

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

Alcohol Free Foaming Hand Sanitizer

☐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 if 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

Drug Facts

Active Ingredient
Benzalkonium Chloride 0.13%.....

Purpose
Antiseptic

Uses

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

Warnings

- **For external use only.**
- Avoid contact with eyes. If contact occurs, rinse thoroughly with water.
- Discontinue use if irritation or redness develops.
- If irritation persists for more than 72 hours, consult a physician.
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Directions

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- Dispense 2 pumps of product onto palm of hand and rub thoroughly over all surfaces of both hands until dry.

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

Datos del Producto

Ingrediente Activo
Cloruro de benzalconio 0.13%.....

Propósito
Antiséptico

Usos

- Desinfectante para las manos destinado a reducir la cantidad de microorganismos de la piel.
- Utilice este producto cuando el jabón y el agua no están disponibles.

Advertencias

- **Sólo para uso externo.**
- Evite el contacto con los ojos. En caso de contacto, enjuáguese los ojos con agua.
- Deje de usarlo si se desarrolla una irritación o enrojecimiento.
- Si la irritación persiste durante más de 72 horas, consulte a un médico.
- **MANTENGA ESTE PRODUCTO FUERA DEL ALCANCE DE LOS NIÑOS.**
- En caso de ingestión, obtenga asistencia médica o diríjase a un Centro de toxicología de inmediato.

Instrucciones

- Lea la etiqueta completa antes de usar este producto.
- Aplique 2 dosis del producto en la palma de la mano y frótelo bien en todas las superficies de ambas manos hasta que se seque.

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888-GO BETCO (888-462-3826)
Betco.com





Alcohol Free Foaming Hand Sanitizer

Freshly scented, Alcohol Free Foaming Hand Sanitizer
Fresca perfumada, Espuma sanitizadora para manos sin alcohol

1 L (33.8 fl. oz.)
1.05 qt.

Item #752
SDS No. 752



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RLB6342 0375216

ALCOHOL FREE FOAMING HAND SANITIZER

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:6560 1-700
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)

