FOAMING INSTANT HAND SANITIZER- benzalkonium chloride solution
Global Equipment Company

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.

Global Equipment F207

Active Ingredient Benzalkonium Chloride 0.13% w/w

Purpose Antibacterial Agent

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

Warnings

For external use only. Avoid contact with eyes. If 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.

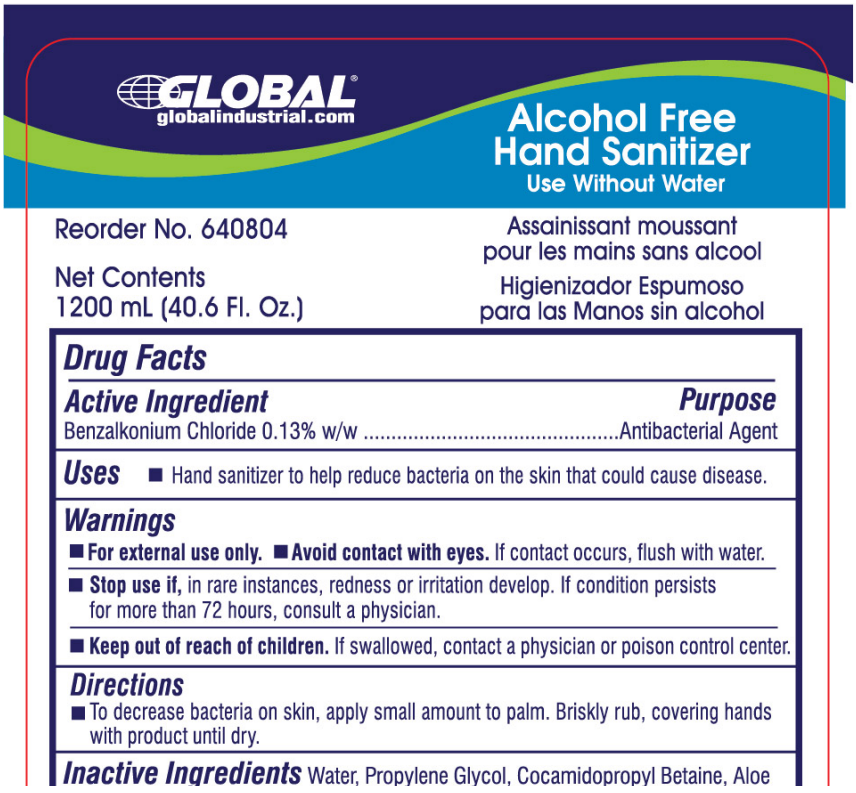
Directions

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

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

Inactive Ingredients

Water, Propylene Glycol, Cocamidopropyl Betaine, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate, PEG-7 Glyceryl Cocoate, Fragrance, Phenoxyethanol, Tetrasodium EDTA.



Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E), PEG-7 Glyceryl Cocoate, Fragrance, Phenoxyethanol, Tetrasodium EDTA.

Ingrédient actif Chlorure de benzalkonium 0,13% p/p (Agent antibactérien) **Utilisations** Pour le lavage des mains, afin de réduire les bactéries sur la peau. **Avertissements** Pour usage externe seulement. Ne pas mettre dans les yeux. En cas de contact avec les yeux, rincer avec de l'eau. En cas rares d'irritation, 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, propylèneglycol, bétaine cocamidopropylique, jus de feuille d'aloès barbadine, acétate tocophérylique (vitamine E), PEG-7 cocoate glycérylique, parfum, phénoxyéthanol, tétrasodium d'EDTA.

Ingrediente Activo Cloruro de benzalkonium 0.13% en peso (Agente antibacteriano) **Usos** Se usa para el lavado de manos a fin de disminuir las bacterias en la piel. **Advertencias** Solamente para uso externo. No lo utilice en los ojos. Si entra en contacto con los ojos, enjuáguelos con agua. En casos raros de irritación, 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, propilén glicol, betaina cocamidopropílica, jugo de hoja de aloe barbadensis, acetato tocoferílico (vitamina E), PEG-7 cocoato gliceril, perfume, fenoxietanol, tetrasodio de EDTA.

LAA120W

Made in the USA

Manufactured for:
Global Industrial / 2505 Mill Center Parkway / Buford, GA 30518
For Reorders: 800-645-1232 / globalindustrial.com

Emergency:
Chemtrec 800-424-9300

KUTOL
Hand Hygiene Specialists

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513-527-5506 fax
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KUTOL INTERNAL
Approved_____
Date_____
Changes:

Label No. LAA120W
Size 3 x 5 Unwind_____
Colors PMS 2755 purple, 375 green, Process blue
Printed on_____
Formula 207
Item No. (Kutol use only) 640804

Printer - please put our formula number on your proof.

Revisions: 3/9/16 - Change formula from 682F to 207, assign new LA number (Old: LA710U).
7/25/19 - Changing from 1250 mL to 1200 mL, (OLD LAA113Q)

Package Label

FOAMING INSTANT HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78641-207
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.013 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE SODIUM (UNII: MP1J8420LU)	
PROPYLENE GLYCOL 1,2-DISTEARATE (UNII: T65PN3O37H)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78641-207-78	950 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/16/2020	
2	NDC:78641-207-50	1200 mL in 1 BAG; Type 0: Not a Combination Product	03/16/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/16/2020	

Labeler - Global Equipment Company (001472216)

Registrant - Kutol Products Company (004236139)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(78641-207)

Revised: 8/2023 Global Equipment Company