CARROLL ULTREX ANTIMICROBIAL WIPES- benzalkonium chloride cloth
Carroll Company

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.

Carroll ® Ultrex Antimicrobial Wipes

Drug Facts

Active ingredient

Benzalkonium chloride 0.13% w/w

Purpose

Antimicrobial

Uses

- for hand cleaning to decrease bacteria on the skin
- kills 99.9% of most common germs

Warnings

For external use only

When using this product

- do not use in or near eyes
- discontinue use if irritation and redness develop

Keep out of reach of children. In case of accidental ingestion, seek medical attention or contact a poison control center immediately.

Directions

- Use textured side of wipe to clean hands
- wet hands thoroughly with product
- allow to dry without rinsing

Other information

- store at room temperature

Inactive ingredients

Water, 2-Phenoxyethanol, Alkylpolyglucoside, C 8-10,C10-16, Fragrance, Lauramine,N,N-dimethyl, N-oxide, Tocopherol acetate, Aloe barbadensis leaf juice, PEG-40 Lanolin, Tetrasodium EDTA, Methylchloroisothiazoline, Methylisothiazoline.

Manufactured in the U.S.A. by:
CARROLL ® COMPANY
2900 W. Kingsley Rd., Garland, Texas 75041

PRINCIPAL DISPLAY PANEL - 70 Cloth Canister Label

WIPES
CARROLL®
ULTREX

Antimicrobial Wipes
-Kills 99.9% of common germs
-Contains Aloe and Vitamin E
-Moisturizes and conditions skin
-Alcohol free

CITRUS SCENTED
Dual Textured
Wipe

KEEP OUT OF REACH OF CHILDREN
WARNING
For external use only.

Contains 70
12" x 10" Pre-Moistened Wipes

ITEM #
681

CARROLL ULTREX ANTIMICROBIAL WIPES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10685-681
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.01417 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10685-681-97	70 in 1 CANISTER; Type 0: Not a Combination Product	09/15/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/15/2012	

Labeler - Carroll Company (007372329)

Registrant - Carroll Company (007372329)

Establishment

Name	Address	ID/FEI	Business Operations
Carroll Company		007372329	manufacture(10685-681)

Revised: 1/2018 Carroll Company