SOLGREAT ALCOHOL-FREE FOAMING HAND SANITIZER- benzalkonium chloride spray 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 Alcohol-Free Hand Sanitizer

Active Ingredient

Benzalkonium Chloride

Purpose

Antibacterial

Uses

- hand sanitizer to help reduce bacteria on the skin that could cause disease.
- recommended for repeated use.

Warnings

For external use only.

When using this product, do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water,

Stop use and ask a doctor if redness or irritation develop and persist for more than 72 hours.

Directions

- place enough product in your palm to thoroughly cover your hands.
- rub hands together until dry.
- children under 6 years of age should be supervised when using this product.

Inactive Ingredients

cetrimonium chloride citric acid, cocamidopropyl betaine, cocamidopropylamine oxide, glycerin, propylene glycol, tocopheryl acetate, water.

Keep out of reach of children.

Children must be supervised in use of this product.

Other information

To prevent discoloration, avoid contact with clothing.

Product Label - 60 mL SolGreat Alcohol-Free Hand Sanitizer Spray





Drug Facts (continued)

When using this product, do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if redness or irritation develop and persist for more than 72 hours.

Keep out of reach of children. Children must be supervised in use of this product.

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Product Label- 354mL SolGreat Alcohol-Free Foaming Hand Sanitizer





benzalkonium chloride spray

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73913-002	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL		

Inactive Ingredients			
Ingredient Name	Strength		
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)			
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)			
COCAMIDO PRO PYL BETAINE (UNII: 50 CF30 11KX)			
CO CAMIDO PRO PYLAMINE O XIDE (UNII: M4SL82J7HK)			
GLYCERIN (UNII: PDC6A3C0OX)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
.ALPHATO COPHERO L ACETATE (UNII: 9E8X80D2L0)			
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73913-002-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/28/2020	
2	NDC:73913-002-02	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/28/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/28/2020		

Labeler - PANATURAL USA, INC. (029572239)

Establishment				
Name	Address	ID/FEI	Business Operations	
Hefei Yatai Daily-Use Chemical Industry Co., Ltd		654641724	manufacture(73913-002)	

Revised: 3/2020 PANATURAL USA, INC.