INDULGENCE/PROPOWER- benzalkonium chloride soap
U S 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 F213F

Benzalkonium Chloride 0.13% w/w

Antibacterial Agent

For handwashing to decrease bacteria on skin.

For external use only.

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 TELEPHONE: 1-866-923-4913**

To decrease bacteria on skin, apply small amount, covering hands with product for 30 seconds. Add water, lather, rinse.

Water, cocamidopropyl betaine, cocamidopropyl PG-dimonium chloride phosphate, PEG-6 cocamide, laurtrimonium chloride, aloe barbadensis leaf juice, tocopheryl acetate (vitamin E), fragrance, iodopropynyl butylcarbamate, methylisothiazolinone, yellow 5, red 33.

ProPower™

FOAMING ANTIMICROBIAL HAND SOAP

Drug Facts

Active Ingredient — **Purpose**

Benzalkonium Chloride 0.13% w/w ..... Antibacterial Agent

Uses ■ For handwashing to decrease bacteria on skin.

Warnings

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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**

Directions

■ To decrease bacteria on skin, apply small amount, covering hands with product for 30 seconds. Add water, lather, rinse.

Inactive Ingredients Water, Cocamidopropyl Betaine, Cocamidopropyl PG-Dimonium Chloride Phosphate, PEG-6 Cocamide, Laurtrimonium Chloride, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E), Fragrance, Iodopropynyl Butylcarbamate, Methylisothiazolinone, Yellow 5, Red 33.

LAA112P

Ingrediente Activo Cloruro de benzalconio 0.13 % en peso (Agentes 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. 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** Aplique una cantidad pequeña de producto sobre las manos, cubriendo las manos durante 30 secundos. Añada agua, forme espuma y enjuague. **Ingredientes Inactivos** Agua,cocamidopropil betaina, cloruro y fosfato de cocamidopropil PG-dimonio, PEG-6 cocamida, cloruro de laurtrimonio, jugo de hoja de aloe barbadensis, acetato tocoferílico (vitamina E), fragancia, yodopropinil butilcarbamato, metilisotiazolinona, amarillo 5, rojo 33.

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Marketed By / Comercializado Por
Independent Marketing Alliance
16000 Memorial Drive, Suite 200
Houston, TX 77079 USA
Emergency Phone 1-866-923-4913

Product Of USA / Producto De EE.UU.

Net contents: 1 L (1000 mL) 1.06 qt

INDULGENCE/PROPOWER

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:61307-213
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	2.5 mg in 1 L

Inactive Ingredients

Ingredient Name	Strength
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
PEG-6 COCAMIDE (UNII: YZ6NLA4O1E)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61307-213-41	1 L in 1 BAG; Type 0: Not a Combination Product	01/30/2017	
2	NDC:61307-213-10	1 L in 1 BAG; Type 0: Not a Combination Product	01/30/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/30/2017	

Labeler - U S Chemical Corporation (031457842)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(61307-213)

Revised: 8/2023 U S Chemical Corporation