FOAMING HAND SANITIZER- ethyl alcohol 62% liquid
RJ Schinner

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.

RJ Schinner F688F

Dosage & Administration Section

Directions Apply a small amount to palm. Briskly rub, covering hands with product until dry.

Inactive Ingredient Section

Inactive Ingredients Water, Aloe Barbadensis Leaf Juice, Disodium EDTA, Glycerin, Polyquaternium-11, Isopropyl Myristate, Tocopheryl Acetate, PEG-10 Dimethicone

Indication & Usage Section

Uses Hand sanitizer to help reduce bacteria on the skin that could cause disease. Recommended for repeated use.

OTC-Keep Out of Reach of Children

Keep out of reach of children. If swallowed, contact a physician or poison control center.

Purpose Section

Purpose Antibacterial Agent

OTC - Active Ingredient Section

Active Ingredient Ethyl Alcohol 62% v/v

Warnings Section

Warnings

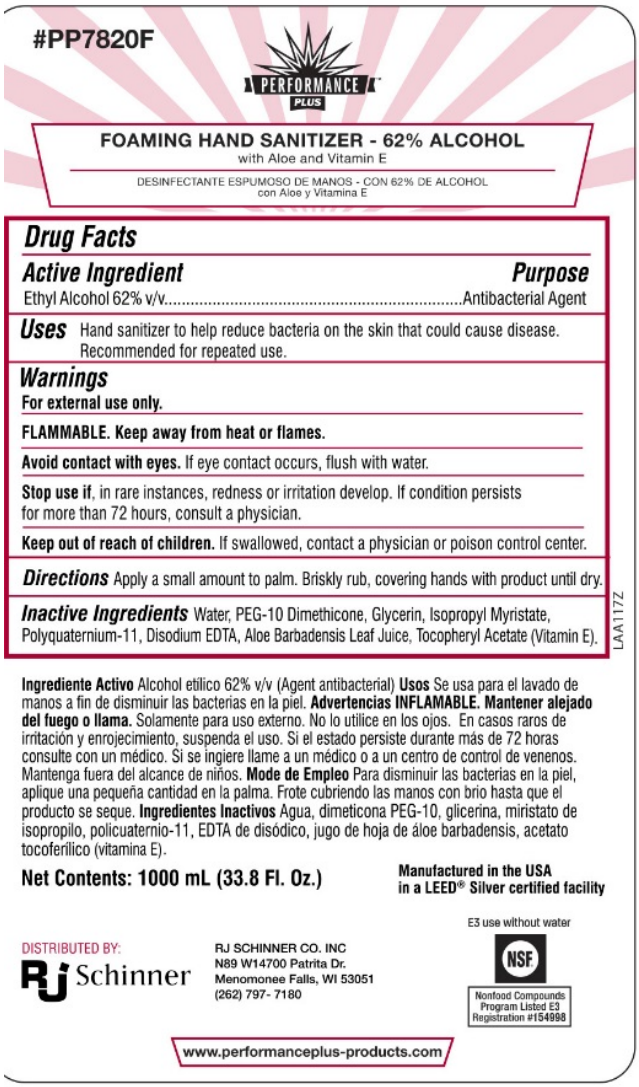
For external use only.

Avoid contact with eyes. If eye contact occurs, flush with water.

Stop use if, in rare instances, redness or irritation develop. If condition persists for more than 72 hours, consult a physician.

Principal Display Panel

PP7820F f688F



FOAMING HAND SANITIZER

ethyl alcohol 62% liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71303-688
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYQUATERNIUM-11 (1000000 MW) (UNII: 0B44BS5IJS)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
WATER (UNII: 059QF0KO0R)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71303-688-31	1000 mL in 1 BAG; Type 0: Not a Combination Product	03/30/2017	
2	NDC:71303-688-41	1000 mL in 1 BAG; Type 0: Not a Combination Product	03/30/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/30/2017	

Labeler - RJ Schinner (023432909)

Registrant - Kutol Products, Inc. (004236139)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products, Inc.		004236139	manufacture(71303-688)

Revised: 8/2023 RJ Schinner