ALCARE ELEVATE ANTISEPTIC HANDRUB- alcohol liquid
SC Johnson Professional USA, Inc.

Alcare® Elevate Antiseptic Handrub

Drug Facts

Active ingredient

Ethyl Alcohol, 70% 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 sanitizer to cover hands
- rub into skin
- no rinsing required

Inactive ingredients

Aqua (Water), Glycerin, Hydroxypropyl Cellulose, Panthenol, Parfum (Fragrance), Trisodium Dicarboxymethyl Alaninate.

PRINCIPAL DISPLAY PANEL - 1 Liter Bottle Label

Alcare®

Antiseptic Handrub
Elevate

SCJ PROFESSIONAL
HEALTHCARE

NDC 11084-034-27

Excellent Moisturization

Net Contents: 1 Liter (33.8 fl oz)
SAP # 4000009648

REORDER #
ALCELV100

deb
SKIN CARE

SCJ PROFESSIONAL HEALTHCARE

Alcare®

NDC 11084-034-27

Antiseptic Handrub

Elevate

Excellent Moisturization

Drug Facts

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Ethyl Alcohol, 70% v/v	Antibacterial

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Alcare® Antiseptic Handrub Elevate

Drug Facts (continued)

Uses ■ for hand sanitizing to reduce bacteria on the skin

Warnings
For external use only

ALCARE ELEVATE ANTISEPTIC HANDRUB

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-034
Route of Administration	TOPICAL		

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)	

GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
DEXPANTHENOL (UNII: 1O6C93RI7Z)	
TRISODIUM DICARBOXYMETHYL ALANINATE (UNII: 784K2O81WY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-034-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/15/2020	12/31/2025
2	NDC:11084-034-18	370 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/15/2020	12/31/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M003	10/15/2020	12/31/2025

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

Establishment

Name	Address	ID/FEI	Business Operations
SC Johnson Professional CA Inc.		203765300	MANUFACTURE(11084-034)

Revised: 12/2024 SC Johnson Professional USA, Inc.