

BENZALKONIUM CHLORIDE- sanitizing hand wipes cloth
ALO NEW YORK LLC

Peppermint Eucalyptus + Aloe

Active ingredients

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- Hand Sanitizer to help reduce bacteria on the skin
- Recommended for repeated use

Warnings

For external use only.

When using this product

avoid contact with eyes. In case of contact, flush with water.

Stop use and ask a doctor

If redness or irritation develops and persists for more than 72 hours

Keep out of reach of children

Children should be supervised when using this product

Directions

Take wipe and rub thoroughly over all surface of both hands. Wet hands thoroughly with wipes. Rub hand together briskly to dry without wiping. Dispose of wipe. Do not flush.

Inactive ingredients

Purified Water, Decyl Glucoside, Aloe Barbadensis (Aloe) Leaf Extract, Phenoxyethanol, Citric Acid, Sodium Benzoate, Potassium Sorbate, Eucalyptus Globulus Oil, Mentha Piperita (Peppermint) Oil.

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0013 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
LAVANDULA ANGUSTIFOLIA FLOWER (UNII: 19AH1RAF4M)	
ANIBA ROSAEODORA WHOLE (UNII: 5ZTZ6VF78R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82355-552-01	20 in 1 PACKAGE	01/04/2024	
1		1 mg in 1 PACKAGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	01/04/2024	

Labeler - ALO NEW YORK LLC (110122374)

Revised: 1/2024

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