

VIROCIDANTIBACTERIAL WIPES- benzalkonium chloride, isopropyl alcohol cloth
Virocide 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.

Virocide Antibacterial Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride 0.25%

Isopropyl Alcohol 5%

Purpose

Benzalkonium Chloride Antimicrobial Agent

Isopropyl Alcohol Antimicrobial Agent

Uses decrease bacteria on skin.

Warnings

For external use only.

When using this product, avoid contact with the eyes.

In case of contact, rinse eyes thoroughly with water.

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 2 years: ask a doctor before using.

Questions or Comments?

(M-F 9AM-5PM)

1(833)494-5459

Inactive Ingredients

Water, Aloe Vera (Aloe Barbadensis)

Leaf, Anhydrous Citric Acid,

Dimethicone, Edetate Sodium

Tetrahydrate, Hydantoin,

Polysorbate 20, Tocopheryl Acetate

(Vitamin E), Glycerin, Fragrance,

Methylchloroithiazolinone,
Methylisothiazolinone.

Package Label

BOX OF 12-PACKETS



PACKET OF 100 WIPES



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VIROCID ANTIBACTERIAL WIPES

benzalkonium chloride, isopropyl alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77005-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	2.5 mg
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	5 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
EDETATE SODIUM TETRAHYDRATE (UNII: L13NHD21X6)	
HYDANTOIN (UNII: I6208298TA)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77005-007-02	12 in 1 BOX	10/07/2020	
1	NDC:77005-007-01	100 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Virocide LLC (117469716)

Registrant - Virocide LLC (117469716)

Revised: 10/2020

Virocide LLC