RAW SPIRIT - MOISTURIZING HAND SANITIZER FOAM- benzalkonium chloride liquid
Cosmetic Solutions 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.

Raw Spirit - Moisturizing Hand Sanitizer Foam

DRUG FACTS:

ACTIVE INGREDIENTS

Benzalkonium chloride 0.13%

Purpose

Antiseptic

USES

To decrease bacteria on the skin.

WARNINGS

For external use only: Hands only.

- **When using this product** avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. Avoid contact with broken skin.
- **Stop use and ask a doctor if** irritation or redness develops - 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 thoroughly with product and allow to dry without wiping - for children under 6, use only under adult supervision - not recommended for infants

INACTIVE INGREDIENTS

Water (Aqua), Propanediol, Glycereth-7 Triacetate, Glycerin, Babassu Oil Glycereth-8 Esters, Decyl Glucoside, Calendula Officinalis Flower Extract, Equisetum Arvense Extract, Geranium Maculatum Extract, Honey, Citric Acid, Panax Ginseng Root Extract, Salvia Officinalis Leaf Extract, Sambucus Nigra Flower Extract, Ethylhexylglycerin, Phenoxyethanol, Sodium Hydroxide.

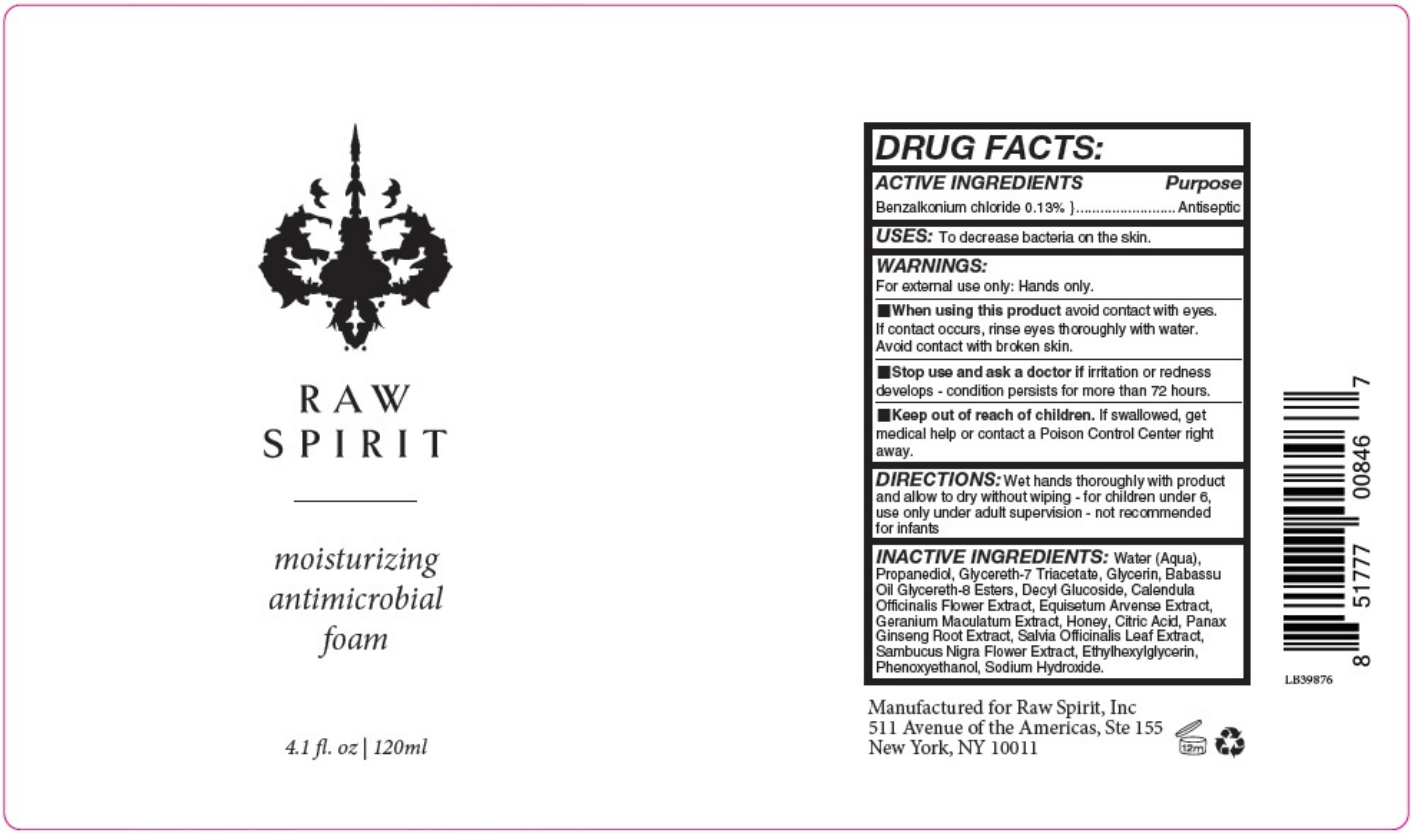
PRINCIPAL DISPLAY PANEL - 120 ml Bottle Label

RAW
SPIRIT

moisturizing
antimicrobial

foam

4.1 fl. oz | 120ml



RAW SPIRIT - MOISTURIZING HAND SANITIZER FOAM

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66163-1901
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 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Propanediol (UNII: 5965N8W85T)	
Glycereth-7 Triacetate (UNII: S9W6Z48HUP)	
Glycerin (UNII: PDC6A3C0OX)	
Babassu Oil Glycereth-8 Esters (UNII: 12XQ76K0YL)	
Decyl Glucoside (UNII: Z17H97EA6Y)	

Calendula Officinalis Flower (UNII: P0M7O4Y7YD)	
Equisetum Arvense Branch (UNII: 1L0VKZ185E)	
Geranium Maculatum Root (UNII: 93IXI5B6OJ)	
Honey (UNII: Y9H1V576FH)	
Citric Acid Monohydrate (UNII: 2968PHW8QP)	
Asian Ginseng (UNII: CUQ3A77YXI)	
Sambucus Nigra Flower (UNII: 07V4DX094T)	
Ethylhexylglycerin (UNII: 147D247K3P)	
Phenoxyethanol (UNII: HIE492ZZ3T)	
Sodium Hydroxide (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66163-1901-1	120 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	06/01/2020	

Labeler - Cosmetic Solutions LLC (807907928)

Revised: 7/2020 Cosmetic Solutions LLC