APPEAL ALCOHOL FOAMING HAND SANITIZER- ethyl alcohol liquid SC Johnson Professional USA, 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 70% ww

Purpose

Antibacterial

Uses

For hand sanitizing to reduce bacteria on the skin

Warnings

For external use only

Flammable: Keep away from fire or flame.

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.

Directions

Apply one shot of foaming sanitizer to dry hands

Rub into skin

No rinsing required

Inactive ingredients

Water, Bis-PEG-12 Dimethicone, Behentrimonium Chloride, PEG-200 Hydrogenated Glyceryl Palmate, PEG-7 Glyceryl Cocoate, Coco-Glucoside, Glyceryl Oleate, Dihydroxypropyl PEG-5 Linoleammonium Chlroide

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1 Liter 33.8 Fluid Ounces

Made in Canada by Deb



Distributed exclusively by:

Interline Brands Jacksonville, FL 32207 www.AppealProducts.com

01830

Rev. 05-12

APP17102

102668

appeal Alcohol Foaming Hand Sanitizer

Certified EcoLogo

Certified Instant Hand Antiseptic CCD-170

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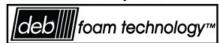
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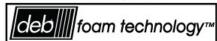
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ethyl alcohol liquid

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Product Type HUMAN OTC DRUG Item Code (Source)

Route of Administration TOPICAL NDC:11084-702

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)				
DIHYDROXYPROPYL PEG-5 LINO LEAMMONIUM CHLO RIDE (UNII: 0 Y0 NQR2GH1)				
BEHENTRIMO NIUM CHLO RIDE (UNII: X7GNG3S47T)				
DIMETHICO NE (UNII: 92RU3N3Y1O)				
COCO GLUCOSIDE (UNII: ICS790225B)				
PEG-200 DILAURATE (UNII: TWV5J70L88)				
GLYCERYL COCOATE (UNII: WVK1CT5994)				
GLYCERYL MONOOLEATE (UNII: C4YAD5F5G6)				

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:11084-702- 27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2012	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	05/15/2012		

Labeler - SC Johnson Professional USA, Inc. (607378015)

Establishment					
Name	Address	ID/FEI	Business Operations		
SC Johnson Professional CA Inc.		203765300	manufacture(11084-702)		

Revised: 12/2020 SC Johnson Professional USA, Inc.