FRESH HAND FOAMING HAND SANITIZER ALCOHOL FOAM- ethyl alcohol solution CWGC LA Inc.

CWGC (as PLD) - Resource Solutions 70% Foaming Hand Sanitizer Alcohol Foam (70415-203)

Active ingredient

Ethyl Alcohol 70 %

Purpose

Antibacterial

PURPOSE

ANTIBACTERIAL

Uses

Hand sanitizer to help reduce bacteria on the skin.

WARNINGS

Flammable. Keep away from fire or flame.

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 irritation or rash appears and lasts.

When using this product

avoid contact with eyes. In case of eye contact, flush with water

Keep out of reach of children.

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

Keep out of reach of children.

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

DIRECTIONS

- Place enough product on your palm to thoroughly cover your hands
- Rub hands together briskly until dry

Inactive ingredients

Water, Isopropyl Alcohol, Propylene Glycol, PEG-12 Dimethicone Crosspolymer, BIS-PFG-12 Dimethicone.



Drug Facts

Active Ingredient Purpose Ethyl alcohol, 70% Antibacterial

Use · Hand sanitizer to help reduce bacteria on the skin

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Directions:

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 • Rub hands together briskly until dry

Other Information

- Store below 110° (43°C)
- · May discolor certain fabrics or surfaces

Inactive Ingredients

Water, Isopropyl Alcohol, Propylene Glycol, PEG-12 Dimethicone Crosspolymer, Bis-PEG-12 Dimethicone.

Resource Solutions of SF Group, Inc. PO BOX 51, South San Francisco, CA 94083 Phone: (855) 742-6100 Fax: (855) 742-6105 www.rsgsf.com

Reorder Item #: RSGFHFS6 Emergency: Chemtel 800-255-3924



WARNING: FLAMMABLE LIQUID, SERIOUS EYE IRRITATION. HARMFUL IF SWALLOWED.

1000ml 34 fl oz

FOAMING HAND SANITIZER

User assumes all other risks and liabilities resulting from the use of this product.

FRESH HAND FOAMING HAND SANITIZER ALCOHOL FOAM

ethyl alcohol solution

Product Information

Product Type HUMAN OTC DRUG NDC:70415-203 Item Code (Source)

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Active ingredient/Active Plotety	Jiety .		
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)				
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)				
BIS-PEG-12 DIMETHICONE (70 MPA.S) (UNII: 2JDK5W22H4)				

Packaging	Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:70415- 203-11	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/04/2020		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/04/2020	

Labeler - CWGC LA Inc. (034967904)

Revised: 9/2023 CWGC LA Inc.