

HAND SANITIZE WIPES- benzalkonium chloride cloth
Landy International

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.

51706-958 HAND SANITIZE WIPES 0.13% Benzalkonium chloride

Active Ingredient(s)

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

Uses sanitizing hand wipes to help decrease bacteria on the skin. recommended for single use.

Warnings

For external use only

Do not use if you are allergic to any of the ingredients.

When using this product do not get into eyes. If contact occurs, flush eyes in water. stop use and ask a doctor if irritation develops and condition for more than 72 hours.

Keep out of reach of children. Children should be supervised when using this product.

Directions

Directions wipe liberally over the hands & body, rub into skin until

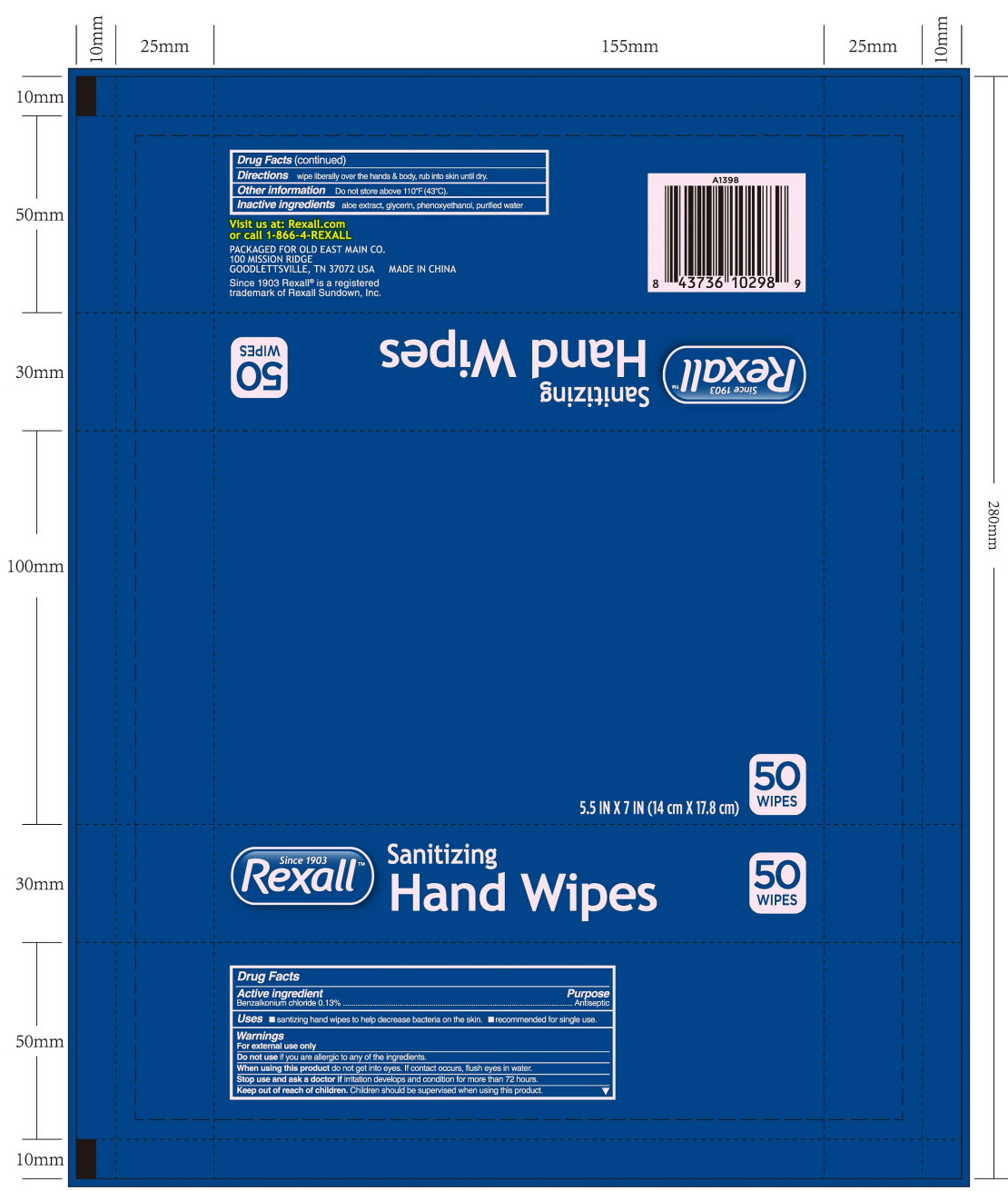
Other information

- Other information Do not store above 110°F(43°C).

Inactive ingredients

aloe extract, glycerin, phenoxyethanol, purified water

Package Label - Principal Display Panel



Drug Facts (continued)
Directions wipe liberally over the hands & body, rub into skin until dry.
Other information Do not store above 110°F (43°C).
Inactive ingredients aloe extract, glycerin, phenoxylethanol, purified water

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 GOODLETTSVILLE, TN 37072 USA MADE IN CHINA
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Sanitizing Hand Wipes



5.5 IN X 7 IN (14 cm X 17.8 cm) 50 WIPES



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HAND SANITIZE WIPES

benzalkonium chloride cloth

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:51706-958 |
| 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 g in 100 g |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| PHENOXYETHANOL (UNII: HIE492ZZ3T) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| WATER (UNII: 059QF0K00R) | |
| ALOE VERA LEAF (UNII: ZY81Z83H0X) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:51706-958-01 | 50 in 1 BAG | 10/30/2020 | |
| 1 | | 0.0048 g in 1 NOT APPLICABLE; 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 | 10/30/2020 | |

Labeler - Landy International (545291775)

Establishment

| Name | Address | ID/FEI | Business Operations |
|--|---------|-----------|------------------------|
| Fujian Yifa Healthcare Products Co.,Ltd. | | 420510793 | manufacture(51706-958) |