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

Ban 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, helianthus annuus (sunflower) seed oil, phellodendron amurense bark extract, hordeum distichon (barley) extract, santalum album (sandalwood) extract

Questions? 1-866-226-3363

www.feelbanfresh.com

BAN is a trademark of Kao Corp.

Dist. by Kao USA Inc. Cincinnati, OH 45214

CANADA MFD LABEL

ban

unscented
ROLL-ON

ANTIPERSPIRANT

DEODORANT

3.5 FL OZ (103 mL)

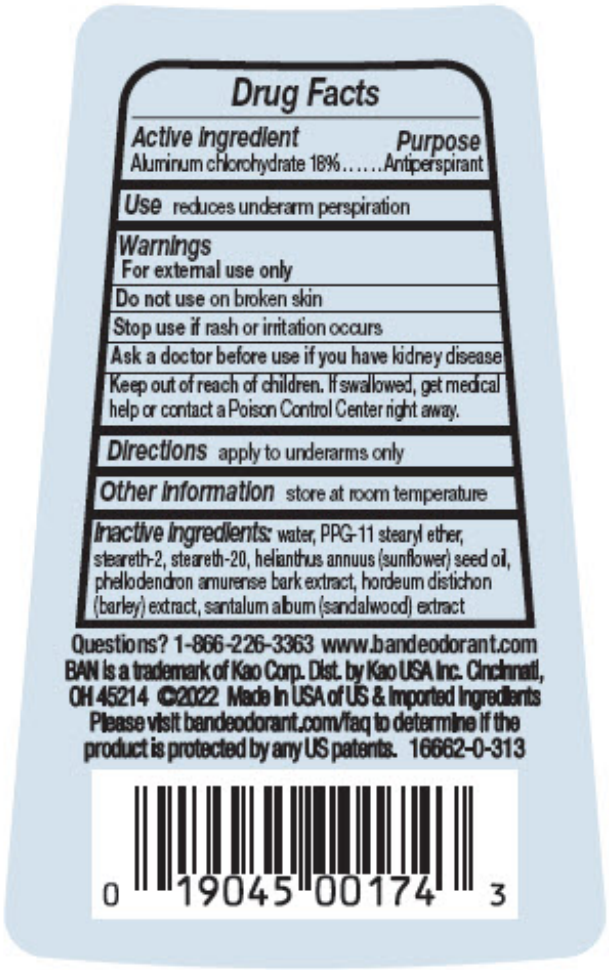
USA MANUFACTURED LABEL

ban

unscented
INVISIBLE ROLL-ON

antiperspirant deodorant

3.5 FL OZ (103 mL)



BAN ROLL-ON ANTIPERSPIRANT DEODORANT UNSCENTED

aluminum chlorohydrate liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10596-338
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)	
PPG-11 STEARYL ETHER (UNII: S4G2J0Y0LG)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10596-338-35	103 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	01/01/2012	
2	NDC:10596-338-70	2 in 1 PACKAGE	09/12/2016	
2		103 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		
3	NDC:10596-338-15	44 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	11/04/2016	
4	NDC:10596-338-11	3 in 1 PACKAGE	06/23/2020	
4		103 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		
5	NDC:10596-338-14	4 in 1 PACKAGE	08/24/2020	
5		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: 1/2025 Kao USA Inc.