

HAND SANITIZER MOJITO MINT- organic ethyl alcohol spray
Whole Foods Market, 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.

Whole Foods 365 Mojito MInt Hand Sanitizer Spray 2 fl oz

Active ingredient Purpose

Organic Ethyl Alcohol 62% Antiseptic

Uses ■ to help reduce bacteria on the skin.

For external use only. Flammable. Keep away from fire or flame.

When using this product avoid contact with eyes. If eye contact occurs, rinse with water.

Stop use and ask a doctor if skin irritation or redness develops.

Keep out of reach of children. In case of accidental ingestion, seek medical help or contact a Poison Control Center immediately.

Directions ■ apply 1 to 2 sprays on hands and rub together until dry

Other information ■ do not store above 104°F (40°C).

Inactive ingredients water, isopropyl alcohol, fragrance (natural)*, glycerin, natural fragrance, helianthus annuus (sunflower) seed oil, rosmarinus officinalis (rosemary) leaf extract, triethyl citrate.

*Complies with ISO 9235, see www.wholefoodsmarket.com for more information.

Questions, Comments? ■ customer.questions@wholefoods.com ■ 1-844-936-8255

NDC 42681-0008-1



PEEL HERE



mojito mint HAND SANITIZER

Kills 99.9% of Germs
With Sunflower Oil & Glycerin
to Replace Moisture

2 FL OZ (59mL)



HAND SANITIZER MOJITO MINT

organic ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42681-0008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
ROSEMARY (UNII: IJ67X351P9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42681-0008-1	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	12/31/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2020	12/31/2024

Labeler - Whole Foods Market, Inc. (196175616)**Registrant** - V Manufacturing & Logistics, Inc. (825176857)

Revised: 2/2020

Whole Foods Market, Inc.