SOLGREAT ANTIBACTERIAL HANDWASH- benzalkonium chloride liquid PANATURAL USA, 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.

SolGreat Antibacterial handwash

Active Ingredient

Benzalkonium Chloride

Purpose

Antibacterial

Uses

• for handwashing to decrease bacteria on the skin.

Warnings

For external use only.

When using this product, avoid contact with eyes. In case of eye contact, flush with water.

Stop use and ask a doctor if irritation and redness develop.

Keep out of reach of children.

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

Directions

- place enough product into your palms, wet as needed.
- lather vigorously for at least 20 seconds.
- wash skin, rinse and dry thoroughly.

Inactive ingredients

Cetrimonium Chloride, Cocamide Methyl MEA, Lauramine Oxide, Methylchoroisothiazolinone, Methylisothiazolinone, Parfum (Fragrance), Propanediol, Sodium Chloride, Tocopheryl Acetate, FD&C Yellow No. 5, FD&C Red No. 4, Water.

Product Label

237mL SolGreat Antibacterial Hand Soap





Active ingredient Purpose Benzalkonium Chloride 0.13%......Antibacterial

Uses for handwashing to decrease bacteria on the skin.

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Directions ■ place enough product into your palms, wet as needed. ■ lather vigorously for at least 20 seconds. ■ wash skin, rinse and dry thoroughly.

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Distributed by: SOLGREAT LABS 3705 Pico Blvd., Los Angeles, CA 90019



Product Label - 500mL SolGreat Antibacterial Hand Soap



Drug Facts

Active ingredient Purpose
Benzalkonium Chloride 0.13%......Antibacterial

Uses for handwashing to decrease bacteria on the skin.

Warnings For external use only.

When using this product, avoid contact with eyes. In case of eye contact, flush with water.

Stop use and ask doctor if irritation and redness develop.

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Made in China

SOLGREAT ANTIBACTERIAL HANDWASH

benzalkonium chloride liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:73913-003 Route of Administration TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII: 7N6 JUD5 X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg

Inactive Ingredients	
Ingredient Name	Strength
Cetrimonium chloride (UNII: UC9PE95IBP)	
COCOYL METHYL MONOETHANOLAMINE (UNII: 79G1T427CF)	
LAURAMINE O XIDE (UNII: 4F6 FC4MI8 W)	
METHYLCHLORO ISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)	
PROPANEDIOL (UNII: 5965N8W85T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:73913-003-01	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/27/2020	
	2 NDC:73913-003-02	$500\ mL$ in $1\ BOTTLE;$ Type $0\colon Nota$ Combination Product	03/27/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/23/2020		

Labeler - PANATURAL USA, INC. (029572239)

Establishment					
Name	Address	ID/FEI	Business Operations		
Hefei Yatai Daily-Use Chemical Industry Co., Ltd		654641724	manufacture(73913-003)		

Revised: 3/2020 PANATURAL USA, INC.