

WET ONES- benzalkonium chloride swab
Edgewell Personal Care Brands 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.

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

decreases 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: Rub on hands for 30 seconds and allow skin to dry without wiping.

children under 2 years: ask a doctor before use.

Inactive Ingredients

Water, Alcohol Denat., Phenoxyethanol, PEG-8 Dimethicone, Caprylyl Glycol, Dihydroxypropyl PEG-5 Linoleammonium Chloride, Potassium Sorbate, Disodium EDTA, Citric Acid, Fragrance, Aloe Barbadensis Leaf Juice.

Questions or Comments?

Call 1-866-WET-111S (1-866-938-1117), M-F

Principal Display Panel

TEAR HERE

Wet

Ones(R)

Kids

ANTIBACTERIAL

HAND WIPE

kills 99.99% of Germs*

Hypoallergenic Pediatrician tested

fruity fresh

1 WIPE

Drug Facts

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www.WetOnes.com

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Wet
Ones
Kids
ANTIBACTERIAL
HAND WIPE
write On Wrapper
kills 99.99% of Germs
tough on germs
gentle on skin
24 INDIVIDUALLY
WRAPPED WIPE
Recyclable
cardboard box



WET ONES

benzalkonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63354-926
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
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALCOHOL (UNII: 3K9958V90M)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63354-926-24	24 in 1 BOX; Type 0: Not a Combination Product	10/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/01/2023	

Labeler - Edgewell Personal Care Brands LLC (151179769)

Revised: 10/2023

Edgewell Personal Care Brands LLC