ALCOHOL FREE FOAMING HAND SANITIZER- 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.

Alcohol Free Foaming Hand Sanitizer

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Active Ingredient

Benzalkonium Chloride 0.13%

Alcohol Free Foaming Hand Sanitizer

Uses

- Hand sanitizer to remove microorganisms on the skin.
- Use this product when soap and water are not available.

Alcohol Free Foaming Hand Sanitizer

Warnings

- **For external use only.**
- Avoid contact with eyes. If contact occurs, rinse thoroughly with water.
- Discontinue use is irritation or redness develops.
- If irritation persists for more than 72 hours, consult a physician.
- **KEEP OUT OF REACH OF CHILDREN**.
- If swallowed, get medical help or contact a Poison Control Center right away.

Alcohol Free Foaming Hand Sanitizer

Directions

- **Read the entire label before using this product.**
- Dispense 2 pumps of product onto palm of hand and rub thoroughly over all surfaces of both hands until dry.

Alcohol Free Foaming Hand Sanitizer

Inactive Ingredients

Deionized Water, Sodium PCA, PEG/PPG-8/3 Laurate, Dimethicone, PEG-3 Cocamide, Fragrance, Methyl Chloro Isothiazolinone, Methyl Isothiazolinone, D&C Green #5.

Alcohol Free Foaming Hand Sanitizer

Questions or Comments?**Phone: (800) 777-9343**

MDS information:**(800) 891-4965**

Alcohol Free Foaming Hand Sanitizer

Purpose

Antiseptic

Alcohol Free Foaming Hand Sanitizer

KEEP OUT OF REACH OF CHILDREN

Alcohol Free Foaming Hand Sanitizer

ALCOHOL FREE FOAMING HAND SANITIZER

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65601-700
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
WATER (UNII: 059QF0KO0R)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PEG-3 COCAMINE (UNII: KTM00873VC)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
2-METHYL-4-ISOTHIAZOLIN-3-ONE HYDROCHLORIDE (UNII: 3G9PNZ5P41)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	

Product Characteristics

Color	blue	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65601-700-53	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2016	
2	NDC:65601-700-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	01/01/2016	
3	NDC:65601-700-04	3780 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2016	
4	NDC:65601-700-57	550 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2016	
5	NDC:65601-700-55	207900 mL in 1 DRUM; Type 0: Not a Combination Product	01/01/2016	
6	NDC:65601-700-03	750 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/15/2016	
7	NDC:65601-700-05	1250 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/15/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/12/2012	

Labeler - Betco Corporation, Ltd. (024492831)

Registrant - Betco corporation, Ltd. (024492831)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd.		024492831	manufacture(65601-700) , pack(65601-700) , label(65601-700)