FOAMING HAND SANITIZER- indulgence hand sanitizer solution
US Chemical Corporation

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.

US Chem F688F

Directions Apply a small amount to palm. Briskly rub, covering hands with product until dry. Children under 6 years of age should be supervised when using this product.

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

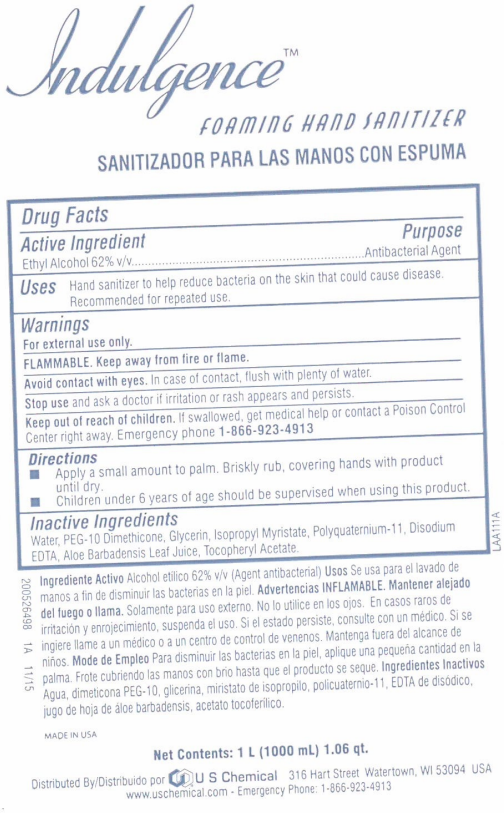
Uses Hand sanitizer to help reduce bacteria on the skin that could cause diease. Recommended for repeated use.

Active Ingredient Ethyl Alcohol 62% v/v

Keep out of reach of children. If swallowed, get medical help, or contact a Poison Control Center right away.

Purpose Antibacterial Agent

Warnings For external use only. Flammable. Keep away from fire or flame. Avoid contact with eyes. In case of contact, flush with plenty of water. Stop use and ask a doctor if irritation or rash appears and persists.



FOAMING HAND SANITIZER

indulgence hand sanitizer solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:61307-688
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)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
POLYQUATERNIUM-11 (1000000 MW) (UNII: 0B44BS5IJS)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61307-688-10	1000 mL in 1 BAG; Type 0: Not a Combination Product	07/03/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/03/2012	

Labeler - US Chemical Corporation (031457842)

Registrant - Kutol Products Company (004236139)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(61307-688)