

**GERMISEPT ANTIBACTERIAL HAND SANITIZING WIPES- benzalkonium chloride cloth
Innovent Inc**

GERMisept Antibacterial Hand Sanitizing Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride 0.1%

Purpose

Antibacterial

Uses:

Decrease bacteria on the skin.

Warnings

- **For external use only.**

When using this product

- avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Do not use

if irritation and redness develop.

Stop use and ask a doctor

if condition persists for more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact Poison Control Center right away.

Directions

- Open the flip top lid, remove one wipe to use.
- Wipe hands thoroughly the product and allow to dry without wiping.
- Close flip top lid after use to retain moisture.

Other information

- Dispose of wipe in the proper container.
- Do not flush down the toilet.

*Kill Claims Against: E.Coli & Staphylococcus.

Inactive ingredients

2-Bromo-2-Nitropropane-1,3-Diol, Alcohol, Aloe Barbadensis Leaf Extract, Chamomilla Recutita Flower Extract, Citric Acid, Fragrance, Iodopropynyl Butylcarbamate, Lauryl Glucoside, Phenoxyethanol, Propylene Glycol, Tetrasodium EDTA, Water.

Package Labeling:

Carry Bag of 4 Packs x 50ct GERMisept Antibacterial Hand Sanitizing Wipes



GERMISEPT ANTIBACTERIAL HAND SANITIZING WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70335-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BRONOPOL (UNII: 6PU1E16C9W)	
ALCOHOL (UNII: 3K9958V90M)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70335-012-01	4 in 1 BAG	02/10/2021	
1		50 in 1 PACKAGE		
1		4.3 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/10/2021	

Labeler - Innovent Inc (079973489)

Revised: 12/2023

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