

**GILLETTE CLEARSHIELD WILD RAIN CLEAR- aluminum zirconium  
octachlorohydrate gly gel  
The Procter & Gamble Manufacturing Company**

-----  
**Gillette® ClearShield Wild Rain Clear Gel (lower strength)**

***Drug Facts***

**Active ingredient**

Aluminum zirconium octachlorohydrate Gly 16% (anhydrous)

**Purpose**

Antiperspirant

**Use**

reduces underarm wetness

**Warnings**

**For external use only.**

**Do not use** on broken skin

**Ask a doctor before use if you have** kidney disease

**Stop use if** rash or irritation occurs

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- apply to underarms only

**Inactive ingredients**

water, alcohol denat., cyclopentasiloxane, propylene glycol, dimethicone, calcium chloride, PEG/PPG-18/18 dimethicone, fragrance

**Questions?**

**1-800-445-5388**

**Dist. by PROCTER & GAMBLE,  
CINCINNATI, OH 45202.**

Gillette®  
CLEARSHIELD  
72HR SWEAT PROTECTION  
POWER RUSH  
CLEAR GEL  
ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY  
ANTIPERSPIRANT/DEODORANT  
NET WT. 3.8 OZ (107 g)



| GILLETTE CLEARSHIELD WILD RAIN CLEAR                                                                                      |                |                                            |                  |
|---------------------------------------------------------------------------------------------------------------------------|----------------|--------------------------------------------|------------------|
| aluminum zirconium octachlorohydrate gly gel                                                                              |                |                                            |                  |
| Product Information                                                                                                       |                |                                            |                  |
| Product Type                                                                                                              | HUMAN OTC DRUG | Item Code (Source)                         | NDC:84126-012    |
| Route of Administration                                                                                                   | TOPICAL        |                                            |                  |
| Active Ingredient/Active Moiety                                                                                           |                |                                            |                  |
| Ingredient Name                                                                                                           |                | Basis of Strength                          | Strength         |
| ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY (UNII: P9D3YP29MY)<br>(ALUMINUM ZIRCONIUM OCTACHLOROXYDREX GLY - UNII:P9D3YP29MY) |                | ALUMINUM ZIRCONIUM<br>OCTACHLOROXYDREX GLY | 16 g<br>in 100 g |
| Inactive Ingredients                                                                                                      |                |                                            |                  |
| Ingredient Name                                                                                                           |                |                                            | Strength         |

|                                                     |  |
|-----------------------------------------------------|--|
| <b>ALCOHOL</b> (UNII: 3K9958V90M)                   |  |
| <b>WATER</b> (UNII: 059QF0KO0R)                     |  |
| <b>CYCLOMETHICONE 5</b> (UNII: 0THT5PCI0R)          |  |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)          |  |
| <b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)               |  |
| <b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)          |  |
| <b>PEG/PPG-18/18 DIMETHICONE</b> (UNII: 9H0AO7T794) |  |

### Packaging

| # | Item Code        | Package Description                                    | Marketing Start Date | Marketing End Date |
|---|------------------|--------------------------------------------------------|----------------------|--------------------|
| 1 | NDC:84126-012-10 | 107 g in 1 CYLINDER; Type 0: Not a Combination Product | 01/15/2024           |                    |
| 2 | NDC:84126-012-01 | 2 in 1 CELLO PACK                                      | 01/15/2024           |                    |
| 2 |                  | 107 g in 1 CYLINDER; Type 0: Not a Combination Product |                      |                    |

### Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|------------------------------------------|----------------------|--------------------|
| OTC Monograph Drug | M019                                     | 01/15/2024           |                    |

**Labeler** - The Procter & Gamble Manufacturing Company (004238200)