# SOFT AND DRI DRIGEL- aluminum zirconium octocholrohydrex stick The Village Company

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.

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#### **Active Ingredient**

Aluminum Zirconium Octachlorohydrex Gly (16.4%)

Antiperspirant

#### Use

reduces underarm perspiration

Warnings For external use only

#### Stop use and ask a doctor if

rash or irritation occurs

#### Keep out of reach of chlidren

If swallowed, get medical help or contact a posion control center right away

#### Ask a doctor before use if

you have kidney disease

#### Do not use

on broken skin

#### **Directions**

Apply to underarms only.

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#### **Inactive ingredients**

Water, (Auqua, EAU), Alcohol denat., Cyclopentsiloxane, Propylene Glycol, Dimethicone, Calcium cholride, Trisiloxane, Pentasiloxane, PEG/PPG-18/18, Dimethicone, T-butyl alcohol, Denaton-ium Benzoate, Fragrance (Parfum), Benzyl Salicylate, Cirtronellol, Benzyl Alcohol, Geraniol, Limonene





### SOFT AND DRI DRIGEL

aluminum zirconium octocholrohydrex stick

| Product Information     |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:69752-102 |
| Route of Administration | TOPICAL        |                    |               |

| Active Ingredient/Active Moiety  |  |                  |
|--|--|------------------|
| Ingredient Name  | Basis of Strength                          | Strength         |
| ALUMINUM ZIRCONIUM OCTACHLOROHYDREX GLY (UNII: P9 D3YP29 MY) (ALUMINUM ZIRCONIUM OCTACHLOROHYDREX GLY - UNII:P9 D3YP29 MY) | ALUMINUM ZIRCONIUM<br>OCTACHLOROHYDREX GLY | 164 mg<br>in 1 g |
|  |  |                  |

| Inactive Ingredients            |          |  |  |
|---------------------------------|----------|--|--|
| Ingredient Name                 | Strength |  |  |
| DIMETHICO NE (UNII: 92RU3N3Y1O) |          |  |  |

| l | Packaging          |   |                             |                    |
|---|--------------------|---|-----------------------------|--------------------|
|   | # Item Code        | Package Description                               | <b>Marketing Start Date</b> | Marketing End Date |
| ı | 1 NDC:69752-102-06 | 1 in 1 TUBE                                       | 11/13/20 15                 |                    |
|   | 1                  | 85 g in 1 TUBE; Type 0: Not a Combination Product |                             |                    |

| Marketing Information |  |                      |                    |
|-----------------------|--|----------------------|--------------------|
| Marketing Category    | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph final   | part350                                  | 11/13/2015           |                    |
|                       |  |                      |                    |

## Labeler - The Village Company (172208105)

### **Registrant -** Apex International (015226132)

| Establishment |         |           |                        |  |
|---------------|---------|-----------|------------------------|--|
| Name          | Address | ID/FEI    | Business Operations    |  |
| VVF           |         | 024177178 | manufacture(69752-102) |  |

Revised: 12/2019 The Village Company