ALCARE ENHANCED FOAMING HAND SANITIZER- 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.

Alcare® Enhanced Foaming Hand Sanitizer

Drug Facts

Active ingredient

ETHYL ALCOHOL, 72% v/v

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) Behentrimonium Chloride, BIS-PEG-12 Dimethicone, Coco-Glucoside, Dihydroxypropyl PEG-5 Linoleammonium Chloride, Glyceryl Oleate, PEG-200 Hydrogenerated Glyceryl Palmate, PEG-7 Glyceryl Cocoate, Propyl Alcohol.

PRINCIPAL DISPLAY PANEL - 1 Liter Bottle Label

SCJ PROFESSIONAL
A family company™

Alcare®
Enhanced

NDC 11084-026-27

Foaming Hand Sanitizer

Excellent Moisturization

15
seconds

Fast-acting

CHG Compatible

REORDER #
107281

Manufactured for:
SC Johnson Professional
Charlotte, NC 28217
1-866-783-0422
www.scjp.com
Pat. www.scjp.com/patents
Made in Canada

1 Liter (33.8 fl oz)
SAP # 4000004994
L-1433 R0

deb
SKIN CARE

Open for
Drug Facts

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Open for Drug Facts

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ECOLOGO

PRODUCTS CERTIFIED FOR REDUCED ENVIRONMENTAL IMPACT. VIEW SPECIFIC ATTRIBUTES EVALUATED: UL.COM/EL
UL 2783

ALCARE ENHANCED FOAMING HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-026
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
PROPYL ALCOHOL (UNII: 96F264O9SV)	
BIS-PEG-12 DIMETHICONE (500 MPA.S) (UNII: 2CNS542YRT)	
COCO GLUCOSIDE (UNII: ICS790225B)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
PEG-200 HYDROGENATED GLYCERYL PALMATE (UNII: W161T051Y1)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-026-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2019	
2	NDC:11084-026-12	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2019	
3	NDC:11084-026-40	400 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/01/2019	
4	NDC:11084-	1000 mL in 1 BOTTLE; Type 0: Not a Combination	02/17/2023	

4	026-66	Product	02/17/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	10/01/2019	

Labeler - SC Johnson Professional USA Inc. (607378015)

Establishment

Name	Address	ID/FEI	Business Operations
SC Johnson Professional CA Inc.		203765300	MANUFACTURE(11084-026)

Revised: 2/2023 SC Johnson Professional USA Inc.