MOONLITE 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.

Moon Lite Antibacterial Soap: Sulfate-Free

Benzalkonium Chloride - 0.13%

Purpose - Antiseptique(skin)cleanser

Uses Effective in destorying(harmful) bacteria to provide antiseptic cleansing.

Warning For external use only

Stop use and ask a doctor if irritation or redness develops

When using the product

- avoid contact wiith 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

Direction

- Adults and children over 2 year
- For occasional and personal domestic use
- Supervise children when use this product
- Lather in hands with water for at least 30 seconds
- Rinse well

- S□tore below 110F(43)
- May discolor certain fabrics or surfaces.

Water,Lauramidopropylamine Oxide (Triaminox LO), Lauryl Glucoside, Citric Acid, PEG-150 distearate, Glycol Stearate, PEG-7 Glyceryl Cocoate,Cetrimonium Chloride, Tetrasodium EDTA,Isopropyl myristate, Aloe Leaf, Tocopheryl Acetate, Methylchloroisothiazolinone, Methylisothiazolinone,

Drug Facts / Informations Médicament

Active Ingredient — **Purpose**
Benzalkonium Chloride 0.13% — Antiseptic (skin) cleanser

Ingredient Actif — **Utilité**
Benzalkonium Chloride 0.13% — 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 procuit, 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/ingredients 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) Methylchbroisothiazolinone, Methylisothiazolinone

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

DISTRIBUTED BY:
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MANUFACTURED BY:
Brands International Corporation
594 Newpark Blvd, Newmarket, ON
L3X 2S2 Canada

MOONLITE ANTIBACTERIAL

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50157-511
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**
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
WATER (UNII: 059QF0KO0R)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50157-511-23	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/23/2020	

Labeler - Brands International Corporation (243748238)

Registrant - Brands International Corporation (243748238)

Establishment

Name	Address	ID/FEI	Business Operations
Brands International Corporation		243748238	manufacture(50157-511)

Revised: 9/2020 Brands International Corporation