BENZALKONIUM CHLORIDE- sanitizing hand wipes cloth
ALO NEW YORK 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.

Sweet Orange Bergamot + Aloe

Active ingredients

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- for hand sanitizing to decrease bacteria on the skin
- recommended for repeated use

Warnings

For external use only.

Do not use

if you are allergic to any of these ingredients.

When using this product

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

Stop use and ask a doctor

if irritation or redness develops, or if condition persists for more than 72 hours.

Keep out of the reach of children.

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

Directions

- adults and children 2 years and over: apply to hands: allow to dry without wiping
- children under 2 years: ask a doctor before use.
- **To dispense**: Peel back fron label slowly. Remove wipes.
- **To reseal**: Firmly rub thumb over label. Dispose of wipe in trash. Do not flush.

Other information

store below 95

- store below 95°F (35°C).
- keep closed tightly
- may discolor certain fabrics or surfaces.

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.

Principal Display Panel

ALO

20 Skin-Softening Sanitizing Hand Wipes

BENZALKONIUM CHLORIDE

sanitizing hand wipes cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82355-551
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)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	
ORANGE OIL (UNII: AKN3KSD11B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82355-551-01	20 in 1 PACKAGE	12/21/2021	
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 not final	part333A	12/21/2021	

Labeler - ALO NEW YORK LLC (110122374)

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