

FOAMING SANITIZING HAND- alcohol soap
Kutol Products Company

F710F

☐ **Active Ingredient**

Ethyl Alcohol 72% v/v

☐ **Purpose**

Antibacterial

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

☐ **Uses** Hand sanitizer to help reduce bacteria on the skin that could cause disease.

☐ **Directions** To decrease bacteria on skin, apply small amount to palm. Briskly rub, covering hands with product until dry.

☐ **Warnings**

For external use only.

FLAMMABLE. Keep away from heat or flames.

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

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

☐ **Inactive Ingredients**

water, cocamidopropyl PG-dimonium chloride phosphate, cocamidopropyl betaine, PEG-6 cocamide, laurtrimonium chloride, iodopropynyl butylcarbamate, methylisothiazolinone.



*Foaming 72% Alcohol
Hand Sanitizer*

*Assainissant moussant à
base d'alcool 72% pour les mains*

*Higienizador Espumoso con
Alcohol 72% para las Manos*

E3 use without water

Drug Facts

Active Ingredient

Ethyl Alcohol 72% v/v.....Antibacterial Agent

Purpose

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

Directions To decrease bacteria on skin, apply 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.

Ingédient actif Alcool éthylique 72% v/v (Agent antibactérien) **Utilisations** Pour le lavage des mains, afin de réduire les bactéries sur la peau. **Avertissements INFLAMMABLE. Ne pas mettre a proximite d'un feu ou d'une flamme.** Pour usage externe seulement. Ne pas mettre dans les yeux. En cas d'irritation ou de rougeur, arrêtez l'utilisation. Si les symptômes persistent plus 72 heures, consultez un médecin. En cas d'ingestion, consultez un médecin ou un centre antipoison. Garder hors de portée des enfants. **Mode d'emploi** Pour réduire les bactéries sur la peau, appliquer une petite quantité sur la paume. Appliquer et faire pénétrer dans la peau en frottant.

Ingédients inactifs Eau, diméthicone PEG-10, glycérine, myristate isopropylique, polyquaternium-11, EDTA de disodique, jus de feuille d'aloès barbadine, acétate tocopherylique.

Ingrediente Activo Alcohol etílico 72% 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 llama.** 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 tocoferílico.

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

Manufactured in the USA
in a LEED® Silver certified facility

MANUFACTURED BY:
KUTOL PRODUCTS COMPANY
CINCINNATI, OH 45241
1-800-543-4641 www.kutol.com

FOAMING SANITIZING HAND

alcohol soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50865-710
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.072 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
POLYQUATERNIUM-11 (100000 MW) (UNII: 0B44BS5JJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50865-710-41	1000 mL in 1 BAG; Type 0: Not a Combination Product	12/01/2015	
2	NDC:50865-710-17	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2015	
3	NDC:50865-710-44	1000 mL in 1 BAG; Type 0: Not a Combination Product	10/01/2015	
4	NDC:50865-710-55	1200 mL in 1 BAG; Type 0: Not a Combination Product	10/01/2015	
5	NDC:50865-710-91	532 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/01/2015	

Labeler - Kutol Products Company (004236139)**Registrant** - Kutol Products Company (004236139)**Establishment**

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(50865-710)