

HAND SANITIZER- foaming alcohol hand sanitizer solution
State Industrial Product

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.

State Industrial F688F

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

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

Active Ingredient Ethyl Alcohol 62% v/v

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

Purpose Antibacterial Agent

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



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.

3 x 5 Foaming bag 1000mL label

FOAMING ALCOHOL HAND SANITIZER
ASSAINISSANT MOUSSANT À BASE D'ALCOOL POUR LES MAINS NPN 80014944

Product #121456

www.statechemical.com

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.	
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Ingrédient actif Alcool éthylique 62% v/v (Agent antibactérien) **Utilisations** Pour le lavage des mains, afin de réduire les bactéries sur la peau. **Avertissements** INFLAMMABLE. Ne pas mettre à proximité 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 de 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. **Ingrédients inactifs** Eau, diméthicone PEG-10, glycérine, myristate isopropylique, polyquaternium-11, EDTA de disodique, jus de feuille d'aloès barbadine, acétate tocophérylique.

Seller shall not be liable for consequential, special, or indirect damages from use of their product and liability is expressly limited to the refund of purchase price or replacement of product.

State Industrial Products
5915 Landerbrook Drive
Mayfield Heights, OH 44124
To Order Call: 1-866-747-2229
www.statechemical.com

Canada
6935 Davand Drive
Mississauga, Ontario L5T 1L5
To Order Call: 1-800-668-8513

Puerto Rico
Royal Industrial Park, Bldg "M"
Local #5, Carr 869, km 1.5 Palmas
Cataño, PR. 00962
To Order Call: 787-275-3185

61N03 **Net Contents/Contenu Net: 1000mL (33.8 fl.oz.)**

Label#121456LBL MADE IN USA

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Principal Label

HAND SANITIZER

foaming alcohol hand sanitizer solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64542-688
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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
POLYQUATERNIUM-11 (1000000 MW) (UNII: 0B44BS5IJS)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64542-688-41	1000 mL in 1 BAG; Type 0: Not a Combination Product	11/21/2017	
2	NDC:64542-688-17	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/21/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/21/2017	

Labeler - State Industrial Product (004520987)**Registrant** - Kutol Products Company (004236139)**Establishment**

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(64542-688)