DEB INSTANTFOAM COMPLETE- 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, 80% w/w

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 foaming sanitizer to cover hands

rub into skin

no rinsing required

Inactive ingredients

AQUA (WATER), BIS-PEG-12 DIMETHICONE, CITRIC ACID, COCO-GLUCOSIDE, DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE, GLYCERYL OLEATE, PANTHENOL, PEG-200 HYDROGENATED GLYCERYL PALMITATE, PEG-7 GLYCERYL COCOATE

deb stoko

NDC 11084-170-27

Deb InstantFOAM Complete

EN Broad Spectrum Alcohol-Based Instant Hand Sanitizer Foam with Moisturizer and Skin Conditioner

Perfume-free Dye-free No water required Kills 99.999% of many types of common germs

ES Amplio espectro antiseptico con alcohol para manos en espuma y con acondicionador

Sin perfume Sin colorante No se requiere agua Elimina el 99.999% de los muchos tipos de germenes mas comunes

SANITIZE

deb

Stock #:

IFC1L

Made in / Hecho en Canada

US Patents 5,445,288 & 6,082,586

Worldwide Patent Pending.

Made by Deb for:

Deb USA, Inc.

Charlotte, NC 28217

1-800-248-7190

www.debgroup.com

DCN9207/0715

1 L

(33.8 fl oz)

Meets food code hand sanitizer criteria (Section 2-301.16) published by the FDA

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DEB INSTANTFOAM COMPLETE

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-170
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.80 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BIS-PEG-12 DIMETHICONE (500 MPA.S) (UNII: 2CNS542YRT)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
COCO GLUCOSIDE (UNII: ICS790225B)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
PANTHENOL (UNII: WV9CM0O67Z)	
PEG-200 HYDROGENATED GLYCERYL PALMATE (UNII: W161T051Y1)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11084-170-01	47 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/18/2015	
2	NDC:11084-170-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/18/2015	
3	NDC:11084-170-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/18/2015	
4	NDC:11084-170-12	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/18/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/18/2015	

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

Registrant - SC Johnson Professional USA, Inc. (078805627)

Establishment

Name	Address	ID/FEI	Business Operations
SC Johnson Professional CA Inc.		203765300	manufacture(11084-170)

Revised: 4/2019

SC Johnson Professional USA, Inc.