

UNICOWIPES- benzethonium chloride cloth
Unicoadds 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.

Active Ingredient	Purpose
Benzethonium Chloride 0.2%	Antibacterial

Uses: decreases bacteria on skin

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

Unicowipes

Warnings:

For external use only

Do not use if you have allergic to any of the ingredients

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

Stop use and ask a doctor if irritation or rash develops and continues for more than 72 hours

Directions:

Tear the pack. Pull out the wet wipe.

Use wet wipe to clean hands, face or legs without soap or water and discard.

Inactive Ingredients

deionized water, cocoamido propyl betain, glycerin pharma, mono propylene glycol, PEG-7 glycery cocoate, polysorbate-20, fragrance, citric acid, benzyl alcohol, methylchloroisothiazolinone, methylisothiazolinone.

Ingredients Facts

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Questions or Comments?

Email info@unicowipes.com

UNICOWIPES

benzethonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69651-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.001 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3011KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69651-000-00	1 in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/19/2015	

Labeler - Unicoadds Corp. (043612349)

Establishment

Name	Address	ID/FEI	Business Operations
Sabah Gida Saglik Temizlik Urunleri Dolum Ve Ambalaj Sana		365616970	manufacture(69651-000)

Revised: 4/2015

Unicoadds Corp.