SINOFRESH FOAMING HAND SANITIZER- foaming hand sanitizer aerosol, foam
ZapVir, 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.

SinoFresh Hand Sanitizer

Warnings

Warnings

For External Use Only

When using this product

•Do not spray in eyes. If in eyes rinse with water

Stop Use and ask a Doctor

•If irritation or rash develops and lasts

Inactive Ingredients Water (Aqua), Carbamide

Active Ingredients Purpose

Benzalkonium Chloride 0.13%……..….Antiseptic

Keep out of reach of children

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

Uses • Hand Sanitizer to help reduce bacteria on the skin that could cause disease

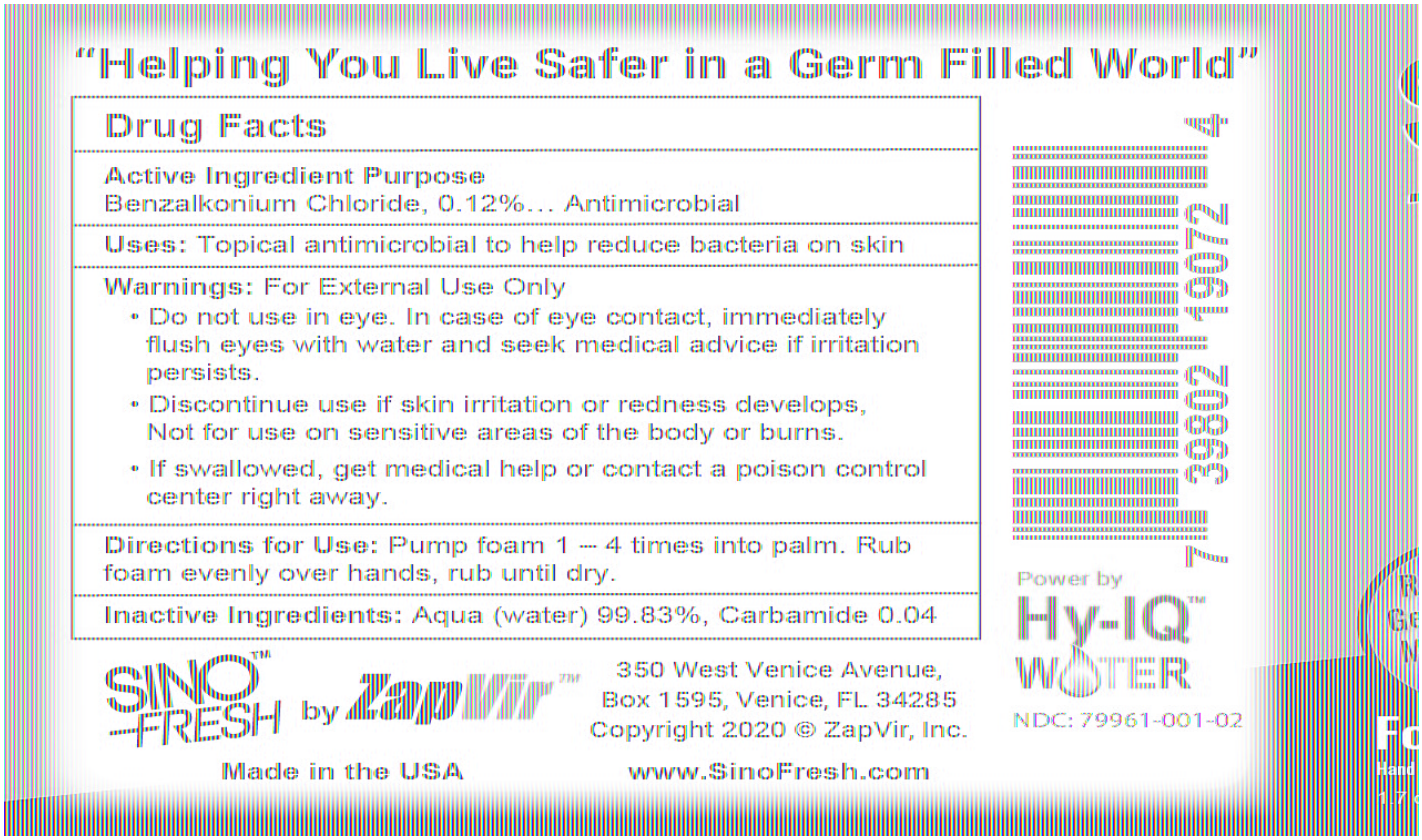
• Recommended for repeated use

Directions • Pump enough product on your palm to thoroughly cover your hands.

• Rub together until dry.

• Children under 6 years should be supervised when using this product.

Uses • Hand Sanitizer to help reduce bacteria on the skin that could cause disease



SINOFRESH FOAMING HAND SANITIZER

foaming hand sanitizer aerosol, foam

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79961-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.12 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
UREA (UNII: 8W8T17847W)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79961-001-01	30000 mg in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	08/17/2020	
2	NDC:79961-	50000 mg in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a	08/17/2020	

2	001-02	Combination Product	08/17/2020	
3	NDC:79961-001-03	207000 mg in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	08/17/2020	
4	NDC:79961-001-04	3800000 mg in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	08/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/17/2020	

Labeler - ZapVir, Inc. (117600850)

Registrant - ZapVir, Inc. (117600850)

Revised: 9/2020 ZapVir, Inc.