

**SURE ORIGINAL SOLID POWDER- aluminum zirconium trichlorohydrate
gly stick
Idelle Labs, Ltd**

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.

Sure Original Solid Antiperspirant Deodorant Powder

**SURE®
48hr CONFIDENCE
POWDER FRESH
original solid
ANTI-PERSPIRANT & DEODORANT
Net Wt 2.7oz (76g)**

Drug Facts

Active Ingredient

Aluminum Zirconium Trichlorohydrate GLY 16% (anhydrous)

Purpose

Antiperspirant

Uses

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

Cyclopentasiloxane, Stearyl Alcohol, Talc, Dimethicone, Hydrogenated Castor Oil, Fragrance, Polyethylene, Silica, Dipropylene Glycol, Behenyl Alcohol.

Questions? 1 800 487-7273

www.suredeodorant.com

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El Paso, TX 79912 USA
Sure[®] is a trademark of Kaz Europe Sarl.

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SUU71607 LBL B01

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aluminum zirconium trichlorohydrate gly stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41595-5516
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY (UNII: T27D6T99LH) (ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY - UNII:T27D6T99LH)	ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY	16 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

TALC (UNII: 7SEV7J4R1U)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
DOCOSANOL (UNII: 9G1OE216XY)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41595-5516-1	76 g in 1 CANISTER; Type 0: Not a Combination Product	12/22/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	12/22/2009	

Labeler - Idelle Labs, Ltd (128822926)

Registrant - Idelle Labs, Ltd (128822926)

Establishment

Name	Address	ID/FEI	Business Operations
KDC/ONE Development Corporation, Inc.		204006464	manufacture(41595-5516) , analysis(41595-5516) , label(41595-5516) , pack(41595-5516)

Revised: 8/2021

Idelle Labs, Ltd