APLICARE ANTISEPTIC HAND RINSE- ethanol gel
Aplicare Products, LLC

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.

0540 Aplicare Antiseptic Hand Rinse

Active ingredient

Alcohol denat. (anhydrous) 62%

Purpose

Antiseptic

Use

instant hand antiseptic

Warnings

For external use only. Keep out of eyes. If contact with eyes occurs, rinse promptly and thoroughly with water.

Flammable. Keep away from fire or flame.

Stop use and ask a doctor if

significant irritation or sensitization ocurs

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.

Directions

- apply to clean, dry hands
- wet hands thoroughly with product and allow to dry

Other information

- not made with natural rubber latex
- recommended for repeated use
- reduces bacteria that can potentially cause disease

Inactive ingredients

acrylates/C10-30 alkyl acrylate crosspolymer, diisopropanolamine, FD&C blue no. 1, FD&C yellow no. 5, fragrance, PPG-20 methyl, glucose ether, water

Questions or comments?

1 800 633-5463

Manufacturing Information

Manufactured by:

Aplicare Products, LLC

550 Research Parkway, Meriden, CT 06450 USA

Made in USA

REF: APLL1540

RJ19APL

Package Label

◀ Tear Here Hold Upright Tear Here ▶

APLICARE®

NDC 52380-0540-1 REF APLL1540

ANTISEPTIC GEL HAND RINSE

Contains 62% Ethyl Alcohol

Light Herbal Fragrance

Net Contents 1.5 mL

Drug Facts

Active ingredient	*Purpose*
Alcohol denat. (anhydrous) 62%	Antiseptic

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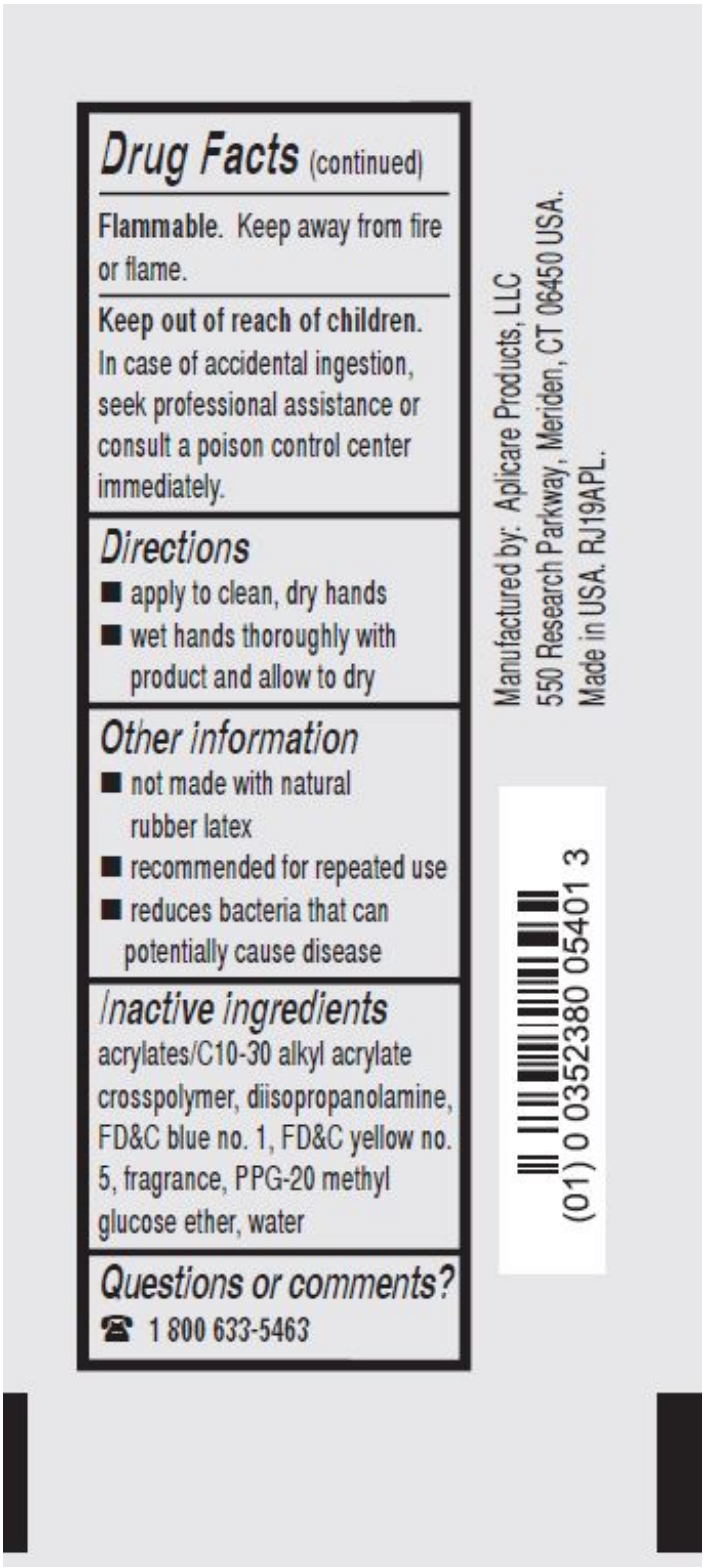

APLICARE ANTISEPTIC HAND RINSE

ethanol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52380-0540
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIISOPROPANOLAMINE (UNII: 0W44HYL8T5)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52380-0540-1	1.5 mL in 1 PACKET; Type 0: Not a Combination Product	02/14/2018	07/31/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2006	07/31/2025

Labeler - Aplicare Products, LLC (081054904)

Registrant - Medline Industries, LP (025460908)

Revised: 12/2022 Aplicare Products, LLC