MORNING FRESH HAND SANITIZER- ethyl alcohol gel
Meijer Distribution, 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 538

Active Ingredients

Ethyl Alcohol 62%

Purpose

Antiseptic

Use

- to decrease bacteria on the skin that could cause disease
- Recommended for repeated use

Warnings

For external use only: hands

Flammable

Keep away from heat and flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor

- Irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children

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

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105º
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

benzopheone-4, blue 1, carbomer, fragrance, glycerin, isopropyl alcohol, isopropyl myristate, tocopheryl acetate, water

claims

Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds.

Adverse Reactions

DISTRIBUTED BY

MEIJER DISTRIBUTION, INC.

GRAND RAPIDS, MI 49544

www.meijer.com

Questions 1-888-593-0593

principal display panel

meijer

Hand

Sanitizer

Morning Fresh

Kills 99.99%

of germs

1.11 FL OZ (33 mL)

MORNING FRESH HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41250-538
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	Alcohol	558 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:41250-538-06	33 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/21/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/21/2016	

Labeler - Meijer Distribution, Inc (006959555)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(41250-538)

Revised: 5/2022 Meijer Distribution, Inc