BAN PURELY GENTLE ROLL-ON ANTIPERSPIRANT DEODORANT UNSCENTED-aluminum chlorohydrate liquid
Kao USA Inc.

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.

Ban Purely Gentle Roll-On Antiperspirant Deodorant Unscented

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, 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

BAN is a trademark of Kao Corp.

Dist. by Kao USA Inc. Cincinnati, OH 45214

ban

purely gentle
non-irritating - unscented

INVISIBLE ROLL-ON

CRUELTY FREE • PARABEN FREE • DYE FREE

antiperspirant deodorant

3.5 FL OZ (103 mL)

BAN PURELY GENTLE ROLL-ON ANTIPERSPIRANT DEODORANT UNSCENTED

aluminum chlorohydrate liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10596-402
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	18 g in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10596-402-35	103 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	01/01/2018	
2	NDC:10596-402-14	4 in 1 PACKAGE	08/24/2020	
2		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 final	M019	01/01/2018	

Labeler - Kao USA Inc. (004251617)

Revised: 3/2023 Kao USA Inc.