SAFMAXX HAND SANITIZER- benzalkonium chloride liquid
G&N Enterprise,LLC

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.

SAFMAXX Hand Sanitizer

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Uses

- handwash to help reduce bacteria that potentially can cause disease
- helps to prevent cross-contamination by hand contact
- helps to prevent drying of the skin
- recommended for repeated use

Warnings

For external use only

When using this product do not use in or near eyes

Stop use and ask a doctor if irritation or redness develop.

Keep out of reach of children.

If swallowed, seek medical attention or contact a Poison Control Center immediately.

Directions

- apply appropriate amount of product into palm of hand
- rub thouroughtly over all surfaces of both hands for at least 20 seconds
- rub hands together until dry

Other information

store in a cool, dry place below 104 ^{o} F (40 ^{o} C)

Inactive ingredients

Water, cocamidopropyl betaine, glycerine, polozamer 124, benzyl alcohol, phenoxyethanol,dehydroacetic acid, sodium hydroxide, citric acid,xanthan gum, FD&C Blue No.1

Questions?

Contact G&N Enterprise, LLC at **800-290-6816** Mon-Fri 9 am-5 pm CST

Kills 99.9% of germs

Protects up to 4 hours

SĀFMAXX™

HAND SANITIZER FOAM

Microbiological Shield

Benzalkonium Chloride 0.13%

Made in the U.S.A.

250 ml (8.45 fl oz)

SAFMAXX

Hand Sanitizer

Non_Alcohol

Microbiological Shield

Made in the USA

Benzalkonium Chloride 0.13%

SAFMAXX HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69731-041
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLOXAMER 124 (UNII: 1S66E28KXA)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69731-041-21	946352 mL in 1 TANK; Type 0: Not a Combination Product	05/01/2020	
2	NDC:69731-041-22	208197 mL in 1 DRUM; Type 0: Not a Combination Product	05/01/2020	
3	NDC:69731-041-23	3785 mL in 1 JUG; Type 0: Not a Combination Product	05/01/2020	
4	NDC:69731-041-25	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
5	NDC:69731-041-28	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
6	NDC:69731-041-20	45 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
7	NDC:69731-041-29	20 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
8	NDC:69731-041-27	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
9	NDC:69731-041-24	60 mL in 1 BOTTLE; 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	part333E	05/01/2020	

Labeler - G&N Enterprise,LLC (079758164)

Registrant - G&N Enterprise,LLC (079758164)

Revised: 5/2020 G&N Enterprise,LLC