EVERWIPE ANTIBACTERIAL- benzalkonium chloride liquid
Brands International Corporation

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.

Everwipe Antibacterial Hand Soap

Benzalkonium Chloride - 0.13%

Purpose: Antibacterial

Effective in destroying bacteria to provide antiseptic cleaning.

Direction
- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse

For external use only

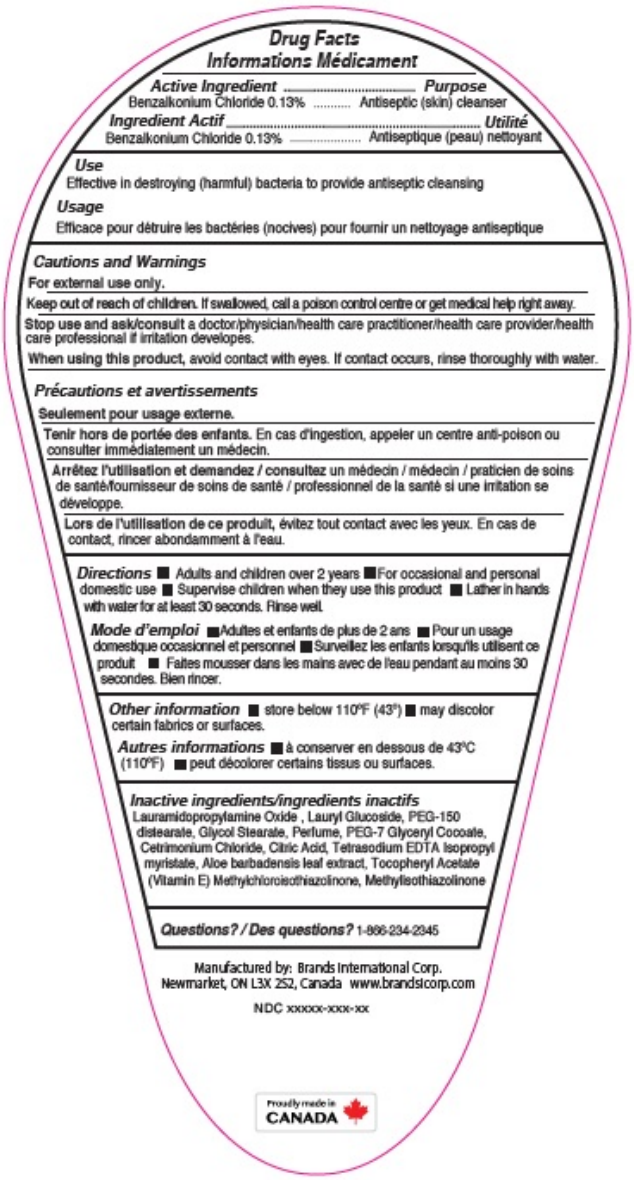
Stop use and ask a doctor if irritation or redness develops

When using this product
- do not get it into eyes. If contact occurs, rinse eye thoroughly with water

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Lauramidopropylamine oxide, Lauryl glucoside, PEG-150 distearate, Glycol Stearate, Perfume, Aloe extract, Tocopheryl Acetate, Peg 7- Glyceryl Cocoate, Cetrimonium, Chloride, Citric Acid, Glycerin, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone.



EVERWIPE ANTIBACTERIAL

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50157-264
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
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	
DITETRACYCLINE TETRASODIUM EDETATE (UNII: WX0A0IT7K5)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50157-264-43	443 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/26/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/16/2021	

Labeler - Brands International Corporation (243748238)

Registrant - Brands International Corporation (243748238)

Establishment

Name	Address	ID/FEI	Business Operations
Brands International Corporation		243748238	manufacture(50157-264)

Revised: 4/2021 Brands International Corporation