CARPE ANTIPERSPIRANT UNDERARM SANDALWOOD SCENT- aluminum sesquichlorohydrate lotion
Cosco International, Inc.

Drug Facts

Active Ingredient ***Purpose***
Aluminum Sesquicholorhydrate 15% .....Anti-perspirant

Anti-perspirant

Use
- Reduces underarm perspiration

Warnings

For external use only

Do not use
- on broken or irritated skin

Stop use and ask a doctor if
- rash or irritation occurs

Ask 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

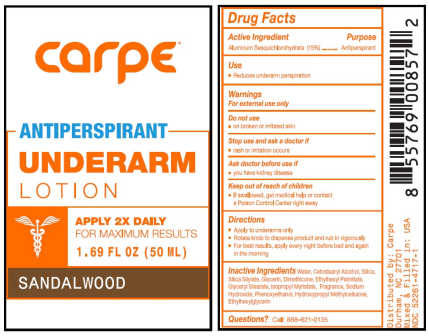
Directions
- Apply to underarms only
- Rotate knob to dispense product and rub in vigorously
- For best results, apply every night before bed and again in the morning

Inactive ingredients

Water, Cetostearyl Alcohol, Silica, Silica Silylate, Glycerin, Dimethicone, Ethylhexyl Palmitate, Glyceryl Stearate, Isopropyl Myristate, Fragrance, Sodium Hydroxide, Phenoxyethanol, Hydroxypropyl Methylcellulose, Ethylhexylglycerin

Questions?

Call 888-621-0135



CARPE ANTIPERSPIRANT UNDERARM SANDALWOOD SCENT

aluminum sesquichlorohydrate lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52261-4717
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Aluminum Sesquichlorohydrate (UNII: UCN889409V) (ALUMINUM SESQUICHLOROHYDRATE - UNII:UCN889409V)	Aluminum Sesquichlorohydrate	0.15 kg in 1 kg

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HEXAMETHYLDISILAZANE (UNII: H36C68P1BH)	
GLYCERIN (UNII: PDC6A3C0OX)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	

dimethicone 350 (UNII: 2Y53S6ATLU)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
phenoxyethanol (UNII: HIE492ZZ3T)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYPROMELLOSE 2208 (100000 MPA.S) (UNII: VM7F0B23ZI)	
ethylhexylglycerin (UNII: 147D247K3P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52261-4717-1	0.050 kg in 1 TUBE; Type 0: Not a Combination Product	05/02/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M019	04/27/2022	

Labeler - Cosco International, Inc. (016433141)

Registrant - Cosco International, Inc. (016433141)

Establishment

Name	Address	ID/FEI	Business Operations
Cosco International, Inc.		016433141	manufacture(52261-4717) , label(52261-4717) , pack(52261-4717)

Revised: 2/2025 Cosco International, Inc.