

**PULLIO HAND SANITIZING WIPES ALOE VERA- benzalkonium chloride cloth
YES SALES**

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 0.13%

INACTIVE INGREDIENTS

Water, Glycerin, Sodium Benzoate, Ethylhexylglycerin, Polysorbate20, Aloe Barbadensis Leaf Extract, Butylene Glycol,1,2-Hexanediol, Disodium EDTA, Citric Acid, Tocopheryl Acetate, Fragrance

PURPOSE

Antiseptic

WARNINGS

For external use only

Stop use and ask a doctor if

- hypersensitivity symptoms such as erythema, itching and dermatitis happen
- skin irritation happens

Do not use

- in combination with soap or antibacterial cleansing agents
- the product for a long time in the same area as swelling, inflammation or sickness may occur due to absorption through the skin

When using this product

- avoid using repeatedly in the same area, skin irritation may occur
- avoid getting into the eyes (if contact occurs, wash well with clean water)
- if following abnormal symptoms persist, discontinue use :

Irritation around the eyes, ears, mucous membranes, including the mouth, under the skin irritation and rashes

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children

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

Uses

- Instant hand antiseptic to decrease bacteria on the skin

Directions

- Clean with wipes and let dry
- Do not flush

Other information

- Read the directions and warnings before use
- Avoid freezing and excessive heat above 40C (104F)

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



PULLIO HAND SANITIZING WIPES ALOE VERA

benzalkonium chloride cloth

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:80618-010 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-----------------------|-------------|
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 0.13 in 100 |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| WATER (UNII: 059QF0KO0R) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| SODIUM BENZOATE (UNII: OJ245FE5EU) | |
| Ethylhexylglycerin (UNII: 147D247K3P) | |
| Polysorbate 20 (UNII: 7T1F30V5YH) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |
| Butylene Glycol (UNII: 3XUS85K0RA) | |
| 1,2-Hexanediol (UNII: TR046Y3K1G) | |
| EDETATE DISODIUM ANHYDRO US (UNII: 8NLQ36F6MM) | |
| CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP) | |
| .ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:80618-010-01 | 60 in 1 PACKET; Type 0: Not a Combination Product | 09/01/2020 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333E | 09/01/2020 | |

Labeler - YES SALES (080733755)**Registrant** - YES SALES (080733755)**Establishment**

| Name | Address | ID/FEI | Business Operations |
|-------|---------|-----------|------------------------|
| NAICO | | 694725335 | manufacture(80618-010) |

Revised: 9/2020

YES SALES