HAND SANITIZER FOAM- anti-bacterial advanced formula liquid
VMP Cosmetics, 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.

Purpose

Antimicrobial

Uses

*To help decrease bacterial on the skin. Recomended for repeated use as needed.

Warnings

For external use only.

Stop Use

*Stop use and consult with doctor if irritation and redness develops, or if condition persists for more than 24 hours.

Keep out of reach of children

In case of accidental ingestion, seek professional assistance or contact a poison Control Center immediately.

Directions

* Pump a sufficient amount of foam into palm of hand.

* Rub thoroughly over all surfaces of both hands.

* Rub hands repeatedly until dry.

Other Information

Do not store above 104 ^{O}F.

Inactive Ingredients

Water, Glycerin, Hyaluronic Acid, Aloe Barbadensis Leaf Juice Powder, Quillaja Saponaria (Soapbark) Bark Extract, PEG-8 Dimethicone, Polyquaternium-7, PPG-26 Buteth-26, PEG-40 Hydrogenated Castor Oil, Sodium Decylglucosides Hydroxypropylsulfonate,

Ethylhexylglycerin, Phenoxyethanol, Fragrance.

Active Ingredient

Benzalkonium Chloride 0.13%

PDP

PANTONE 2349 C

PANTONE 2184 C

PANTONE 2747 C

Label width size reduced .125”

RECTANGLE 3.5 X 5.625”

MEDILAB

365

HAND SANITIZER FOAM

ANTI-BACTERIAL

ADVANCED FORMULA

KILLS 99.99%

OF GERMS

REGISTERED CERTIFIED

FDA GMP

FORMULA FACILITY

5 FL.OZ (150 ML)

Label width size reduced .125″

HAND SANITIZER FOAM

anti-bacterial advanced formula liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72471-100
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 g

Inactive Ingredients	
Ingredient Name	**Strength**
QUILLAJA SAPONARIA BARK (UNII: 8N0K3807ZW)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600 KD) (UNII: 0L414VCS5Y)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM DECYLGLUCOSIDES HYDROXYPROPYLSULFONATE (UNII: 55P9UXJ9C3)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
WATER (UNII: 059QF0KO0R)	
HYALURONIC ACID (UNII: S270N0TRQY)	
PPG-26-BUTETH-26 (UNII: 2II1K6TZ4P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72471-100-20	1 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/2020	
2	NDC:72471-100-21	1 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/18/2020	
3	NDC:72471-100-22	1 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/20/2020	

Labeler - VMP Cosmetics, LLC (022451337)

Registrant - VMP Cosmetics, LLC (022451337)

Establishment

Name	Address	ID/FEI	Business Operations
VMP Cosmetics, LLC		022451337	manufacture(72471-100)

Revised: 1/2022 VMP Cosmetics, LLC