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.

RI 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

#PP7820F



FOAMING HAND SANITIZER - 62% ALCOHOL

with Aloe and Vitamin E

DESINFECTANTE ESPUMOSO DE MANOS - CON 62% DE ALCOHOL

Drug Facts

Active Ingredient

Purpose

Ethyl Alcohol 62% v/v.....

Antibacterial Agent

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

Warnings

For external use only.

FLAMMABLE. Keep away from heat or flames.

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.

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

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

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

Ingrediente Activo Alcohol etilico 62% v/v (Agent antibacterial) Usos Se usa para el lavado de manos a fin de disminuir las bacterias en la piel. Advertencias INFLAMABLE. Mantener alejado del fuego o Idama. Solamente para uso externo. No lo utilice en los ojos. En casos raros de irritación y enrojecimiento, suspenda el uso. Si el estado persiste durante más de 72 horas consulte con un médico. Si se ingiere llame a un médico o a un centro de control de venenos. Mantenga fuera del alcance de niños. Mode de Empleo Para disminuir las bacterias en la piel, aplique una pequeña cantidad en la palma. Frote cubriendo las manos con brio hasta que el producto se seque. Ingredientes Inactivos Agua, dimeticona PEG-10, glicerina, miristato de isopropilo, policuaternio-11, EDTA de disódico, jugo de hoja de áloe barbadensis, acetato tocoferilico (vitamina E).

Net Contents: 1000 mL (33.8 Fl. Oz.)

Manufactured in the USA in a LEED® Silver certified facility

E3 use without water



RJ SCHINNER CO. INC N89 W14700 Patrita Dr. Menomonee Falls, WI 53051 (262) 797- 7180



www.performanceplus-products.com

FOAMING HAND SANITIZER

ethyl alcohol 62% liquid

D	Inform	
Produci	INTORM	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71303-688

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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

Inactive	Ingredients
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Ingredient Name Strength

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

l	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			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
part333E	03/30/2017		
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date	

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