

**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.*

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**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 disease. 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.

# Indulgence™

FOAMING HAND SANITIZER

SANITIZADOR PARA LAS MANOS CON ESPUMA

Drug Facts	
Active Ingredient	Purpose
Ethyl Alcohol 62% v/v.....	Antibacterial Agent
<b>Uses</b> Hand sanitizer to help reduce bacteria on the skin that could cause disease. Recommended for repeated use.	
<b>Warnings</b>	
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.	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Emergency phone 1-866-923-4913	
<b>Directions</b>	
<ul style="list-style-type: none"> <li>■ Apply a small amount to palm. Briskly rub, covering hands with product until dry.</li> <li>■ Children under 6 years of age should be supervised when using this product.</li> </ul>	
<b>Inactive Ingredients</b>	
Water, PEG-10 Dimethicone, Glycerin, Isopropyl Myristate, Polyquaternium-11, Disodium EDTA, Aloe Barbardensis Leaf Juice, Tocopheryl Acetate.	

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**Ingrediente Activo** Alcohol etílico 62% v/v (Agente antibacteriano) **Usos** Se usa para el lavado de manos a fin de disminuir las bacterias en la piel. **Advertencias INFLAMABLE. Mantener alejado del fuego o flama.** Solamente para uso externo. No lo utilice en los ojos. En casos raros de irritación y enrojecimiento, suspenda el uso. Si el estado persiste, 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. **Modo de Empleo** Para disminuir las bacterias en la piel, aplique una pequeña cantidad en la palma. Frote cubriendo las manos con brío hasta que el producto se seque. **Ingredientes Inactivos** Agua, dimeticona PEG-10, glicerina, miristato de isopropilo, poliquaternio-11, EDTA de disódico, jugo de hoja de aloe barbadensis, acetato tocoferílico.

MADE IN USA

Net Contents: 1 L (1000 mL) 1.06 qt.

Distributed By/Distribuido por **U S Chemical** 316 Hart Street Watertown, WI 53094 USA  
www.uschemical.com - Emergency Phone: 1-866-923-4913

indulgence hand sanitizer solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:61307-688
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
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

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

### Marketing Information

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

**Labeler** - US Chemical Corporation (031457842)

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

### Establishment

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

