NATURALS VANILLA 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.

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

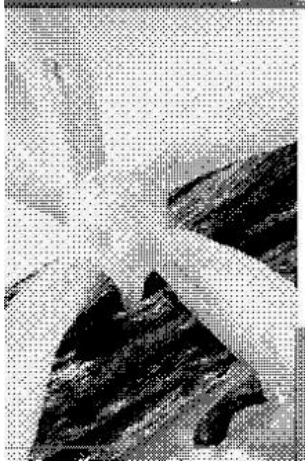
- Wet hands thoroughly with product and allow to dry without wiping.

Inactive ingredients

WATER/EAU
GLYCERIN
VANILLA PLANIFOLIA FRUIT EXTRACT
GLYCINE SOJA (SOYBEAN) SEED EXTRACT
PARFUM/FRAGRANCE
HYDROXYPROPYLCELLULOSE
CARBOMER
TRIETHANOLAMINE

Questions? Call **1-800-FOR-AVON**

AVON
naturals



vanilla
antibacterial
hand gel

30 ml 1 fl. oz.

hand

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

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

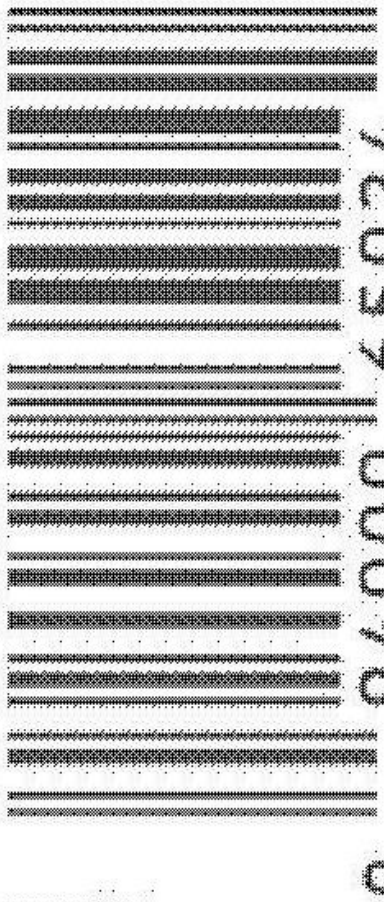
Inactive Ingredients WATER/EAU, GLYCERIN, VANILLA PLANIFOLIA FRUIT EXTRACT, GLYCINE SOJA (SOYBEAN) SEED EXTRACT, PARFUM/FRAGRANCE, HYDROXYPROPYLCELLULOSE, CARBOMER, TRIETHANOLAMINE.

Questions? Call **1-800-FOR-AVON**

LABEL PATENT #7,601,410

Do not use if seal is broken.

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



Drug Facts

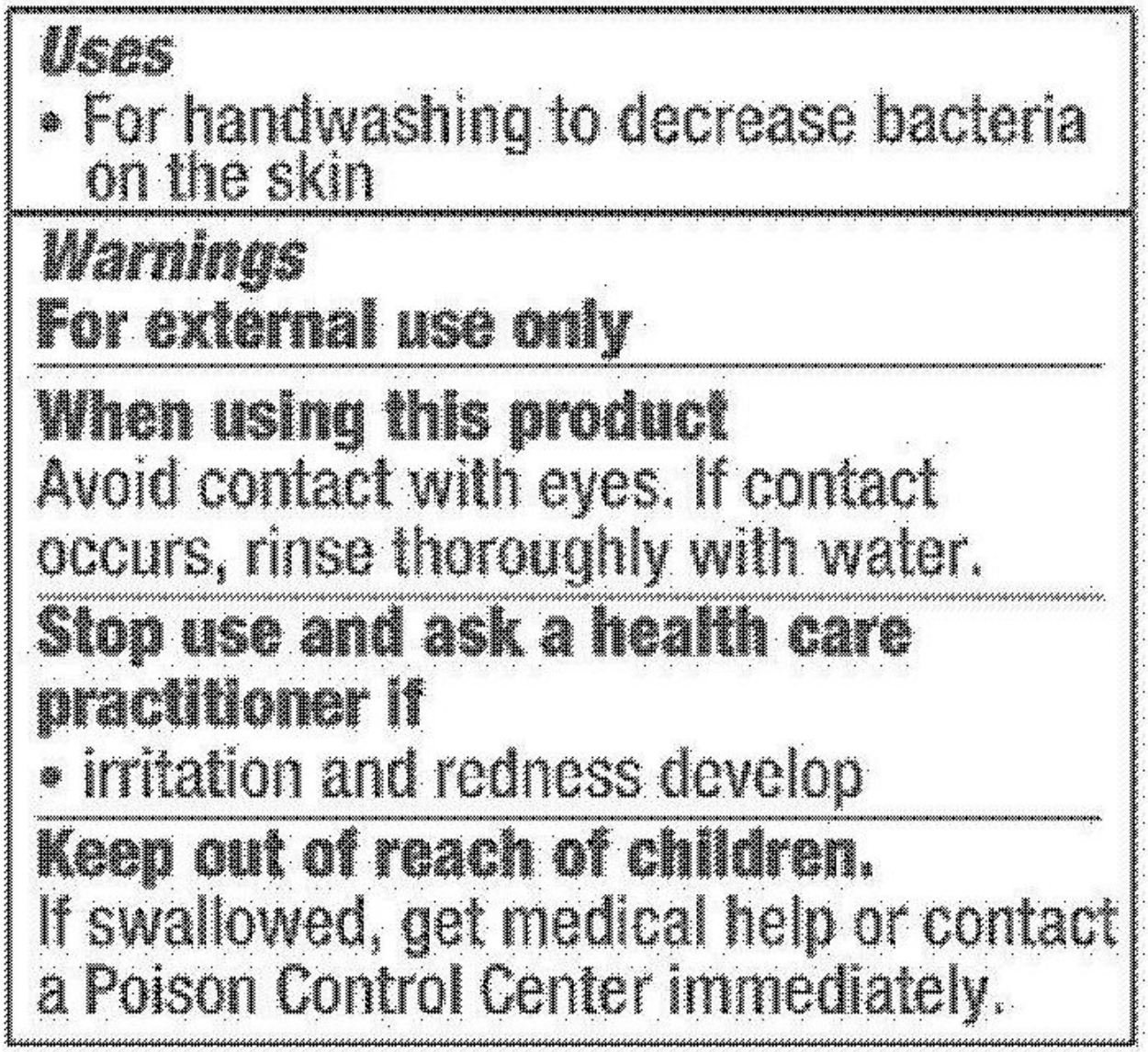
Active ingredient	***Purpose***
Ethyl Alcohol 64.76%	Antiseptic

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

2091699

PULL



NATURALS VANILLA ANTIBACTERIAL HAND

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10096-0251
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

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

1	NDC:10096-0251-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:10096-0251-2	75 mL in 1 TUBE; Type 0: Not a Combination Product		

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
Avon Products, Inc.		005149471	manufacture(10096-0251)

Revised: 2/2016 New Avon LLC