WET WIPES- benzalkonium chloride swab
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 Wipes Package

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antiseptic

Uses

•decreases bacteria on the skin

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

•do not get into eyes. •If contact occurs, rinse thoroughly with water.

Stop use

Stop use and ask a doctor if irritation or rash develops and continues for more than 72 hours

Keep out of reach of children

Keep out of reach of children. If swallowed get medical help or contact a Poison Control Center right away.

Directions

• for adults and children 2 years and over use on hands and face to clean and refresh, allow skin to air dry. For children under 2 years of age ask a doctor before use.

Inactive Ingredients

water, propylene glycol, cocamidopropyl betaine, peg-7 glyceryl cocoate, fragrance, benzyl alcohol, methylchloroisothiazolinine, methylisothiazolinone, tetrasodium EDTA, peg-40 hydrogenated castor oil, cetrimonium chloride, citric acid, aloe vera extract, alphatocopherol acetate

Package Label

WET WIPES

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:20276-431
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:20276-431-30	30 in 1 PACKAGE; Type 0: Not a Combination Product	01/23/2007	
2	NDC:20276-431-36	36 in 1 PACKAGE; Type 0: Not a Combination Product	01/23/2007	
3	NDC:20276-431-40	40 in 1 PACKAGE; Type 0: Not a Combination Product	06/06/2016	
4	NDC:20276-431-16	16 in 1 PACKAGE; Type 0: Not a Combination Product	06/06/2016	
5	NDC:20276-431-80	80 in 1 PACKAGE; Type 0: Not a Combination Product	10/04/2019	
6	NDC:20276-431-60	60 in 1 PACKAGE; Type 0: Not a Combination Product	04/27/2020	
7	NDC:20276-431-50	50 in 1 PACKAGE; Type 0: Not a Combination Product	09/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/23/2007	

Labeler - Delta Brands, Inc (102672008)