

**MELAO WHITENING ANTIPERSPIRANT DEODORANT STICK (ROSE)- melao
whitening antiperspirant deodorant stick (rose) stick
Guangzhou Haishi Biological Technology Co., Ltd.**

MELAO WHITENING ANTIPERSPIRANT DEODORANT STICK (ROSE)

Antiperspirant

ALUMINUM ZIRCONIUM TETRACHLOROXYDREX GLY 15.5%

DIRECTION: Spin out the paste, apply evenly under the armpit.

Ask a doctor before use if you have kidney disease.

If swallowed, get medical help or contact a Poison Control Center right away.

Stop use if rash or irritation occurs.

Do not use on broken skin.

Dry first before putting on clothes. Make sure your armpits are clean, dry, and not injured or recently shaved. Clean the armpits the next day. There are no restrictions if you want to use perfume or other deodorants. Keep out of eyes.

Keep out of reach of children

Use regularly every day for the first 1 week of use and every 3-5 days thereafter as needed.

OZOKERITE 10%
CETEARYL ALCOHOL 4%
BUTYROSPERMUM PARKII (SHEA BUTTER) 18%
CAPRYLIC/CAPRIC TRIGLYCERIDE 10%
ETHYLHEXYL PALMITATE 24.8%
GLYCERYL STEARATE 4%
SORBITAN STEARATE 4%
CERA ALBA 6%
ASCORBIC ACID 0.5%
TOCOPHEROL 1%
ASCORBYL TETRAISOPALMITATE 0.2%
PARFUM 2%



MELAO WHITENING ANTIPERSPIRANT DEODORANT STICK (ROSE)

melao whitening antiperspirant deodorant stick (rose) stick

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:60771-0014

Route of Administration

TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY (UNII: 8O386558JE) (ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY - UNII:8O386558JE)	ALUMINUM ZIRCONIUM TETRACHLOROHYDREX GLY	15.5 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
ASCORBYL TETRAISOPALMITATE (UNII: 47143LT58A)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
SORBITAN STEARATE (UNII: NVZ4I0H58X)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60771-0014-1	68 g in 1 BOX; Type 0: Not a Combination Product	05/27/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	05/27/2025	

Labeler - Guangzhou Haishi Biological Technology Co., Ltd. (421262738)

Establishment			
Name	Address	ID/FEI	Business Operations
Guangzhou Haishi Biological Technology Co., Ltd.		421262738	manufacture(60771-0014)

Revised: 5/2025

Guangzhou Haishi Biological Technology Co., Ltd.