SOFT N SURE ANTISEPTIC HAND- alcohol gel 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.

Soft 'N Sure® Antis eptic Hand Gel

Drug Facts

Active ingredient

Ethyl alcohol 65%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable. Keep away from fire or flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation and redness develop.
- condition persists more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use under adult supervision
- not recommended for infants

Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive Ingredients

benzophenone-4, carbomer, fragrance, glycerin, isopropyl myristate, propylene glycol, tocopheryl acetate, water

Questions or comments?

1-866-783-0422

PRINCIPAL DISPLAY PANEL - 1 Liter Bottle Label

SCJ PROFESSIONAL HEALTHCARE

Soft 'N Sure®

NDC 11084-812-21

Hand Sanitizer

Antiseptic Hand Gel

Moisturizing

15

seconds

Fast-Acting

Manufactured for:

SC Johnson Professional USA, Inc.

Charlotte, NC 28217

1-866-783-0422

www.scjp.com

Pat. www.scjp.com/patents

deb

SKIN CARE

1 Liter (33.8 fl oz) (1.05 qt) SAP # 400000065

REORDER#

1445-87

1445-86H(F)(118)



HEALTHCARE

Soft 'N Sure®

NDC 11084-812-21

Hand Sanitizer

Antiseptic Hand Gel



Moisturising



Fast-Acting



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*Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds.

DSP-MO-34 L0000376BC SDA WI-2059 1445-86H(B)(118) SDS-MO-15036

61873

SOFT N SURE ANTISEPTIC HAND

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NI

NDC:11084-812

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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

Inactive Ingredients				
Ingredient Name	Strength			
SULISOBENZONE (UNII: 1W6L629B4K)				
Glycerin (UNII: PDC6A3C0OX)				
Isopropyl Myristate (UNII: 0 RE8 K4LNJS)				
Propylene Glycol (UNII: 6DC9Q167V3)				
ACETATE ION (UNII: 569 DQM74SC)				
Water (UNII: 059QF0KO0R)				

Packaging					
Item Code	Package Description	Marketing Start Date	Marketing End Date		
NDC:11084-812- 41	1000 mL in $1BOTTLE,$ PLASTIC; Type $0\colon Nota$ Combination Product	0 2/0 1/20 18			
NDC:11084-812- 13	444 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 2/0 1/20 18	03/31/2022		
NDC:11084-812- 21	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 2/0 1/20 18	03/31/2022		
NDC:11084-812- 25	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 2/0 1/20 18	0 1/3 1/2 0 2 2		
	NDC:11084-812- 41 NDC:11084-812- 13 NDC:11084-812- 21 NDC:11084-812-	Ttem Code Package Description NDC:11084-812- 41 NDC:11084-812- 13 NDC:11084-812- 13 Product NDC:11084-812- 21 Product Product Product Product NDC:11084-812- 21 Product Product Product Product Product Product Product Product Product NDC:11084-812- 21 Product Separation Product Product Product NDC:11084-812- 21 Product NDC:11084-812- 21 Product NDC:11084-812- 21 Product NDC:11084-812- 21 Product Product NDC:11084-812- 21	Item CodePackage DescriptionMarketing Start DateNDC:11084-812-411000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product02/01/2018NDC:11084-812-13444 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product02/01/2018NDC:11084-812-21946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product02/01/2018NDC:11084-812-59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination02/01/2018		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	0 2/0 1/20 18			

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

Establishment					
Name	Address	ID/FEI	Business Operations		
Vi-Jon		088520668	MANUFACTURE(11084-812)		

Revised: 12/2020 SC Johnson Professional USA, Inc.