

WET WIPES- benzethonium chloride swab
Kareway Product, Inc.

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.

Pure-Aid Wet Wipes

Active Ingredients

Benzethonium Chloride 0.3%

Purpose

Antibacterial

Use

decrease bacteria on skin

Warnings

For external use only

Do not use

if you are 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

Keep out of reach of children

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

Directions

adults and children 2 years and over

- apply to hands
- allow to dry without wiping

Children under 2 years

- ask a doctor before use

Inactive Ingredients

Water, Butylene Glycol, Alcohol, Hyaluronic Acid, Polysorbate 80, Sodium Lauryl Sulfate, Disodium

Lareth Sulfosuccinate,

Glycerin, Phytosqualane, Anthemis Nobilis Flower Extract, Camellia Sinensis Leaf Extract, Cyclopia Intermedia Leaf Extract,

Tocopheryl Acetate, Phenoxyethanol, Disodium EDTA, Fragrance

Principle Display Panel

Fresh Scent

Wet wipes

Antibacterial formula

30 wipes



WET WIPES

benzethonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67510-0888
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.3 in 100

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
HYALURONIC ACID (UNII: S270N0TRQY)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DISODIUM LAURETH SULFO SUCCINATE (UNII: D6DH1DTN7E)	
GLYCERIN (UNII: PDC6A3C0OX)	
CYCLOPIA INTERMEDIA LEAF (UNII: 29UXG2GG8O)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67510-0888-3	30 in 1 POUCH; Type 0: Not a Combination Product	06/30/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/30/2015	

Labeler - Kareway Product, Inc. (121840057)

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Kareway Product, Inc.