

MEIJER HAND SANITIZER 59ML 01- alcohol gel
Meijer, Inc.

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.

meijer hand sanitizer 59ml

Drug Facts



52mm



47mm

52mm



47mm

Drug Facts (continued)

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

Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

Inactive Ingredients
Water, Isopropyl Alcohol, Aloe Barbardensis Leaf Juice, Glycerin, Carbomer, Aminomethyl Propanol, Fragrance, Propylene Glycol, Isopropyl Myristate, Tocopherol Acetate, FD&C Yellow #5 (Tartazine), FD&C Blue #1

Drug Facts (continued)

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

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 doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Active Ingredient

Active Ingredient Purpose

Ethyl Alcohol 62% Antiseptic

Use(s)

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.

Warnings

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

Aqua, Isopropyl Alcohol, Aloe Barbadensis Leaf Water, Glycerin, Carbomer, Aminomethyl Propanol, Fragrance, Propylene Glycol, Isopropyl Myristate, Tocopheryl Acetate, FD&C Yellow #5 (Tartazine), FD&C Blue #1.

Directions

Place enough product on hands to cover all surfaces. Rub hands together until dry. Supervise children under 6 years old 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.

Do not use

in children less than 2 months of age.

on open skin wounds.

Keep out of reach of children

If swallowed, get medical help or contact a poison control center right away.

Stop use and ask doctor if

irritation or rash occurs, These may be signs of serious condition.

Other information

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alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-0060
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
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	0.001 mL in 100 mL
ALOE (UNII: V5VD430YW9)	0.1 mL in 100 mL
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	0.000024 mL in 100 mL
WATER (UNII: 059QF0KO0R)	37.197916 mL in 100 mL
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.05 mL in 100 mL
CARBOMER 940 (UNII: 4Q93RCW27E)	0.26 mL in 100 mL
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.09 mL in 100 mL
GLYCERIN (UNII: PDC6A3C0OX)	0.1 mL in 100 mL
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	0.001 mL in 100 mL
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.00006 mL in 100 mL
ISOPROPYL ALCOHOL (UNII: ND2M416302)	0.1 mL in 100 mL
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	0.1 mL in 100 mL

Product Characteristics

Color	green (light green)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79481-0060-1	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/04/2020	

Labeler - Meijer, Inc. (006959555)

Registrant - LANTERN HEALTH&BEAUTY LAB INC (086860340)

Establishment

Name	Address	ID/FEI	Business Operations
Meijer, Inc.		006959555	label(79481-0060)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(79481-0060)

Revised: 12/2020

Meijer, Inc.