## SUBIR EYELASH- biotin cream Dream Polymer

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.

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## **Active Ingredient(s)**

Biotin 10%

D-Panthenol 8%

## **Purpose**

Anti-static, Hair conditioning, Skin conditioning

### Use

Subir Eyelash Biotin Cream for eyelash damage care

## Warnings

- Be careful not to get in your eyes.
- If it gets into your eyes, immediately wash it off with clean water several times.
- Wear appropriate gloves during the procedure.
- Do not eat.
- Keep out of reach of children.
- Store in a cool place away from direct sunlight.
- Recommended for single use only.
- For professional use only.
- Follow the recommendations.
- After use, if skin irritation such as rash or itching persists, stop use and consult a doctor or pharmacist

### **Directions**

- Using a Brush follow the same steps used for applying the perm and setting creams to applying the Essence (nourishing agent).
- Let nourishing set for five minutes.
- Prevents eyelash damage and promotes growth.

#### Other information

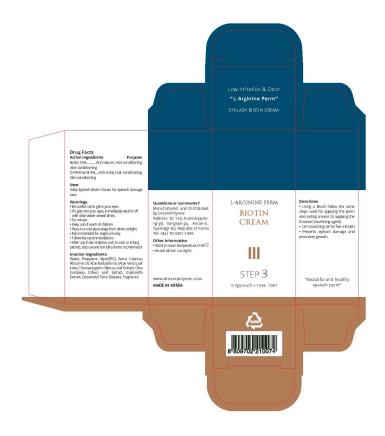
☐ Store in room temperature 0~30°C

☐ Avoid direct sunlight.

## **Inactive ingredients**

Water, Propylene Glycol(PG), Acrus Calamus Rhizome Oil, Aloe Barbadensis (Aloe Vera) Leaf Juice, Chamaecyparis Obtusa Leaf Extract, Olea Europaea (Olive) Leaf Extract, Espinosilla Extract, Decametyl Tetra Siloxane, Fragrance

## **Package Label - Principal Display Panel**



SUBIR EYELASH biotin cream				
Product Information				
Product Type	HUMAN OTC DRUG	Item C	ode (Source)	NDC:81716-004
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength

Inactive Ingredients			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
Propylene Glycol (UNII: 6DC9Q167V3)			
OLEA EUROPAEA LEAF (UNII: MJ95C3OH47)			
CHAMAECYPARIS OBTUSA LEAF (UNII: 70L154J5XB)			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:81716-004- 01	0.7 g in 1 POUCH; Type 0: Not a Combination Product	03/30/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/30/2021	

# Labeler - Dream Polymer (695501990)

# Registrant - Dream Polymer (695501990)

Establishment				
Name	Address	ID/FEI	Business Operations	
Dream Polymer		695501990	manufacture(81716-004)	

Revised: 3/2021 Dream Polymer