

SAFE SOURCE HAND SANITIZER- alcohol gel
Cadence Keen Innovations, 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.

Safe Source Hand Sanitizer

Drug Facts

Active Ingredient:

Ethyl Alcohol 75%

Purpose

Antiseptic

Uses

Hand washing to reduce bacteria on skin, suitable when water is unavailable

Warnings

External Use only. Flammable, keep away from fire or flame.

Avoid contact with eyes, rinse thoroughly with water if contact occurs

Stop use if:

Skin irritation or redness develop or increase contact a doctor.

Keep out of reach of children.

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

Directions

- Spray product in your palm and rub together until dry.
- Children under 8 years of age should be used under supervision.

Inactive Ingredients

Water, Fragrance, Propanediol, Polysorbate 20, Glycerin

Store 59°-86°F. Avoid freezing heat above 104°F

CKI Solutions is USA distributor for O'ustane

Package Labeling:



Kills 99.9% of germs



www.CkiSolutions.us
1.69 fl oz (50ml)

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Use: Hand washing to reduce bacteria on skin, suitable when water is unavailable
Stop use if: Skin irritation or redness develop or increase contact a Doctor. Keep out of reach of children. If swallowed, get medical attention or contact a Poison Control Center right away.
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SAFE SOURCE HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79128-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.75 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPANEDIOL (UNII: 5965N8W85T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79128-001-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/22/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/22/2020	

