

**ALCARE EXTRA HAND SANITIZER- alcohol solution**  
**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.*

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**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

**PRINCIPAL DISPLAY PANEL - 1 Liter Bottle Label**

SCJ PROFESSIONAL  
HEALTHCARE

Alcare®

NDC 11084-006-27

Hand Sanitizer  
Foaming Antiseptic Handrub  
Extra

Excellent Moisturization

15  
seconds  
Fast-acting  
CHG Compatible

SC Johnson Professional USA, Inc.  
Charlotte, NC 28217  
1-866-783-0422  
Pat. [www.scjp.com/patents](http://www.scjp.com/patents)  
[www.scjp.com](http://www.scjp.com)  
Made in Canada

1 Liter (33.8 fl oz)  
SAP # 400000076  
L-1398 R0

REORDER #  
101561

deb  
SKIN CARE

Open for  
Drug Facts

# Alcare®

NDC 11084-006-27

## Hand Sanitizer

Foaming Antiseptic Handrub **Extra**



Excellent Moisturization



Fast-acting



CHG Compatible

### Drug Facts

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REORDER #  
**101561**



Open for  
Drug Facts



• Perfume-free



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

**ECOLOGO**

Product certified to EcoLogo UL 2783 Standard for reduced environmental impact which sets metrics for environmental and other criteria, including: materials, packaging, human health, environment, product performance and labeling. View specific attributes evaluated: [ul.com/el](http://ul.com/el)



Nonfood Compounds  
Program Listed E3  
155160



**Drug Facts (continued)**

**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 Hydrogenerated Glyceryl Palmate, PEG-7 Glyceryl Cocoate

**ALCARE EXTRA HAND SANITIZER**

alcohol solution

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11084-006
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BIS-PEG-12 DIMETHICONE (500 MPA.S) (UNII: 2CNS542YRT)	

<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>COCO GLUCOSIDE</b> (UNII: ICS790225B)	
<b>DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE</b> (UNII: 0Y0NQR2GH1)	
<b>GLYCERYL OLEATE</b> (UNII: 4PC054V79P)	
<b>PANTHENOL</b> (UNII: WW9CM0067Z)	
<b>PEG-200 HYDROGENATED GLYCERYL PALMATE</b> (UNII: W161T051Y1)	
<b>PEG-7 GLYCERYL COCOATE</b> (UNII: VNX7251543)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-006-01	47 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2017	
2	NDC:11084-006-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2017	
3	NDC:11084-006-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2017	
4	NDC:11084-006-12	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2017	
5	NDC:11084-006-66	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/17/2023	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/01/2017	

**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(11084-006)

Revised: 2/2023

SC Johnson Professional USA, Inc.