ALCARE ELEVATE ANTISEPTIC HANDRUB- 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® 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

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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	
2	NDC:11084-034-18	370 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	10/15/2020	

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

Establishment

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

Revised: 9/2020 SC Johnson Professional USA, Inc.