FOAMING HAND SANITIZER- hand sanitizer aerosol, foam Sanitor Corporation

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.

Foaming Hand Sanitizer

Active Ingredient:

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Uses

for hand sanitizing to decrease bacteria on the skin recommended for repeated use

Warnings

For external use only

When using this product

avoid contact with eyes.

In case of contact, flush eyes with water.

Stop use and ask doctor

if irritation or redness develops, or if condition persists for than 72 hours.

Keep out of reach of children

if swallowed get medical help or contact a Poison Control Center right away.

Directions

pump enough foam to thoroughly cover your hand rub thoroughly over all surfaces of both hands rub hands together briskly until dry

Inactive Ingredients:

Citric Acid, Cocamidopropyl Betaine, Fragrance, Phenoxyethanol, Water

Foaming Hand Sanitizer Label

Kills germs and keeps you safe Alcohol-free, non-drying Non-irritating to the skin Rinse-free formula Made in USA

Alcohol-FREE

DRUG FACTS

Benzalkonium chloride 0,1% Active Ingredient

Purpose Antimicrobial

· for hand sanitizing to decrease bacteria on the skin

recommended for repeated use

Made in USA

Stop use and ask a doctor if irritation or redness develops, or

if condition persists for more than 72 hours.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

For external use only

Keep out of reach of children if swallowed get medical help

or contact a Poison Control Center right away

Directions





citric acid, cocamidoproyl betaine, fragrance, phenoxyethanol,

pump enough foam to thoroughly cover your hand
rub thoroughly over all surfaces of both hands

rub hands together briskly until dry

Inactive Ingredients





FOAMING HAND SANITIZER

hand sanitizer aerosol, foam

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:23261-100

Route of Administration TOPICAL

Active Ingredient/Active Moiety

•		
Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZ ALKONIUM	0.1 g
UNII:7N6IUD5X6Y)	CHLORIDE	in 100 a

Inactive Ingredients			
Ingredient Name	Strength		
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
FRAGRANCE ORANGE ORC2000765 (UNII: GX5JI7UXQ7)			
WATER (UNII: 059QF0KO0R)			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:23261- 100-01	3785 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	05/01/2020		

Labeler - Sanitor Corporation (797472792)

Registrant - Sanitor Corporation (797472792)

Establishment				
Name	Address	ID/FEI	Business Operations	
Sanitor Corporation		797472792	manufacture(23261-100)	

Revised: 12/2021 Sanitor Corporation