

WINTER APPLE- hand sanitizer 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.

Hand Sanitizer Winter Apple
548.000/548AA

Active ingredient

Ethyl alcohol 62%

Purpose

Antiseptic

Uses

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

Warnings

For external use only: hands

Flammable. Keep away from fire or 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 if

- 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 hand 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 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glycerin, carbomer, fragrance, red 40, red 33

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

DISTRIBUTED BY

MIEJER DISTRIBUTION, INC

GRAND RAPIDS, MI 49544

www.meijer.com

Questions 1-999-593-0593

principal display panel

meijer

Hand Sanitizer

Winter Apple

Kills 99.99% of germs

2 fl oz (59 mL)



WINTER APPLE

hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-119
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:41250-	59 mL in 1 BOTTLE, PLASTIC, Type 0; Net		

1	NDC:41250-119-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/24/2016	
2	NDC:41250-119-10	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/24/2016	
3	NDC:41250-119-06	33 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/24/2016	
4	NDC:41250-119-34	237 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/24/2016	
5	NDC:41250-119-38	296 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/24/2016	
6	NDC:41250-119-32	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/24/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/24/2016	

Labeler - Meijer Distribution, Inc (006959555)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(41250-119)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(41250-119)

Revised: 12/2022

Meijer Distribution, Inc