LAFLARE SANTIX ANTIBACTERIAL MOISTURIZING WIPES- benzalkonium chloride liquid
NAICO

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.

Drug Facts

ACTIVE INGREDIENT

Benzalkonium Chloride 0.1%

INACTIVE INGREDIENTS

Water, Phenoxyethanol, Glycerin, Sodium Benzoate, Polysorbate 20, Disodium EDTA, Citric Acid, Tocopheryl Acetate, Macadamia Integrifolia/Tetraphylla Seed Oil, Anthemis Nobilis Flower Extract, Fragrance

PURPOSE

Antimicrobial

WARNINGS

For external use only.

Do not use
• in children less than 2 months of age
• on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Posion Control Center right away

Uses

Instant hand antimicrobial to decrease bacteria on the skin

Directions

■ Wet hands thoroughly with product and allow to dry without wiping

■ Do not flush

Other information

Storage in a cool, dry place, Avoid freezing and excessive heat above 40 □ (104 □)

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

for 15 sec. and discard. Be sure to reseal the front label.

CAUTION: FOR EXTERNAL USE ONLY
Do not use on eyes, mouth and ears. Keep out of reach of children unless under adult supervision.

STORAGE INSTRUCTIONS
Store in a cool, dry place, away from direct sunlight.

LAFLARE SANTIX ANTIBACTERIAL MOISTURIZING WIPES

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75536-0005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
Polysorbate 20 (UNII: 7T1F30V5YH)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CHAMAEMELUM NOBILE FLOWER (UNII: O2T154T6OG)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75536-0005-1	81 g in 1 CONTAINER; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/01/2020	

Labeler - NAICO (694725335)

Registrant - NAICO (694725335)

Establishment

Name	Address	ID/FEI	Business Operations
NAICO		694725335	manufacture(75536-0005)

Revised: 5/2020 NAICO