NATURALS POMEGRANATE AND MANGO ANTIBACTERIAL HAND- alcohol gel New Avon LLC

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.

Naturals Pomegranate Mango Antibacterial Hand Gel

Active Ingredient

Ethyl Alcohol 64.76%.....

Purpose

.....Antiseptic

Uses

• For handwashing to decrease bacteria on the skin

Warnings

For external use only

When using this product

Avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a health care practitioner if

• irritation and redness develop

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

FLAMMABLE. Keep away from open flame and sources of heat.

Directions

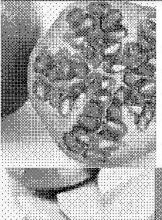
• Wet hands thoroughly with product and allow to dry without wiping.

Inactive ingredients

WATER/EAU
GLYCERIN
PUNICA GRANATUM FRUIT EXTRACT
MANGIFERA INDICA (MANGO) FRUIT EXTRACT
CARBOMER
PARFUM/FRAGRANCE
HYDROXYPROPYLCELLULOSE
TRIETHANOLAMINE

Questions? Call 1-800-FOR-AVON

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pomegranate & mango antibasterial hand gel

30 ml 1 fl. oz.

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Olrections

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Questions? Call 1-800-FDR-AVON

LABEL PATENT #7,601,410

Do not use if seal is broken.

Kills 99.99% of bacteria. Wild and gentle to skin.

To order, see your Avon Independent
Sales Representative or call,
1-800-FOR-AVON.
Www.avon.com
MADE IN THE USA
AVON PRODUCTS, INC.
NEW YORK, N.Y. 10017
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alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10096-0252
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	19.428 mL in 30 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10096-0252-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/14/2011	
2	NDC:10096-0252-2	75 mL in 1 TUBE; Type 0: Not a Combination Product	04/14/2011	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	04/14/2011		

Labeler - New Avon LLC (080143520)

Establishment				
Name	Address	ID/FEI	Business Operations	
Fareva Morton Grove, Inc.		116752326	manufacture(10096-0252)	

Revised: 1/2019 New Avon LLC