

**BRUT SIGNATURE ANTI-PERSPIRANT AND DEODORANT- aluminum zirconium trichlorohydrate gly stick**  
**Sodalis USA LLC**

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**Brut Signature Anti-Perspirant and Deodorant**

***Drug Facts***

***Active ingredient***

Aluminum Zirconium Trichlorohydrate Gly 16% (anhydrous)

***Purpose***

antiperspirant

***Use***

reduces underarm wetness

***Warnings***

**For external use only.**

**Stop use if**

rash or irritation occurs.

**Do not use**

on broken skin.

**Keep out of reach of children.**

In case of ingestion, get medical help or contact a Poison Control Center right away.

**Ask a doctor before use if you have**

kidney disease.

***Directions***

apply to underarms only.

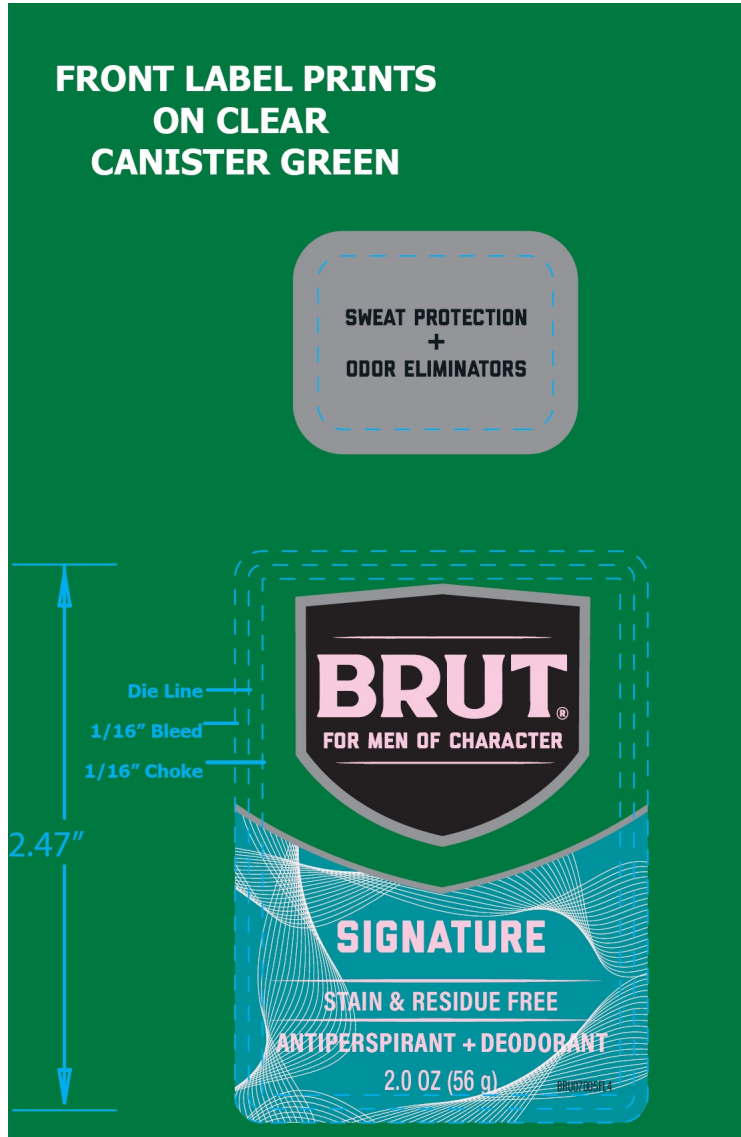
***Inactive ingredients:***

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

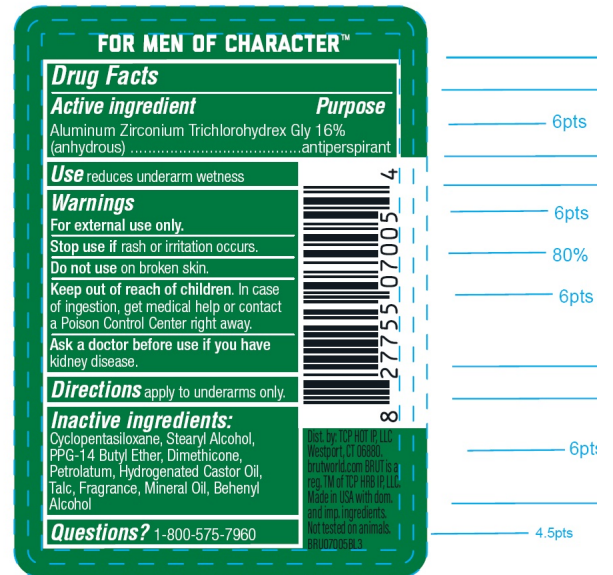
# Questions?

1-800-575-7960

## Label 81277-001-01

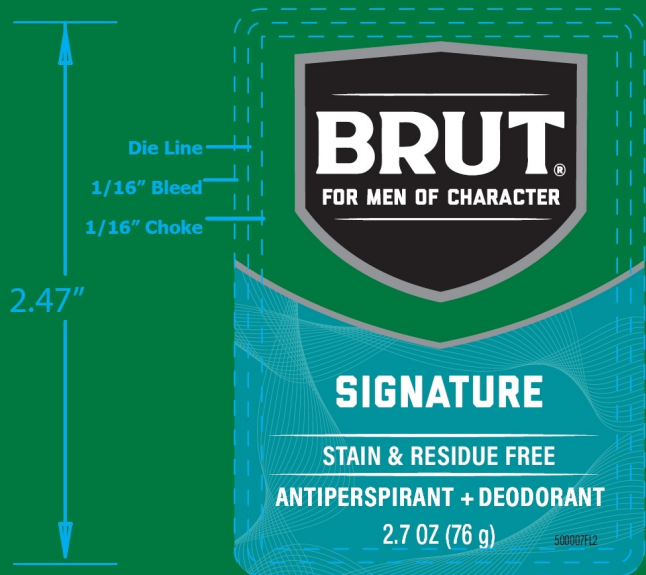


## BACK LABEL PRINTS ON WHITE



## Label 81277-001-02

**FRONT LABEL PRINTS  
ON CLEAR  
CANISTER GREEN**



**BRUT SIGNATURE ANTI-PERSPIRANT AND DEODORANT**

aluminum zirconium trichlorohydrate stick

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:81277-001
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY</b> (UNII: T27D6T99LH) (ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY - UNII:T27D6T99LH)	ALUMINUM ZIRCONIUM TRICHLOROXYDREX GLY	19 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>CYCLOMETHICONE 5</b> (UNII: 0THT5PCI0R)	
<b>STEARYL ALCOHOL</b> (UNII: 2KR89I4H1Y)	

<b>PPG-14 BUTYL ETHER</b> (UNII: R199TJT95T)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>HYDROGENATED CASTOR OIL</b> (UNII: ZF94AP8MEY)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>DOCOSANOL</b> (UNII: 9G1OE216XY)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81277-001-01	56 g in 1 CONTAINER; Type 0: Not a Combination Product	01/01/2010	
2	NDC:81277-001-02	76 g in 1 CONTAINER; Type 0: Not a Combination Product	01/01/2010	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	01/01/2010	

**Labeler** - Sodalis USA LLC (119375257)

Revised: 11/2024

Sodalis USA LLC