

**ODORONO ANTIPERSPIRANT AND DEODORANT ROLL-ON ORIGINAL- aluminum chlorohydrate liquid
Omega & Delta Co**

Odorono Antiperspirant Deodorant Roll-on Original

☐ *Drug Facts*

Active Ingredients

Aluminum Chlorohydrate 20%

Purpose

Antiperspirant

Use: Reduces underarm wetness

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.

Directions: Apply evenly to underarms only

Inactive Ingredients: Water, PPG-15 Stearyl Ether, Steareth-2, Steareth-21, Fragrance, Disodium EDTA, BHT, Triclosan

Hecho en Puerto Rico por:

☐ ***Omega & Delta Co., Inc.***

P.O. Box 1831, Carolina, P.R. 00984



Odorono[®]

Original

ANTI-PERSPIRANT
Deodorant

ALCOHOL FREE

2.5oz (74ml)

USO: Reduce la humedad en las axilas. **PRECAUCIONES:** Solo para uso externo. No usar sobre piel irritada o lastimada. Detenga su uso si presenta irritación o alguna molestia. Consulte a su médico antes de usar si padece de los riñones. Manténgase fuera del alcance de los niños. **INSTRUCCIONES DE USO:** Aplique únicamente en las axilas. **INGREDIENTE ACTIVO:** Clorhidrato de aluminio **OTROS INGREDIENTES:** Agua, PPG-15 Estearil Eter, Estearat-2, Estearat-21, Fragancia, EDTA disódico, BHT.

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Distributed by:
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Carolina, P.R. 00984
www.omegadeltaco.com





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aluminum chlorohydrate liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51048-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZ W13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZ W13M)	ALUMINUM CHLOROHYDRATE	20 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PPG-15 STEARYL ETHER (UNII: 1II18XLS1L)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	

EDETATE DISODIUM (UNII: 7FLD91C86K)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
TRICLOSAN (UNII: 4NM5039Y5X)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51048-201-25	74 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	05/08/2006	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	05/08/2006	

Labeler - Omega & Delta Co (090317793)

Establishment			
Name	Address	ID/FEI	Business Operations
Omega & Delta Co		090317793	manufacture(51048-201)

Revised: 11/2024

Omega & Delta Co