WET NAPS ANTIBACTERIAL HAND WIPES- antibacterial hand wipes swab
Westrock Rkt 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.

Warning section

For external use only. Keep out of reach of children. If swallowed get medical help or contact a posion control center right away. Do not use in eyes. Discontinue use if irritation develop. If condition persists for more than 72 hours consult a doctor.

Purified water, SD alcohol 40, Sorbic Acid, PPG-2 Hydroxyethyl Cocamide, Disodium EDTA, Aloe Barbadensis leaf join, Fragrance

Directions

To dispense, life cover, remove seal, pull center sheet from toll, twist to point, feed through dispenser hole in cover. Keep lid closed to prevent moisture loss. Use as a part of your cleaning routine. Rub product onto hands and allow to dry. Discard after single use.

Keep out of reach of children.

Antiseptic handwash

Benzalkonium chloride 0.13% mg/ml

Strength Benzalkonium chloride 0.13% mg/ml

Cloth wipe administraion

handwipe.jpg

Consumer facing image of hand wipes.


The Original
Wet-Nap
ANTIBACTERIAL
HAND WIPES
Moisturizes with Aloe
Kills 99.99% of Germs
Fresh Scent
40

WET NAPS ANTIBACTERIAL HAND WIPES

antibacterial hand wipes swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71928-7011(NDC:10819-7011)
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 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
SORBIC ACID (UNII: X045WJ989B)	
PPG-2 HYDROXYETHYL COCAMIDE (UNII: 34N07GUJ3X)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71928-7011-1	193 mL in 1 CANISTER; Type 0: Not a Combination Product	01/01/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2014	

Labeler - Westrock Rkt Company (603215539)

Establishment

Name	Address	ID/FEI	Business Operations
WESTROCK RKT COMPANY		603215539	relabel(71928-7011) , repack(71928-7011)

Revised: 12/2017 Westrock Rkt Company