

ALCARE 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® Liquid 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

SCJ PROFESSIONAL
HEALTHCARE

Alcare®

NDC 11084-035-27

Antiseptic Handrub Liquid

Excellent Moisturization

Net Contents: 1 Liter (33.8 fl oz)

SAP # 4000009648

REORDER #

ALCELV100

L-1448 RO

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SKIN CARE

SCJ PROFESSIONAL
HEALTHCARE

Alcare®

NDC 11084-035-27

Antiseptic Handrub

Liquid



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L-1449 RO

Manufactured for:
SC Johnson Professional USA, Inc. Charlotte, NC 28217
1-866-783-0422 www.scjp.com

ALCARE ANTISEPTIC HANDRUB

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11084-035
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62.94 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
PANTHENOL (UNII: WW9CM0O67Z)	
TRISODIUM DICARBOXYMETHYL ALANINATE (UNII: 784K2O81WY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-035-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2021	
2	NDC:11084-035-18	370 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	06/01/2021	

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

Establishment

Name	Address	ID/FEI	Business Operations
SC Johnson Professional CA Inc.		203765300	MANUFACTURE(11084-035)

Revised: 6/2021

SC Johnson Professional USA, Inc.