JUICEY ANTIBACTERIAL WIPES- benzalkonium chloride cloth Fuson LLC

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.

Juicey Antibacterial Wipes Pouch

Active Ingredient

Benzalkonium Chloride .13% v/v - Antiseptic

Inactive Ingredients

Aqua, Parfum, C12-15 Pareth-12, Glycerin, Polysorbate 20, Propylene Glycol, Tetrasodium EDTA, Piroctone Olamine, Tocopheryl Acetate (Vitamin E), Panthenol, Citric Acid, Aloebarbadensis Leaf Extract

Purpose

Antiseptic- antibacterial hand towelettes

Use

Wipes used on hands to reduce bacteria on the skin

Warnings

Keep away from flame and direct sunlight. Children should not use this product without adult supervision. For external use only. If ingested, contact a poison control center or seek medical attention. Keep away from eyes.

Do not use

Do not use if allergic to any of its ingredients. Do not use near eyes. Do not use on children under 2 months of age. Consult a doctor before use if pregnant, breastfeeding, are under the age of 6, or have sensitibe skin.

When Using

If eye contact occurs, remove any contact lenses and flush eyes with water repeatedly. If swallowed, contact a poinson control center immediately or seek medical attention.

Store product out of reach of children. Children under the age of 12 should be supervised during use.

Directions

Open the hard plastic snap cover to reveal the adhesive label that reads, "OPEN". Peel back the adhesive label and remove one wipe at a time. Wipe your hands with the antibacterial wipe until all of the solution is absorbed. Seal the packaging tightly closed when done.

Storage

Store out of reach of children. Store in a cool, dry place. Do not store near food, drinks, or animal feed. Store away from direct sunlight and open flame.

Package Label

60 Count Antibacterial Wipes Pouch NDC: 78141-901



JUICEY ANTIBACTERIAL WIPES

benzalkonium chloride cloth

Product Information	roduct Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78141-901			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	0.007 g in 5.33 g	

Inactive Ingredients		
Ingredient Name	Strength	
EDETATE SO DIUM (UNII: MP1J8420LU)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		
WATER (UNII: 059QF0KO0R)		
POLYSORBATE 20 (UNII: 7T1F30 V5YH)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
GLYCERIN (UNII: PDC6A3C0OX)		
PANTHENOL (UNII: WV9CM0O67Z)		
C12-15 PARETH-12 (UNII: 131316 X18 L)		
PIROCTONE OLAMINE (UNII: A4V5C6R9FB)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:78141-901-60	60 in 1 POUCH	10/27/2020		
1		5.33 g in 1 PACKET; 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	10/27/2020		

Labeler - Fuson LLC (113376279)

Revised: 10/2020 Fuson LLC