

NIMBUS 9 HAND SANITIZER 3.78L 01- alcohol liquid
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.

Nimbus 9 hand sanitizer 3.78L

Drug Facts



Active Ingredient

Active Ingredient Purpose

Ethyl Alcohol 71% w/w Antiseptic

USE

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

Recommended for repeated use.

use anywhere without water.

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 poison control center immediately.

Inactive ingredients

Water(Aqua),Glycerin,Propylene Glycol,Carbomer,Aminomethyl Propanol,Parfum.

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.

Keep out of reach of children

If swallowed,get medical help or contact a Poison Control Center right away.

Do not use

in children less than 2 months of age.

on open skin wounds.

Stop use and ask a doctor

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

Other information

Store between 15-30 C(59-86F)

Avoid freezing and excessive heat above 40C(104F)

材质 白PE过光油



155 mm

银色效果

135 mm

银色效果



NIMBUS 9 HAND SANITIZER 3.78L 01

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-263
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	71 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
LEMON (UNII: 24RS0A988O)	0.1 g in 100 g
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.01 g in 100 g
CARBOMER 940 (UNII: 4Q93RCW27E)	0.35 g in 100 g
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.11 g in 100 g
WATER (UNII: 059QF0KO0R)	28.42 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	0.01 g in 100 g

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-263-01	3780 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/11/2020	

Labeler - Shenzhen Lantern Science Co Ltd (421222423)

Registrant - Lantern Beauty America,INC. (117371139)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-263)

Revised: 5/2020

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