

keep out of reach of children

If swallowed,get medical help or contact a Poison Control Center right away. Recommended for repeated use. use anywhere without water.

Directions

place enough product in your palm rub hands vigorously until dry children under 6 years of age should be supervised when using this product. When using this product keep out of eyes. In case of contact with eyes, flush thoroughly with water. Do not inhale or ingest. Avoid contact with broken skin.

Other information

Do not store above 105F. May discolor some fabrics. Harmful to wood finishes and plastics.

Inactive ingredients

Water(Aqua), Glycerin, Propylene Glycol,, Carbomer, Aminomethyl propanol Fragrance,

Other information

Store between 106°F[]41°C). May discolor certain fabrics or surfaces.

Purpose

Antiseptic



KEKE HAND SANITIZER 750ML 01 alcohol liquid									
Product Information									
Product Type	HUMAN OTC DRUG	ltem Code (Source)							
Route of Administration	TOPICAL								
Active Ingredient/Active Moiety									
Ingredient Name			Basis of Stren	ngth					
ALCOHOL (UNII: 3K9958V90M) (AI	COHOL - UNII:3K9958V90M)		ALCOHOL						

Inactive Ingredients				
Ingredient Name	Strength			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.01 mL in 100 mL			
CARBOMER 940 (UNII: 4Q93RCW27E)	0.35 mL in 100 mL			

NDC:54860-372

Strength 75 mL in 100 mL

	AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			0.11 mL in 100 mL					
WATER (UNII: 059QF0KO0R)				24.42 mL in 100 mL					
GLYCERIN (UNII: PDC6A3C0OX)				0.01 mL in 100 mL					
LEMON (UNII: 24RS0A9880)				0.1 mL in 100 mL					
Product Characteristics									
Color		white (transparant)	S	Score					
Sł	hape		S	Size					
Flavor			I	Imprint Code					
Co	Contains								
Packaging									
				Marketing Start					
#	Item Code	Package Description		Date	Marketing End Date				
	NDC:54860- 7	Package Description 50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a combination Product							
	NDC:54860- 7	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a		Date					
1	NDC:54860- 7 372-01 C	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a		Date					
1	NDC:54860- 7 372-01 C	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a combination Product		Date					
1	NDC:54860- 372-01	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		Date 08/03/2020 Marketing Start	Date Marketing End				

Labeler - Shenzhen Lantern Scicence Co.,Ltd. (421222423)

Registrant - LANTERN HEALTH&BEAUTY LAB INC. (086860340)

Establishment									
Name	Address	ID/FEI	Business Operations						
Shenzhen Lantern Science Co., Ltd.		421222423	manufacture(54860-372)						

Revised: 2/2021

Shenzhen Lantern Scicence Co.,Ltd.