AQUAX DEO- aluminum chlorohydrate cream
Pella Pharmaceuticals Co. Ltd

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.

Deo

Forms and presentation

Cream: Tube of 75 g

Active Ingredient

Aluminum Chlorohydrate

Inactive Ingredients

Aqua, Cetearyl Alcohol, Ceteareth-22, Glycerin, Petrolatum, Ceteareth-25, Sodium Citrate, Methylparaben, Citric Acid, Parfum, Propylparaben.

Purpose

Antiperspirant

Properties

Aquax ® Deo cream reduces excessive sweating (hyperhidrosis)

Indications

Aquax ® Deo cream is used to reduce excessive perspiraton (hyperhidrosis)

Precautions

keep out of reach of children

Warnings

- For external use only
- Do not apply to broken or irritated skin
- Do not shave auxiliaries for 24 hours before application

Contraindications

Hypersensitivity to any of the components

Side effects

Aquax ® Deo cream has no known side effect, its us is without risk

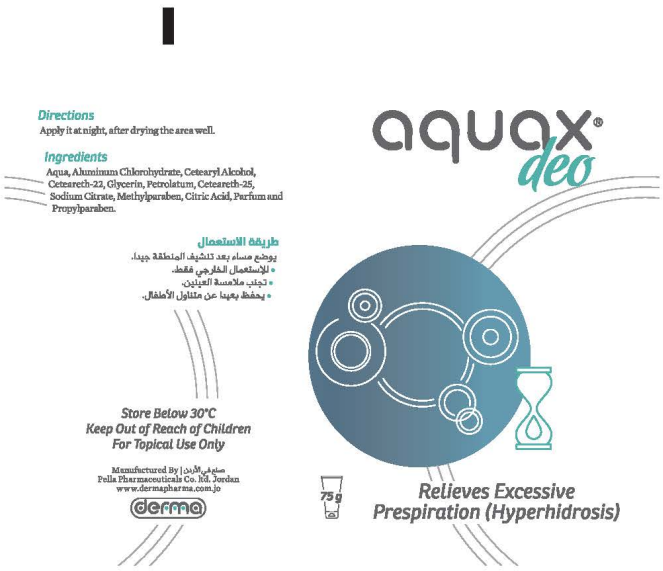
Dosage and administration

Dry the area to be treated, and then apply Aquax ® Deo cream at night, at bed time

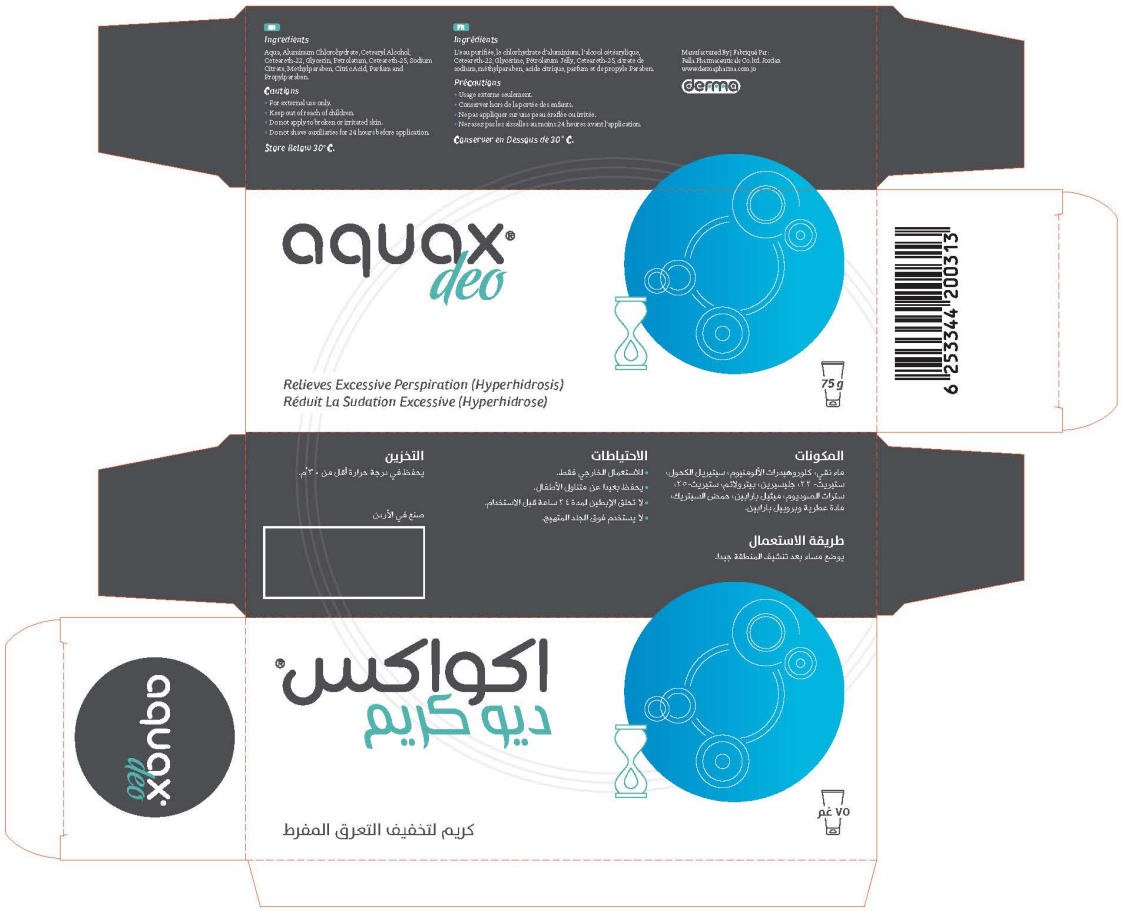
Storage conditions

Store at a temperature belwo 30 °C

Primary Package



Secondary Package



AQUAX DEO

aluminum chlorohydrate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82160-313
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALUMINUM CHLOROHYDRATE (UNII: HPN8MZW13M) (ALUMINUM CHLOROHYDRATE - UNII:HPN8MZW13M)	ALUMINUM CHLOROHYDRATE	150 mg in 75 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CETEARETH-22 (UNII: 28VZG1E234)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PETROLATUM (UNII: 4T6H12BN9U)	
CETEARETH-25 (UNII: 8FA93U5T67)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82160-313-01	1 in 1 CARTON	03/15/2011	
1		75 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part350	03/15/2011	

Labeler - Pella Pharmaceuticals Co. Ltd (562370925)

Revised: 11/2021 Pella Pharmaceuticals Co. Ltd