

**DRY IDEA ADVANCED DRY ANTIPERSPIRANT ROLL-ON, POWDER FRESH- dry
idea advanced dry antiperspirant roll-on, powder fresh liquid
Thriving Brands LLC**

82699-201



82699-201

Drug Facts

Dry Idea®^{MD}
Never Let Them See You Sweat®

Active Ingredient**Purpose**

Aluminum Zirconium Pentachlorohydrex Gly 16.3%.....Antiperspirant

Use • Reduces underarm perspiration • Extra effective

Warnings

For external use only.

Do not use on broken skin.

Stop use and ask a doctor if rash or irritation occurs.

Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

Ask a doctor before use if you have kidney disease.

Directions Shake well. Apply to underarms only.

Inactive ingredients Cyclomethicone, Distearidimonium Hectorite, Propylene Carbonate, Fragrance, Tocopheryl Acetate (Vitamin E Acetate), Isopropyl Myristate, Gossypium Herbaceum (Cotton) Seed Oil

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Cincinnati, OH USA 45242 1-888-480-9571

www.DryIdea.com
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DRY IDEA®

AdvancedDry®

Fast Drying
Silky Smooth Feel
With Vitamin E

POWDER FRESH

3.25 FL OZ (96 ml)

300000

Roll-On
Antiperspirant
& Deodorant

UP TO
72 HR
ODOR PROTECTION

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dry idea advanced dry antiperspirant roll-on, powder fresh liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82699-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name**Basis of Strength****Strength**

ALUMINUM ZIRCONIUM

ALUMINUM ZIRCONIUM PENTACHLOROHYDREX GLY (UNII: 94703016SM) (ALUMINUM ZIRCONIUM PENTACHLOROHYDREX GLY - UNII:94703016SM)	ALUMINUM ZIRCONIUM PENTACHLOROHYDREX GLY	16.3 g in 16.3 g
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Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE (UNII: NMQ347994Z)	

Product Characteristics

Color	brown (Light Brown)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82699-201-03	96.1 g in 1 BOTTLE, WTH APPLICATOR; Type 0: Not a Combination Product	03/31/2023	
2	NDC:82699-201-04	576.6 g in 1 PACKAGE; Type 0: Not a Combination Product	09/28/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	03/31/2023	

Labeler - Thriving Brands LLC (118346160)

Establishment

Name	Address	ID/FEI	Business Operations
KDC/One Development Corporation, Inc.		204006464	analysis(82699-201) , manufacture(82699-201) , pack(82699-201) , label(82699-201)

Revised: 12/2023

Thriving Brands LLC