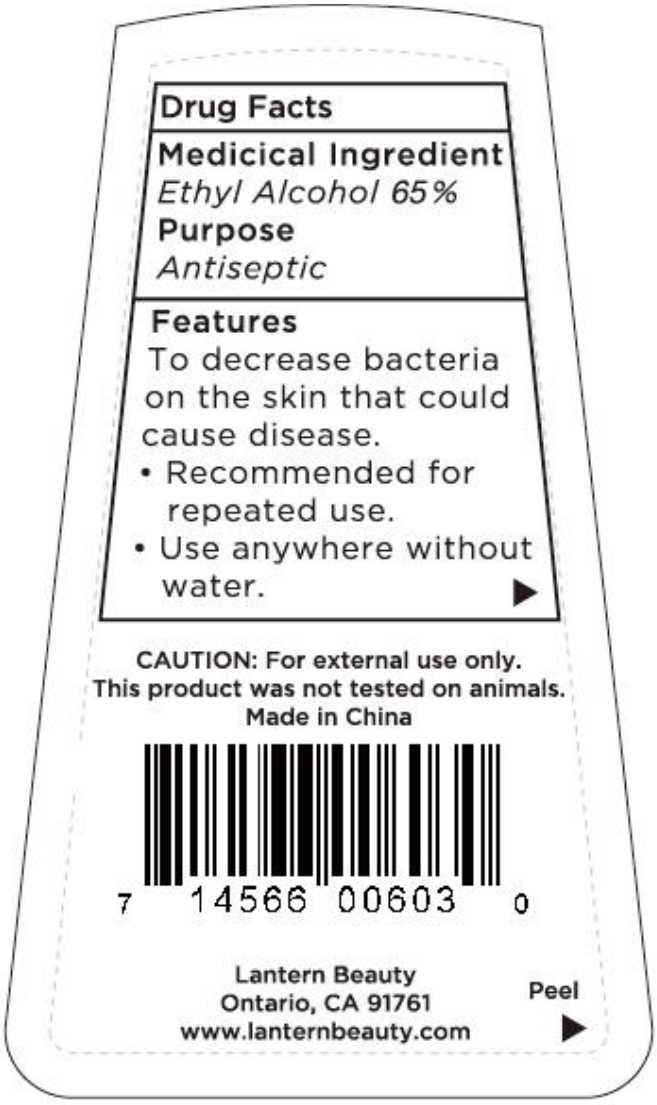
HAND SANITIZER 29ML 01- alcohol liquid
Shenzhen Lantern Scicence Co.,Ltd.

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.

Hand sanitizer 29ml

Drug Facts

drug facts



Active Ingredient

Active Ingredient Purpose

Ethyl Alcohol 65%□V/V□ Antiseptic

Use

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

Use(s)

To decrease bacteria on the skin that could cause disease.Recommended for repeated use.

Warning

For external use only-hands

Flammable.Keep away from heat and flame.

For external use only.

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.

keep out of reach of children

In case of accidental ingestion,seek professional assistant or contact a Poison Control Center immediately.

Recommended for repeated use.

use anywhere without water.

Inactive ingredients

Water(Aqua),Fragrance,Carbomer,CI 19140,CI 16255.

Directions

Wet hands thoroughly with product and rub until dry without wiping.Children under 6,use only with adult supervision.

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

Do not use in or near eyes,In case of eye contact ,flush eyes thoroughly with water. Discontinue if skin becomes irritated and ask for a doctor.

Other information

store below 105℉

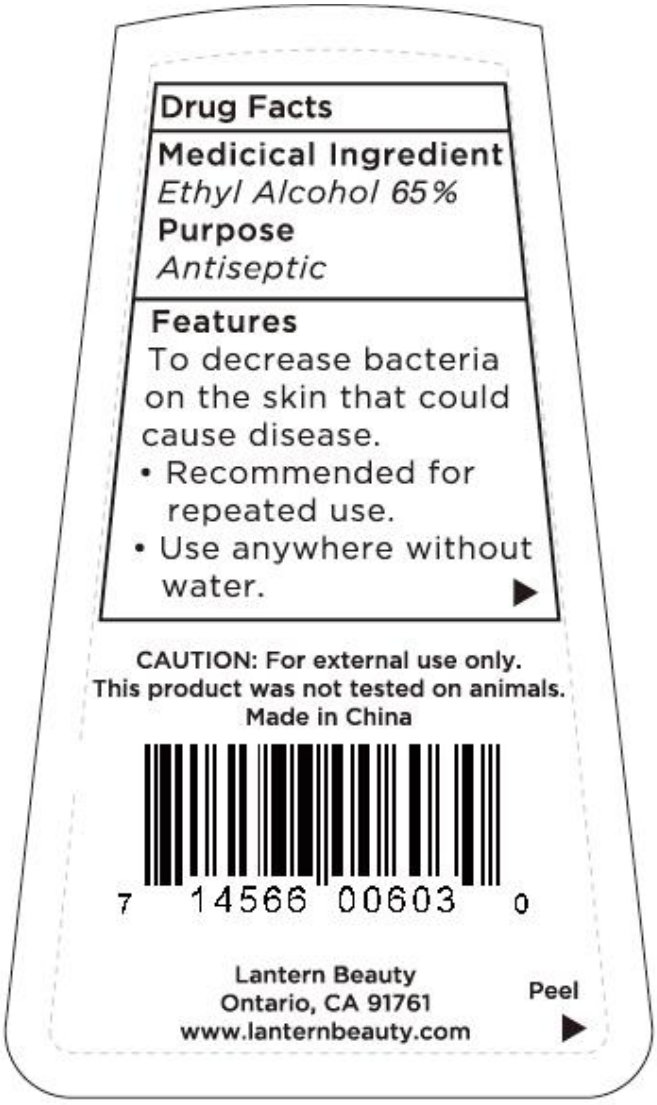
May discolor some fabrics and surfaces.

Purpose

Antiseptic

Packaging

Packaging



HAND SANITIZER 29ML 01

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-414
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	34.7397 mL in 100 mL
CARBOMER 940 (UNII: 4Q93RCW27E)	0.26 mL in 100 mL

FD&C RED NO. 4 (UNII: X3W0AM1JLX)	0.0002 mL in 100 mL
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.0001 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-414-01	29 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/23/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/23/2023	

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

Registrant - Lantern Beauty America,INC. (117371139)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-414)

Revised: 5/2023 Shenzhen Lantern Scicence Co.,Ltd.