CB ALCOHOL WIPES- benzalkonium chloride swab swab Nutritech Pharmaceuticals, 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.

Active Ingredient

Benzalkonium chloride 0.1%

Purpose

Antibacterial

Use

decreases bacteria on skin

Directions

For adults and children of 2 years and over.

Children under 2 years ask a doctor before use.

Allow to dry without wiping.

Warnings

For external use only.

Do not use

over large areas of the body

if you are allergic to any of the ingredients

when using this product, avoid contact with eyes and face.

If contact occurs, flush 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 aPoisonControlCenter right away.

Inactive Ingredients

water, sodium lauryl sulfate, fragrance, 2-bromo-2-nitropropane-1, 3-diol, aloe barbadensis leaf extract, disodium EDTA, trithanolamine, glycerin, to copheryl acetate









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Drug Facts (continued)

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Other information store at room temperature

Inactive ingredient

Water, Sodium Lauryl sulfate, Fragrance,

2-Bromo-2-Nitropropane-1, 3-Diol, Aloe Barbadensis Leaf Extrad-Disodium EDTA, Triethanolamine, Glycerin, Tocopheryl Acetate

CB ALCOHOL WIPES

benzalkonium chloride swab swab

Product Information

Product Type HUMAN OTC DRUG NDC:73529-352 Item Code (Source)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength

BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -UNII:7N6JUD5X6Y)

BENZALKONIUM CHLORIDE

 $0.1\,\mathrm{g}$ in 100 g

Inactive Ingredients				
Ingredient Name	Strength			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)				
CUPRIC BIS(TRIETHANO LAMINE) (UNII: YBM44X0B6H)				
2-BROMO-2-NITROETHANOL (UNII: FA22WV2B2Z)				
WATER (UNII: 059QF0KO0R)				
DISO DIUM EDTA-COPPER (UNII: 6 V475AX06U)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
GLYCERIN (UNII: PDC6 A3C0 O X)				

l	Packaging				
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
l	1 NDC:73529-352-01	0.1 g in 1 POUCH; Type 0: Not a Combination Product	03/16/2020		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/16/2020		

Labeler - Nutritech Pharmaceuticals, Inc (117342868)

Revised: 3/2020 Nutritech Pharmaceuticals, Inc