

E2 ANTIBACTERIAL FOAMING SKIN CLEANSER- benzalkonium chloride soap
Betco Corporation, Ltd.

E2 Antibacterial Foaming Skin Cleanser

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Active Ingredient

Benzalkonium Chloride 0.13%

E2 Antibacterial Foaming Skin Cleanser

Uses

- For handwashing to decrease the bacteria on the skin.
- Recommended for repeated use.

E2 Antibacterial Foaming Skin Cleanser

Warnings

- **For external use only.**
- When using this product avoid contact with eyes. In case of contact flush with water
- Discontinue use if irritation or redness develops.
- Stop use and ask a doctor if irritation persists for more than 72 hours, or if condition persists for more than 72 hours.
- **KEEP OUT OF REACH OF CHILDREN.**
- If swallowed, get medical help or contact Poison Control Center right away.

E2 Antibacterial Foaming Skin Cleanser

Directions

- **Read the entire label before using this product.**
- Dispense 2 pumps of product in hands and scrub thoroughly over all surfaces of both hands for 15 seconds. Rinse with clean water.

E2 Antibacterial Foaming Skin Cleanser

Inactive Ingredients

Water, coco-glucoside, laurtrimonium chloride, cocamidopropylamine oxide, citric acid.

E2 Antibacterial Foaming Skin Cleanser

Purpose

Antimicrobial

E2 Antibacterial Foaming Skin Cleanser

KEEP OUT OF REACH OF CHILDREN

E2 Antibacterial Foaming Skin Cleanser



E2 Antibacterial Foaming Skin Cleanser

E2 Registered Antibacterial Foaming Skin Cleanser for Food Service and Healthcare Applications

Producto de limpieza para la piel en espuma antibacterial tipo E2 para servicios alimenticios y aplicaciones de atención médica

E2 SKIN CLEANSER 717

55 gal. (208 L)



Nonfood Compounds Program Listed E2 153670

Drug Facts	
Active Ingredient Benzalkonium Chloride 0.13%.....	Purpose Antimicrobial
Uses <ul style="list-style-type: none"> For hand washing to decrease bacteria on the skin. Recommended for repeated use. 	
Warnings For external use only. When using this product avoid contact with eyes. If contact occurs, rinse thoroughly with water. Stop use and ask a doctor if irritation or rash occurs. Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions <ul style="list-style-type: none"> Read the entire label before using this product. Place enough product on your palm and scrub thoroughly over all surfaces of both hands. Rinse with clean water. 	
Inactive Ingredients Water, Lauramine Oxide, Caprylyl/Capryl Glucoside, Glycerine, Sodium Benzoate, Tetrasodium EDTA, Citric Acid.	
Questions? ¿Preguntas? 888-GO BETCO (888-462-3826)	

Datos del Producto	
Ingrediente Activo Cloruro de benzalconio 0,13%.....	Propósito Antimicrobiano
Usos <ul style="list-style-type: none"> Para lavar las manos y reducir las bacterias en la piel. Recomendado para uso reiterado. 	
Advertencias Sólo para uso externo. Al utilizar este producto evite el contacto con los ojos. En caso de contacto, enjuáguese los ojos con agua. Deje de usar y consulte a un médico si se produce irritación o sarpullido. Mantener fuera del alcance de los niños. En caso de ingestión, obtenga asistencia médica o diríjase a un centro de toxicología de inmediato.	
Instrucciones <ul style="list-style-type: none"> Lea toda la etiqueta antes de usar este producto. Poner suficiente cantidad del producto en la palma y fríeguelo bien en todas las superficies de ambas manos. Enjuague con agua limpia. 	
Ingredientes Inactivos Agua, óxido de lauramina, glucósido de caprilo/caprililo, glicerina, benzoato de sodio, EDTA tetrasódico, ácido cítrico.	

NDC# 65601-817-55

SDS No. 717



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E2 ANTIBACTERIAL FOAMING SKIN CLEANSER

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65601-817
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
WATER (UNII: 059QF0KO0R)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
TETRASODIUM EDTA (UNII: MP1J8420LU)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65601-817-04	3780 mL in 1 JUG; Type 0: Not a Combination Product	01/17/2014	
2	NDC:65601-817-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	01/17/2014	
3	NDC:65601-817-03	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/17/2014	09/11/2020
4	NDC:65601-817-55	208000 mL in 1 DRUM; Type 0: Not a Combination Product	01/17/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/17/2014	

Labeler - Betco Corporation, Ltd. (005050158)

Registrant - Betco corporation, Ltd. (005050158)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd.		005050158	label(65601-817) , manufacture(65601-817)

Revised: 3/2025

Betco Corporation, Ltd.