

**CARE PLUS HAND SANITIZER 16OZ- ethyl alcohol gel
OSTL, 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.

Drug Facts

Active ingredient

Ethyl Alcohol 62 percent

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product,

avoid contact with face, eyes and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask a doctor if irritation or redness develops.

Keep out of reach for children.

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

Directions

- wet hands thoroughly with product and rub into skin until dry.
- Children under 6 years of age should be supervised by an adult when using this product.

Inactive ingredients:

Water (Aqua), Glycerin, Propylene Glycol, Carbomer, Aminomethyl Propanol, Fragrance, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E)

Care Plus[®]

KILLS 99.9% OF GERMS*

Hand Sanitizer

ANTIBACTERIAL

with Moisturizers
& Vitamin E

16FL OZ (473ML)

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*Kills 99.9% of many common germs.
Imported by: O.S.T.L. Inc
Made in P.R.C.
www.ostltrade.com

CARE PLUS HAND SANITIZER 16OZ

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69950-026
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 934 (UNII: Z135WT9208)	
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging

Marketing Start Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69950-026-01	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/01/2020	
Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC monograph not final	part333A	03/01/2020	

Labeler - OSTL, INC. (020117798)

Registrant - OSTL, INC. (020117798)

Revised: 6/2020

OSTL, INC.