**DAWNMIST ANTIPERSPIRANT DEODORANT- aluminum chlorohydrate lotion**
**Dukal Corporation**

*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.*

---

**DawnMist Antiperspirant Deodorant**

**Active Ingredient**

Aluminum Chlorohydrate 4.5%

**Purpose**

Antiperspirant

**Use**

- Reduces underarm perspiration.

**Warnings**

For external use only.

**Do not use**

on broken skin.

**Discontinue use if**

■irritation and redness
develop ■ If condition persists for more than 72
hours consult a doctor.

**Ask a doctor before use**

if you have kidney disease.

**Keep out of reach of children**

■If swallowed, get medical help and contact Poison Control Center right away ■ Use only as directed

**Directions**

Apply to underarms only.

**Inactive Ingredients**

Water, Propylene Glycol, Hydroxypropyl Methylcellulose, Diazolidinyl Urea, Fragrance, PEG-20 Sorbitan Stearate, Triethanolamine, Menthol, Iodopropynyl Butylcarbamate

**Principal Display Panel - Bottle Label**

NDC 65517-1010-1

**DawnMist®**

**DEODORANTS**

**Roll-On**
**Antiperspirant**

REDUCES WETNESS & ODOR

FRESH SCENT

1.5 FL. OZ. (44ml)


## DAWNMIST ANTIPERSPIRANT DEODORANT

aluminum chlorohydrate lotion

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:65517-1010 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ALUMINUM CHLOROHYDRATE** (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M) | ALUMINUM CHLOROHYDRATE | 45 mg in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **WATER** (UNII: 059QF0KO0R) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **HYPROMELLOSES** (UNII: 3NXW29V3WO) | |
| **DIAZOLIDINYL UREA** (UNII: H5RIZ3MPW4) | |
| **IODOPROPYNYL BUTYLCARBAMATE** (UNII: 603P14DHEB) | |
| **TROLAMINE** (UNII: 9O3K93S3TK) | |
| **MENTHOL** (UNII: L7T10EIP3A) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:65517-1010-1 | 44 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product | 01/10/2013 | 09/28/2021 |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph final | part350 | 01/10/2013 | 09/28/2021 |

## Labeler - Dukal Corporation (791014871)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| KEN PRIMA COSMECEUTICALS SDN. BHD. | | 865792209 | manufacture(65517-1010) |

Revised: 11/2018 Dukal Corporation