PERSONAL CARE WET WIPES CITRUS- benzalkonium chloride cloth Delta Brands, 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.

Lucky Wet WipesBzlknmHCl0.13indivpckt

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Uses

for hand sanitizing to 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, ethyl alcohol, aloe barbadensis leaf juice, PEG-60 lanolin, fragrance, tetrasodium EDTA, phenoxyethanol, potassium sorbate, quaternium-52, citric acid

Package Label

Drug Facts

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Manufactured for DELTA BRANDS INC.
1890 Palmer Avenue
Larchmont, New York 10538 USA

ITEM NO: 11873 11873/01(419)



Antibacterial Kills 99.9% of Germs

PERSONAL CARE WET WIPES CITRUS

benzalkonium chloride cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:20276-436	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g	

Inactive Ingredients		
Ingredient Name	Strength	
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
WATER (UNII: 059QF0KO0R)		
ALCOHOL (UNII: 3K9958V90M)		
PEG-60 LANOLIN (UNII: K20I1D27ET)		
QUATERNIUM-52 (UNII: 588EQF3H1P)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
EDETATE SODIUM (UNII: MP1J8420LU)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:20276-436- 15	15 in 1 BOX	01/07/2021		
1	NDC:20276-436- 01	1 in 1 PACKET; Type 0: Not a Combination Product			
2	NDC:20276-436- 18	18 in 1 BOX	01/07/2021		
2	NDC:20276-436- 01	1 in 1 PACKET; Type 0: Not a Combination Product			
3	NDC:20276-436- 30	30 in 1 BOX	01/07/2021		
3	NDC:20276-436- 01	1 in 1 PACKET; Type 0: Not a Combination Product			
4	NDC:20276-436- 40	40 in 1 BOX	01/07/2021		
4	NDC:20276-436- 01	1 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	01/07/2021	

Labeler - Delta Brands, Inc (102672008)

Revised: 12/2022 Delta Brands, Inc