

**WALGREENS ANTIBACTERIAL HAND WIPES- benzalkonium chloride cloth
Walgreen Co.**

Walgreens Antibacterial Hand Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride 0.11%

Purpose

Antibacterial

Uses

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 immediately.

Directions

•**Adults and children 2 years and older:**

Wipe hands thorough with product and allow to dry without wiping.

•**Children under 2 years of age:** Consult a doctor before use.

Inactive ingredients

Water, Alcohol, Propylene Glycol, Phenoxyethanol, Lauryl Glucoside, Tetrasodium EDTA, 2-Bromo-2-Nitropropane-1,3-Diol, Citric Acid, Fragrance, Iodopropynyl

Butylcarbamate, Aloe Barbadensis Leaf Extract, Chamomilla Recutita (Matricaria) Flower Extract.

Gentle cleansing

Kills 99.9% of germs

Hypoallergenic

DO NOT FLUSH

DISTRIBUTED BY: **WALGREEN CO.**

DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED

walgreens.com 1-877-274-8358

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Packaging

20 wipes

Walgreens Antibacterial Hand Wipes

- Gentle cleansing
- Kills 99.9% of germs
- Hypoallergenic

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ANTIBACTERIAL Hand Wipes NEW

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Hypoallergenic

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OPEN

20 wipes
5.5 IN x 7.5 IN (14 cm x 19 cm)

PLASTIC POUCH

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DEERFIELD, IL 60015

Walgreens
100% SATISFACTION GUARANTEED

walgreens.com 1-877-274-8358
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ITEM 744179 W10376-0524-G

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W3ORG0424-F

75 Wipes



WALGREENS ANTIBACTERIAL HAND WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-5189
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.11 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
EDETATE SODIUM (UNII: MP1J8420LU)	
BRONOPOL (UNII: 6PU1E16C9W)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CHAMOMILE (UNII: FGL3685T2X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-5189-01	75 in 1 POUCH	05/21/2024	
1		3.83 mL in 1 PATCH; Type 0: Not a Combination Product		
2	NDC:0363-5189-02	20 in 1 POUCH	05/21/2024	
2		3.83 mL in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/21/2024	

Labeler - Walgreen Co. (008965063)