DAWNMIST ANTIPERSPIRANT DEODORANT- aluminum chlorohydrate liquid
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, Mineral Oil, Cetyl Alcohol, Ceteareth-24, Isopropyl Palmitate, Magnesium Aluminum Silicate, Methylparaben, Sodium Benzoate, Propylparaben

Package Label

DAWNMIST ANTIPERSPIRANT DEODORANT

aluminum chlorohydrate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65517-1005
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)	
MINERAL OIL (UNII: T5L8T28FGP)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65517-1005-1	59 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	01/10/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	01/10/2013	

Labeler - Dukal Corporation (791014871)

Revised: 10/2018 Dukal Corporation