AQUAX WHITENING- oligopeptide-68 cream Pella Pharmaceuticals Co. Ltd

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Whitening

Forms and presentation

Cream: Tube of 50 g.

Active Ingredient

Oligopeptide-68

Inactive Ingredients

Aqua, Glycerin, Celyl Alcohol, Polysorbate 80, Cera Alba, Butylene Glycol, Xanthan Gum, Carbomer, Methylparaben, Triethanolamine, Propylparaben, Parfum, Hydrogenated Lecithin, Glycine Soja, Sodium Oleat, Disodium EDTA and BHA.

Purpose

Whitening

Properties

Aquax $^{\circledR}$ Whitening is a cream that moisturizes, nourishes and lightens skin.

Indications

- Whitening of skin.
- Reduction of spots due to sunlight exposure.
- Pregnancy Mask (Melasma).
- Reduction of secular pigmentations of acne.
- Can be used on sensitive areas (armpits, genital region, knees, elbows .. etc.)
- The Product is Safe and Effective.

Precautions

Keep out of reach of children

Warnings

- For external use only
- Avoid contact with the eyes
- If you are allergic to any of the ingredients listed, you should check with your doctor or pharmacist before you use the product.

Contraindications

Hypersensitivity to any of the components.

Side effects

Aquax ® Whitening Cream has no known side effect, because using it is safe.

Dosage and administration

Apply Aquax [®] Whitening Cream twice daily (morning and evening) on a cleaned skin.

Storage conditions

Store at a temperature below 30 ° C.

Primary Package



Secondary Package



AQUAX WHITENING

oligopeptide-68 cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82160-245
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
AMINO ACIDS (UNII: 0072R8RF8A) (AMINO ACIDS - UNII:0072R8RF8A)	AMINO ACIDS	0.2 mg in 50 g	

Inactive Ingredients		
	Ingredient Name	Strength

SOYBEAN OIL (UNII: 241ATL177A)	
WHITE WAX (UNII: 7G1J5DA97F)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
TROLAMINE (UNII: 903K93S3TK)	
SODIUM OLEATE (UNII: 399SL044HN)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
HYDROGENATED SOYBEAN LECITHIN (UNII: H1109Z9J4N)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
XANTHAN GUM (UNII: TTV12P4NEE)	
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82160-245- 01	1 in 1 CARTON	09/13/2010	
1		50 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
unapproved drug other		09/13/2010	

Labeler - Pella Pharmaceuticals Co. Ltd (562370925)

Revised: 12/2021 Pella Pharmaceuticals Co. Ltd