URBAN STREET OCEAN BREEZE- ocean breeze roll on antiperspirant/deodorant stick
Blue Cross Laboratories LLC

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.

Active Ingredient

Aluminum Chloride 25.0%

Purpose

Antiperspirant

Use

Reduces underarm perspiration. Extra effective

Warnings

For external use only

Do not use on broken skin.

Stop use and ask a doctor if rash or irritation occurs

Keep out of reach of children. If swallowed, get medical help or contact poison control center right away.

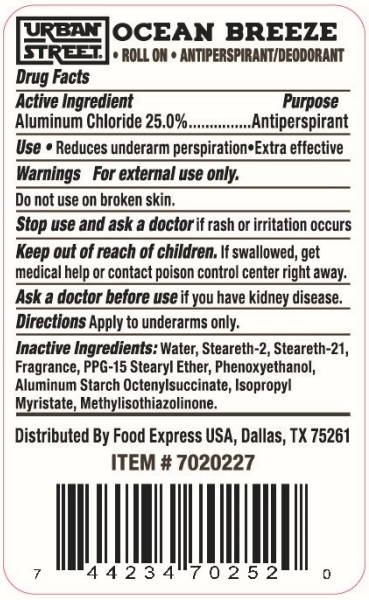
Ask a doctor before use if you have kidney disease.

Directions

Apply to underarms only.

Inactive Ingredients:

Water, Steareth-2, Steareth-31, Fragrance, PPF-15 Stearyl Ether, Phenoxyethanol, Aluminum Starch Octenylsuccinate, Isopropyl Myristate, Methylisothiazolinone



70*35 MM 35*57MM

Urban Street

Ocean Breeze

Roll On

Antiperspirant/Deodorant

3.25 fl. oz. (96.1 ml)

URBAN STREET OCEAN BREEZE

ocean breeze roll on antiperspirant/deodorant stick

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:22431-227
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLORIDE (UNII: 3CYT62D3GA) (ALUMINUM CATION - UNII:3XHB1D032B)	ALUMINUM CHLORIDE	250 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**

STEARETH-21 (UNII: 53J3F32P58)	
PPG-15 STEARYL ETHER (UNII: 1II18XLS1L)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)	
WATER (UNII: 059QF0KO0R)	
STEARETH-2 (UNII: V56DFE46J5)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

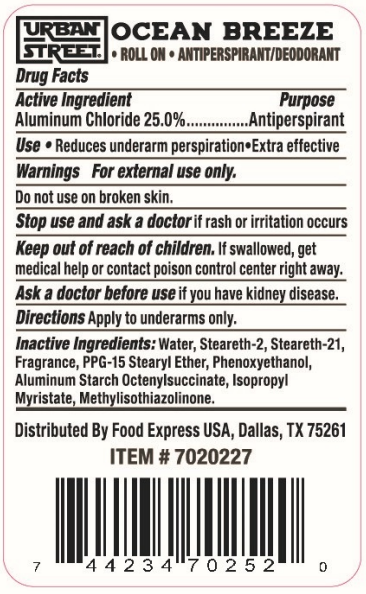
Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22431-227-01	9.6 g in 1 TUBE; Type 0: Not a Combination Product	07/19/2021	

70*35 MM



35*57MM

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	07/19/2021	

Labeler - Blue Cross Laboratories LLC (008298879)

Registrant - Blue Cross Laboratories LLC (008298879)

Establishment

Name	Address	ID/FEI	Business Operations
Blue Cross Laboratories		008298879	manufacture(22431-227)

Revised: 7/2021 Blue Cross Laboratories LLC