WET WIPES- 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 Wipes Package BZK.13

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

decreases bacteria on the 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 ■ apply to hands

2 years and over ■ allow to dry

without wiping

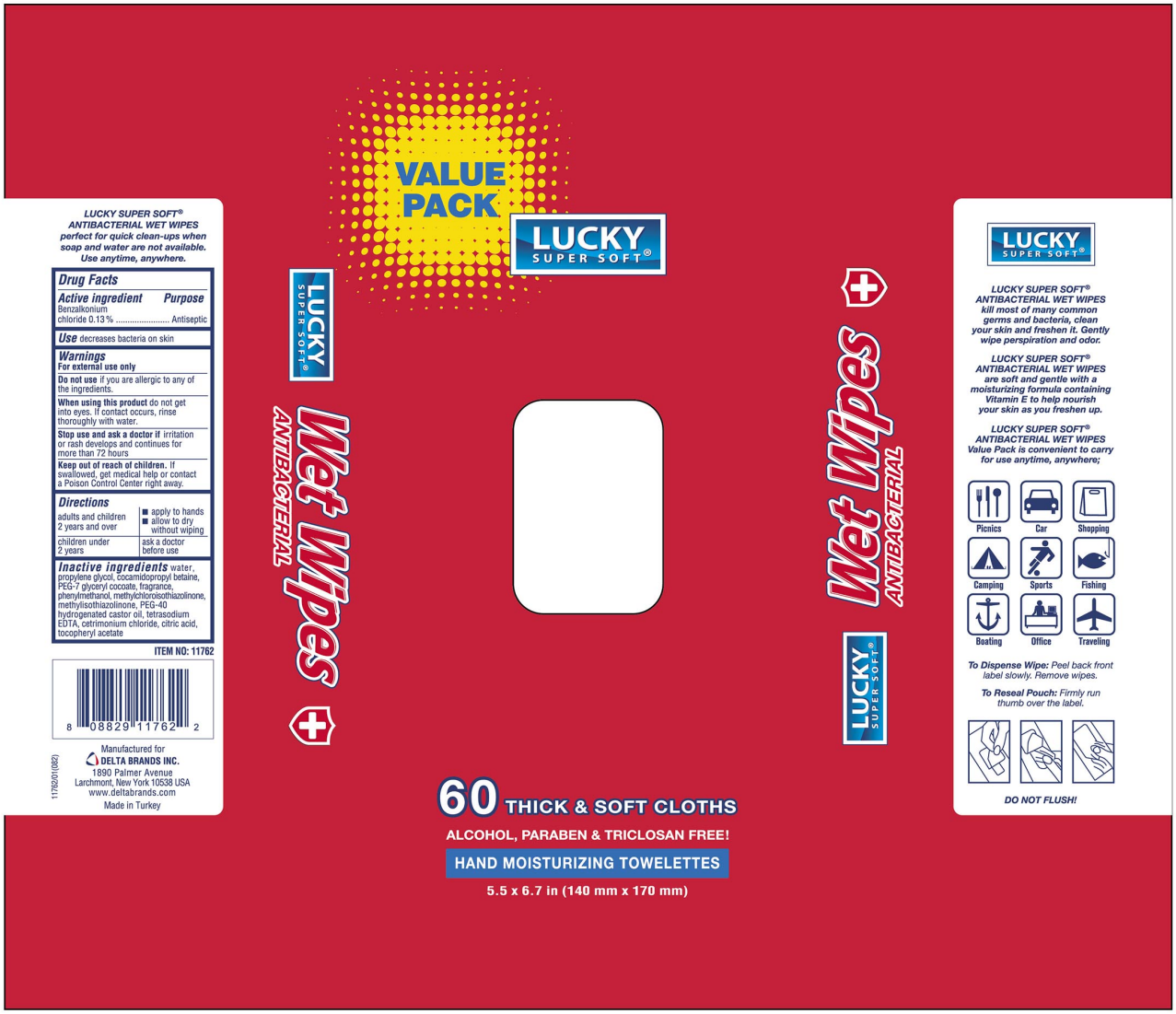
children under ask a doctor

2 years before use

Inactive ingredients

water, propylene glycol, cocamidopropyl betaine,PEG-7 glyceryl cocoate, fragrance, phenylmethanol, methylchloroisothiazolinine, methylisothiazolinone, PEG-40 hydrogenated castor oil, tetrasodium EDTA, cetrimonium chloride, citric acid, tocopherol acetate

Package Label



WET WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:20276-435
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
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20276-435-30	30 in 1 PACKAGE; Type 0: Not a Combination Product	05/18/2020	
2	NDC:20276-435-40	40 in 1 PACKAGE; Type 0: Not a Combination Product	05/18/2020	
3	NDC:20276-435-16	16 in 1 PACKAGE; Type 0: Not a Combination Product	05/18/2020	
4	NDC:20276-435-80	80 in 1 PACKAGE; Type 0: Not a Combination Product	05/18/2020	
5	NDC:20276-435-60	60 in 1 PACKAGE; Type 0: Not a Combination Product	05/18/2020	
6	NDC:20276-435-12	12 in 1 PACKAGE; Type 0: Not a Combination Product	06/25/2020	
7	NDC:20276-435-36	36 in 1 PACKAGE; Type 0: Not a Combination Product	06/25/2020	
8	NDC:20276-435-35	35 in 1 CANISTER; Type 0: Not a Combination Product	06/30/2020	
9	NDC:20276-435-42	42 in 1 CANISTER; Type 0: Not a Combination Product	06/30/2020	
10	NDC:20276-435-50	50 in 1 PACKAGE; Type 0: Not a Combination Product	03/04/2021	

11	NDC:20276-435-24	24 in 1 PACKAGE; Type 0: Not a Combination Product	07/21/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/18/2020	

Labeler - Delta Brands, Inc (102672008)

Revised: 7/2023

Delta Brands, Inc