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

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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#### 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 is 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 benzalkonium chloride soap Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:65601-817

TOPICAL

**Route of Administration** 

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
<b>BENZALKO NIUM CHLO RIDE</b> (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
WATER (UNII: 059QF0KO0R)		
LAURTRIMO NIUM CHLO RIDE (UNII: A8 1MS 10 FIC)		

F	Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
1	NDC:65601-817-04	3780 mL in 1 JUG; Type 0: Not a Combination Product	0 1/17/20 14		
2	NDC:65601-817-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	0 1/17/20 14		
3	NDC:65601-817-03	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/17/20 14	09/11/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	0 1/17/20 14		

## Labeler - Betco Corporation, Ltd. (024492831)

### Registrant - Betco corporation, Ltd. (024492831)

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

Establishment			
Name	Address	ID/FEI	Business Operations
Betco Corportation, Ltd.		024492831	label(65601-817), pack(65601-817), manufacture(65601-817)

Revised: 9/2020 Betco Corporation, Ltd.