QUALITY CHOICE ANTIBACTERIAL MOIST TOWELETTES - benzalkonium chloride cloth
Chain Drug Marketing Association

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.

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.10%

Purpose

Antimicrobial

Uses
For hand sanitizing to decrease bacteria on skin.
Recommended for repeated use.

Warnings

For external use only

When using this product

avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor

if irritation or redness develpos, or if condition persists for more than 72 hours.

Keep out of reach of children.

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

Directions
Open cap, remove one wipe and unfold.
Wet hands throughly with product and allow to air dry.

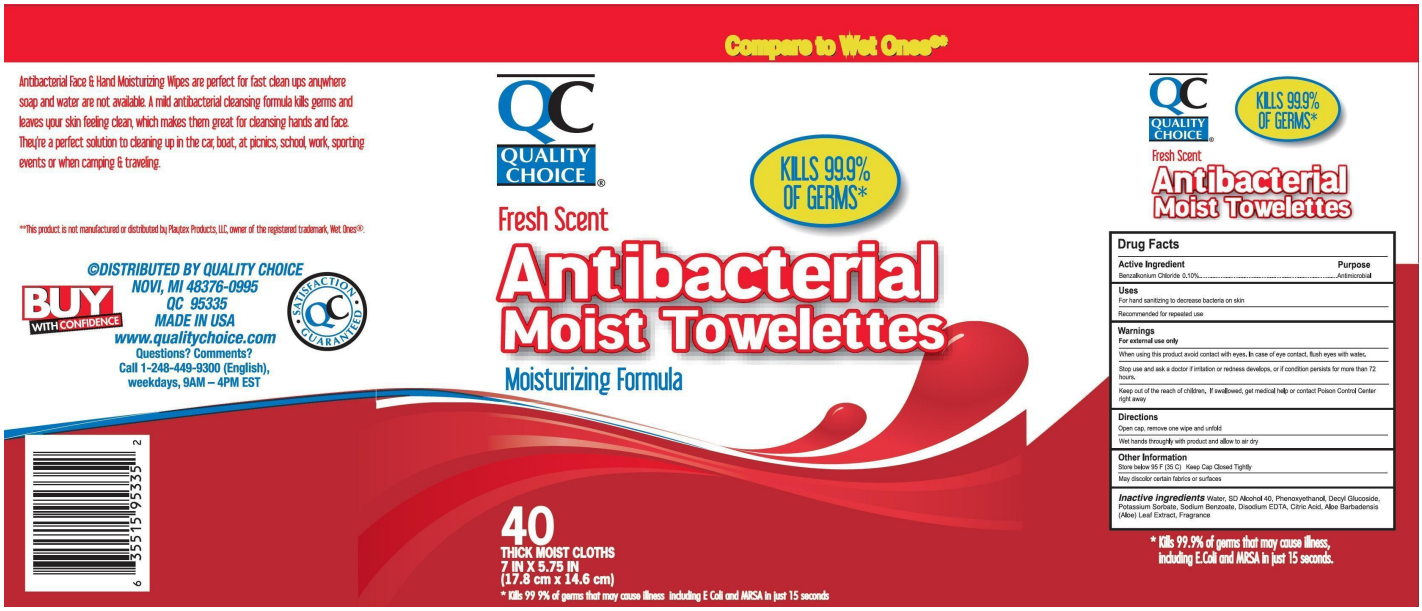
Other Information
Store below 95 F(35C) Keep Cap-Closed Tightly
May discolor certain fabrics or surfaces.

Inactive ingredients

Water, SD Alcohol 40, Phenoxyethanol, Decyl Glucoside, Potassium Sorbate, Sodium Benzoate, Disodium EDTA, Citric Acid, Aloe Barbadensis (Aloe) Leaf Extract, Fragrance

Principal Display Panel
Compare to Wet Ones**

QC
QUALITY
CHOICE
KILLS 99.9%
OF GERMS*
Fresh Scent
Antibacterial
Moist Towelettes
Moisturizing Formula
40
THICK MOIST CLOTHS
7 IN X 5.75 IN
(17.8 cm x 14.6 cm)
*Kills 99.9 %of germs that may cause illness including E Coli and MRSA in just 15 seconds



QUALITY CHOICE ANTIBACTERIAL MOIST TOWELETTES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.10 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-100-40	40 in 1 CANISTER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/03/2013	

Labeler - Chain Drug Marketing Association (011920774)

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