INSTANT FOAM NON-ALCOHOL PURE HAND SANITIZER- benzalkonium chloride 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.

Instant Foam™ Non-alcohol PURE Hand Sanitizer

Drug Facts

Active ingredient

Benzalkonium Chloride, 0.13%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease

Warnings

For external use only

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 for 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 thouroughly with product and allow to dry without wiping.
- For children under 6, use only under adult supervision.

Inactive ingredients

Aqua (Water), Aloe Barbadensis Leaf Juice, Caprylyl Glycol, Cocamidopropyl Betaine,

Lauramine Oxide, Phenoxyethanol, Propylene Glycol, Tetrasodium EDTA

PRINCIPAL DISPLAY PANEL - 1 L Bottle Label

SCJ PROFESSIONAL
A Family Company™

NDC 11084-300-27

InstantFOAM™ Non-Alcohol PURE

HAND SANITIZER
Alcohol-Free FOAM Hand sanitizer - Dye & Perfume-Free

Stock #: 55857/4000007087

SC Johnson Professional USA, Inc.
Charlotte, NC 28217,
1-800-248-7190
www.scjp.com

1 L (33.8 fl oz)

deb
SKIN CARE

1000003036/0220

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INSTANT FOAM NON-ALCOHOL PURE HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-300
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE SODIUM (UNII: MP1J8420LU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-300-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2020	
2	NDC:11084-300-12	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2020	
3	NDC:11084-300-97	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2020	
4	NDC:11084-300-01	47 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2020	
5	NDC:11084-300-40	400 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2020	
6	NDC:11084-300-66	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	04/15/2015	

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

Establishment

Name	Address	ID/FEI	Business Operations
APEX International, Inc.		015226132	MANUFACTURE(11084-300)