PURE SOURCE HAND SANITIZER- alcohol gel Pure Source, LLC

PURE SOURCE Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 70%

Purpose

Topical Antimicrobial

USES

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

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

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product

keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if

irritation or rash occurs. These may be signs of a serious condition

Keep out of reach of children.

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

Directions:

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing

Other information:

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aminomethylpropanediol, Aqua (Deionized Water), Ethylhexylglycerin, Glycerin, Phenoxyethanol, Polysorbate-20

Questions? (866) 565-575

Package Labeling: 65121-054-02



Package Labeling: 65121-054-16





Manufactured By: Pure Source, LLC

9750 NW 17th Street, Miami, FL 33172 Phone: (305) 477-8111 Email: info@thepuresource.com www.thepuresource.com

REFRESHING GEL

16 fl. oz (473ml)

Drug Facts

Active ingredient Ethyl Alcohol 70% v/v.

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Purpose

Antiseptic

Warnings For external use only. Flammable. Keep away from heat or flame.

Do not use

- in children less than 2 months of ageon open skin wounds

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Questions? (866) 565-5750

PURE SOURCE HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:65121-054

Route of Administration TOPICAL

Active Ingredient/Active Moiety

PHENOXYETHANOL (UNII: HIE492ZZ3T)

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) **ALCOHOL** 0.7 mL in 1 mL

Inactive Ingredients

1				
Ingredient Name	Strength			
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)				
WATER (UNII: 059QF0KO0R)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
GLYCERIN (UNII: PDC6A3C0OX)				

PC) I V	SORI	RATE	20	(HIMIII)	7T1F30V5YH)

F	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:65121-054- 02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020				
2	NDC:65121-054- 16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/10/2020				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	505G(a)(3)	04/01/2020			

Labeler - Pure Source, LLC (080354456)

Revised: 11/2023 Pure Source, LLC