

DAWNMIST ANTIPERSPIRANT DEODORANT- aluminum chlorohydrate liquid
Dukal LLC

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-20, Isopropyl Palmitate, Magnesium Aluminum Silicate, Methylparaben, Fragrance, Sodium Benzoate, Propylparaben

Package Label

DawnMist

DEODERANTS

Roll-On

Antiperspirant

REDUCES WETNESS & ODOR

FRESH SCENT

1.5 FL. OZ. (44ml)

NDC 65517-1009-1


DEODORANTS

Roll-On Antiperspirant

REDUCES WETNESS & ODOR
FRESH SCENT

1.5 FL. OZ. (44 ml)

Drug Facts	
Active Ingredients	Purpose
Aluminum Chlorohydrate 4.5%	Antiperspirant
Use: Reduce underarm perspiration.	
Warnings: Do not apply to broken skin.	
■ If a skin rash develops, discontinue use.	
■ Keep out of the reach of children.	
Inactive Ingredients: Water, Paraffinum Liquidum, Cetearyl Alcohol, Ceteareth-24, Isopropyl Palmitate, Magnesium Aluminum Silicate, Methylparaben, Propylparaben, Sodium Benzoate, Fragrance.	

REF RD15 

Manufactured for:
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Made in Malaysia
D10161216 Rev2


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DAWNMIST ANTIPERSPIRANT DEODORANT

aluminum chlorohydrate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65517-1009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZ W13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZ W13M)			ALUMINUM CHLOROHYDRATE	45 mg in 1 mL
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
MINERAL OIL (UNII: T5L8T28FGP)				
CETYL ALCOHOL (UNII: 936JST6JCN)				
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)				
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)				
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65517-1009-1	44 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	01/10/2013	11/02/2026
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M019	01/10/2013	11/02/2026	

Labeler - Dukal LLC (791014871)