CORALITE ANTIBACTERIAL MOIST- benzalkonium chloride solution
United Exchange Corp.

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.

Coralite Antibacterial Moisturizing Pouch Wipe, 30ct 83103, 2019

Drug Facts

Active Ingredient	Purpose
Benzalkonium Chloride 0.115%.....	Antibacterial

Uses

- Decreases bacteria on skin and surfaces

Warnings

For external use only

When using this product avoid contact with the eyes, if contact occurs rinse thoroughly with water.

Stop use and ask a doctor if

- Irritation or rash develops and persists for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poision Control Center right away.

Directions

- Peel back seal using tab. Do not pull seal all the way off. Pull sheet from center and remove as needed. Reseal pouch by pressing seal back down to retain moisture after each use.
- Apply thoroughly to hands and face as desired and allow to dry without wiping.
- Dispose of wipe in the proper container. Do not flush down the toilet.

Inactive ingredients

2-Bromo-2-Nitropropane-1,3-Diol,3-Iodo-2-propynyl Butyl Carbamate, Aloe Gel Powder 100:1, Aqua, Chamomilla Recutita Flower Extract (Propylene Glycol, Aqua), Citric Acid, Lauryl Glucoside, Phenoxyethanol, Polysorbate 20, Propylene Glycol, Sodium Citrate, Terasodium EDTA

Distributed by:

UNITED EXCHANGE CORP.

17211 Valley View Ave.

Cerritos, CA 90703

MADE IN CHINA

CORALITE ANTIBACTERIAL MOIST

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65923-831
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.115 g

Inactive Ingredients	
Ingredient Name	**Strength**
BRONOPOL (UNII: 6PU1E16C9W)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
CITRIC ACID ACETATE (UNII: DSO12WL7AU)	

LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-831-30	30 in 1 POUCH; Type 0: Not a Combination Product	09/26/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/26/2013	

Labeler - United Exchange Corp. (840130579)

Revised: 5/2020 United Exchange Corp.