

**CARE PLUS HAND SANITIZER- 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.

Care Plus® Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 70.0% v/v

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 of 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, Aloe Barbadosis Leaf Juice, Carbomer, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E), Triethanolamine

KILLS GERMS*

ANTIBACTERIAL

with Moisturizers & Vitamin E

*Kills 99.9% of many common germs.

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Imported by **O.S.T.L.Inc**

Made in P.R.C.

Packaging

Care Plus®

KILLS GERMS*

Hand Sanitizer

ANTIBACTERIAL

**with Moisturizers
& Vitamin E**

16.9 FL OZ/500mL

DRUG FACTS LABEL

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CARE PLUS HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69950-028
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69950-028-02	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/17/2020	

Labeler - Ostl, Inc. (020117798)

Revised: 6/2020

Ostl, Inc.