

GLACIER BLUE ANTIBACTERIAL FOAMING SKIN CLEANSER- benzalkonium chloride soap

Lawson Products, Inc

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.

Glacier Blue Antibacterial Foaming Skin Cleanser

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Warnings

- **For external use only. Do not ingest.**
- Avoid contact with eyes. If contact occurs, rinse thoroughly with water.
- Stop use and ask a doctor if irritation and redness develop.
- 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.

Directions

- **?Read the entire label before using this product.**
- ?Dispense 2 pumps of product onto palm of hand and scrub thoroughly over all surfaces of both hands.
- Rinse with clean water.

Inactive Ingredients

Water, Phenoxyethanol, Cocamine Oxide, Propylene Glycol, Decyl Glucoside, PEG-4 Rapeseedamide, Cocamidopropyl Hydroxysultaine, Glycerin, Fragrance, Sodium Chloride, Tetrasodium EDTA, Citric Acid, Ethanol, Iodopropynyl Butylcarbamate, Benzyl Acetate, Sodium Glycolate, Sodium Hydroxide, Trisodium Nitrilotriacetate, Diethyl Phthalate, FD&C Blue #1.

Questions? 888-GO BETCO (888-462-3826)

Glacier Ultra Blue Antibacterial Foaming Skin Cleanser

Purpose

Antibacterial

Glacier Blue Antibacterial Foaming Skin Cleanser

KEEP OUT OF REACH OF CHILDREN

Glacier Blue Antibacterial Foaming Skin Cleanser

Drug Facts	
Active Ingredient Benzalkonium Chloride 0.13%.....	Purpose Antibacterial
Uses • Antibacterial hand cleaner. • Use in daycare centers, hospitals, nursing homes, physician offices, dental offices and clinics.	
Warnings • For external use only. Do not ingest. • Avoid contact with eyes. If contact occurs, rinse thoroughly with water. • Stop use and ask a doctor if irritation and redness develop. • 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.	
Directions • Read the entire label before using this product. • Dispense 2 pumps of product onto palm of hand and scrub thoroughly over all surfaces of both hands. • Rinse with clean water.	
Inactive Ingredients Water, Phenoxyethanol, Cocamine Oxide, Propylene Glycol, Decyl Glucoside, PEG-4 Rapeseedamide, Cocamidopropyl Hydroxysultaine, Glycerin, Fragrance, Sodium Chloride, Tetrasodium EDTA, Citric Acid, Ethanol, Iodopropynyl Butylcarbamate, Benzyl Acetate, Sodium Glycolate, Sodium Hydroxide, Trisodium Nitrotriacetate, FD&C Blue #1.	

Sold by:
Lawson Products, Inc.
 866-LAWSON4U (866-529-7664)
 lawsonproducts.com
 8770 W. Bryn Mawr Ave., Suite 900
 Chicago, IL 60631-3515
 USA



Made In U.S.A.
1000 mL (33.8 fl. oz.)
 15739570119
 RLB370A 0275919

bag-1000ml 759-29

GLACIER BLUE ANTIBACTERIAL FOAMING SKIN CLEANSER

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62428-759
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: 059QF0K00R)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PEG-4 RAPESEEDAMIDE (UNII: 89575CN928)	

COCAMIDOPROPYL HYDROXYSLTAINE (UNII: 62V75NI93W)
GLYCERIN (UNII: PDC6A3C0OX)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
EDETATE SODIUM (UNII: MP1J8420LU)
ALCOHOL (UNII: 3K9958V90M)
BENZYL ACETATE (UNII: 0ECG3V79ZJ)
SODIUM GLYCOLATE (UNII: B75E535IM)
SODIUM HYDROXIDE (UNII: 55X04QC32I)
TRISODIUM NITRILOTRIACETATE (UNII: E3C8R2M0XD)
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)
N-ALKYL DIMETHYL BENZYL AMMONIUM CHLORIDE (C12-C18) (UNII: 9U1Q4T4ZYS)
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)
2,4-DIMETHYL-3-CYCLOHEXENE CARBOXYALDEHYDE (UNII: 452GFV2AFS)
METHYL DIHYDROJASMONATE (SYNTHETIC) (UNII: 3GW44CIE3Y)
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)
MAGNESIUM NITRATE (UNII: 77CBG3UN78)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
GERANIOL (UNII: L837108USY)
LIME OIL (UNII: UZH29XGA8G)
2-METHYLUNDECANAL (UNII: S94QNS2VY5)
.BETA.-CITRONELLOL, (+/-)- (UNII: 565OK72VNF)
ALLYL CYCLOHEXANEPROPIONATE (UNII: H4W9H3L241)
GERANYL ACETATE (UNII: 3W81YG7P9R)
CITRONELLYL ACETATE (UNII: IZ420RT3OY)
LINALOOL, (+/-)- (UNII: D81QY6I88E)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
CITRAL (UNII: T7EU0O9VPP)
FORMALDEHYDE (UNII: IHG84L3525)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62428-759-29	1000 mL in 1 BAG; Type 0: Not a Combination Product	11/12/2012	
2	NDC:62428-759-03	750 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/12/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC mono graph not final	part333E	11/12/2012	

Labeler - Lawson Products, Inc (005438890)

Registrant - Betco Corporation, Ltd (024492831)

Establishment

Name	Address	ID/FEI	Business Operations
Betco Corporation, Ltd		024492831	manufacture(62428-759) , label(62428-759)

Revised: 7/2020

Lawson Products, Inc