

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.



Drug Facts
Informations Médicament

Active Ingredient Benzalkonium Chloride 0.13%	Purpose Antiseptic (skin) cleanser
Ingredient Actif Benzalkonium Chloride 0.13%	Utilité Antiseptique (peau) nettoyant

Use
Effective in destroying (harmful) bacteria to provide antiseptic cleansing

Usage
Efficace pour détruire les bactéries (nocives) pour fournir un nettoyage antiseptique

Cautions and Warnings
For external use only.
Keep out of reach of children. If swallowed, call a poison control centre or get medical help right away.
Stop use and ask/consult a doctor/physician/health care practitioner/health care provider/health care professional if irritation develops.
When using this product, avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Précautions et avertissements
Seulement pour usage externe.
Tenir hors de portée des enfants. En cas d'ingestion, appeler un centre anti-poison ou consulter immédiatement un médecin.
Arrêtez l'utilisation et demandez / consultez un médecin / médecin / praticien de soins de santé/fournisseur de soins de santé / professionnel de la santé si une irritation se développe.
Lors de l'utilisation de ce produit, évitez tout contact avec les yeux. En cas de contact, rincer abondamment à l'eau.

Directions ■ Adults and children over 2 years ■ For occasional and personal domestic use ■ Supervise children when they use this product ■ Lather in hands with water for at least 30 seconds. Rinse well.

Mode d'emploi ■ Adultes et enfants de plus de 2 ans ■ Pour un usage domestique occasionnel et personnel ■ Surveillez les enfants lorsqu'ils utilisent ce produit ■ Faites mousser dans les mains avec de l'eau pendant au moins 30 secondes. Bien rincer.

Other information ■ store below 110°F (43°) ■ may discolor certain fabrics or surfaces.

Autres informations ■ à conserver en dessous de 43°C (110°F) ■ peut décolorer certains tissus ou surfaces.

Inactive ingredients/ingrédients inactifs
Lauramidopropylamine Oxide, Lauryl Glucoside, PEG-150 distearate, Glycol Stearate, Perfume, PEG-7 Glyceryl Cocoate, Cetrimonium Chloride, Citric Acid, Tetrasodium EDTA Isopropyl myristate, Aloe barbadensis leaf extract, Tocopheryl Acetate (Vitamin E) Methylchloroisothiazolinone, Methylisothiazolinone

Questions? / Des questions? 1-866-234-2345

Manufactured by: Brands International Corp.
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NDC xxxxxx-xxxx-xxx

Proudly made in
CANADA

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