

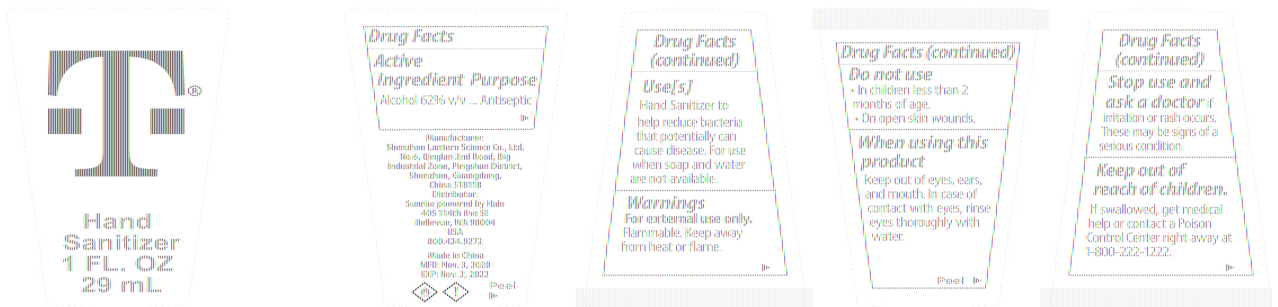
HAND SANITIZER 29ML 02- alcohol gel

Shenzhen Lantern Science 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.

Hand sanitizer 29ml

Drug Facts



Active Ingredient

Active Ingredient Purpose

Ethyl Alcohol 62% V/V Antiseptic

keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away at 1-800-222-1222

Recommended for repeated use.

use anywhere without water.

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 poison control center immediately.

Inactive ingredients

Aminomethyl propanol, Water, Carbomer, Deatonium Benzoate, Glycerin, Parfum, Propylene glycol.

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.

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.

When using this product

Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Purpose

Antiseptic

Stop use and ask a doctor

if irritation or rash occurs, these may be signs of a serious condition.

Use

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

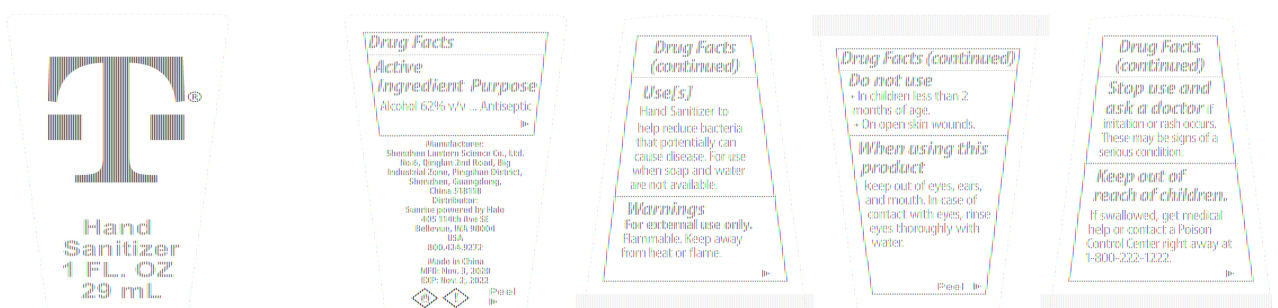
Other information

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

Do not use

in children less than 2 months of age

on open skin wounds.



alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-347
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER 940 (UNII: 4Q93RCW27E)	0.28 mL in 100 mL
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	0.0005 mL in 100 mL
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.1 mL in 100 mL
LEMON (UNII: 24RS0A988O)	0.1 mL in 100 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.1 mL in 100 mL
WATER (UNII: 059QF0KO0R)	37.3195 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	0.1 mL in 100 mL

Product Characteristics

Color	white (transparent)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-347-01	29 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/28/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333A	01/28/2021	

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

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

Establishment			
Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-347)

Establishment			
Name	Address	ID/FEI	Business Operations
Greater Pacific Industries,Inc.		962000253	label(54860-347)

Revised: 1/2021

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