INSTANT FOAM NON-ALCOHOL 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 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, Citric Acid, Cocamidopropyl Betaine, Lauramine Oxide, Parfum (Fragrance), Phenoxyethanol, Propylene Glycol, Tetrasodium EDTA, Blue 1 (CI 42090), Red 33 (CI 17200)

PRINCIPAL DISPLAY PANEL - 1000 mL Bottle Label

SCJ PROFESSIONAL
A Family Company™

NDC 11084-301-27

InstantFOAM™ Non-Alcohol

HAND SANITIZER
Alcohol-Free FOAM Hand Sanitizer - Fragranced

INSTANT FOAM NON-ALCOHOL HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-301
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)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE SODIUM (UNII: MP1J8420LU)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-301-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2020	
2	NDC:11084-301-97	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2020	

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
SC Johnson Professional USA, Inc.		078805627	MANUFACTURE(11084-301)

Revised: 3/2020 SC Johnson Professional USA, Inc.