BAN ROLL-ON ANTIPERSPIRANT DEODORANT SATIN BREEZE- aluminum chlorohydrate liquid
Kao USA Inc.

Ban Roll-On Antiperspirant Deodorant Satin Breeze

Drug Facts

Active ingredient

Aluminum chlorohydrate 18%

Purpose

Antiperspirant

Use

reduces underarm perspiration

Warnings

For external use only

Do not use on broken skin

Stop use if rash or irritation occurs

Ask a doctor before use if you have kidney disease

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

apply to underarms only

Other information

store at room temperature

Inactive ingredients

water, PPG-11 stearyl ether, steareth-2, steareth-20, fragrance, disodium EDTA, helianthus annuus (sunflower) seed oil, phellodendron amurense bark extract, hordeum distichon (barley) extract, santalum album (sandalwood) extract

Questions? 1-866-226-3363

www.bandeodorant.com

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Dist. by Kao USA Inc. Cincinnati, OH 45214

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ban

satin breeze
INVISIBLE ROLL-ON

CRUELTY FREE • PARABEN FREE • DYE FREE

antiperspirant deodorant

3.5 FL OZ (103 mL)

BAN ROLL-ON ANTIPERSPIRANT DEODORANT SATIN BREEZE

aluminum chlorohydrate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10596-339
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	20 g in 103 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-20 (UNII: L0Q8IK9E08)	
SANDALWOOD (UNII: 3641YW25N2)	
PHELLODENDRON AMURENSE BARK (UNII: PBG27B754G)	
BARLEY (UNII: 5PWM7YLI7R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PPG-11 STEARYL ETHER (UNII: S4G2J0Y0LG)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10596-339-35	103 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	01/01/2012	
2	NDC:10596-339-14	4 in 1 PACKAGE	08/24/2020	
2		103 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		
3	NDC:10596-339-70	2 in 1 PACKAGE	08/24/2020	
3		103 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	01/01/2012	

Labeler - Kao USA Inc. (004251617)

Revised: 9/2024 Kao USA Inc.