SURE INVISIBLE SOLID REGULAR- aluminum zirconium tricholorohydrex 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 Invisible Solid Regular

SURe ® 48hr CONFIDENCE REGULAR invisible solid ANTI-PERSPIRANT & DEODORANT Net Wt 2.6oz (73g)

Drug Facts

Active ingredient

Aluminum Zirconium Trichlorohydrex GLY 19% (anhydrous)

Purpose

Antiperspirant

Use

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, Dimethicone, Petrolatum, PPG-14 Butyl Ether, Hydrogenated Castor Oil, Talc, Fragrance, Mineral Oil, Behenyl Alcohol

Questions?

1-800-487-7273 www.suredeodorant.com

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SURE INVISIBLE SOLID REGULAR

aluminum zirconium tricholorohydrex gly stick

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41595-5510
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALUMINUM ZIRCONIUM TRICHLOROHYDREX GLY (UNII: T27D6T99LH) (ALUMINUM ZIRCONIUM TRICHLOROHYDREX GLY - UNII:T27D6T99LH)	ALUMINUM ZIRCONIUM TRICHLOROHYDREX GLY	19 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
CYCLOMETHICONE 5 (UNII: 0THT5PCIOR)		

STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PETROLATUM (UNII: 4T6H12BN9U)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
TALC (UNII: 7SEV7J4R1U)	
MINERAL OIL (UNII: T5L8T28FGP)	
DOCOSANOL (UNII: 9G10E216XY)	

Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:41595- 5510-2	73 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(41595-5510), analysis(41595-5510), label(41595-5510), pack(41595-5510)

Revised: 8/2021 Idelle Labs, Ltd