SUBIR EYELASH- setting 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.

Active Ingredient(s)

Sodium Bromate 12%

Purpose

For permanent wave

Use

Subir Eyelash Setting Cream for eyelash permanent wave

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

 Keep out of reach of children.

Directions

• After desired time has passed, remove perm cream (step1). Use a newly moistened cotton pad to remove all traces of perm cream (step1)
• Follow the same application technique to apply setting cream(step2), as you did to (step1).
• After removing the perm cream, apply the setting cream and after 10 ~ 12 minutes, remove the set cream using a newly moistened cotton pad (depending on the texture of the lashes).

Other information

 Store in room temperature 0~30℃
 Avoid direct sunlight.

Inactive ingredients

Water, PolyPropyleneGlycol-7, Cetearyl Alcohol, Mineral Oil, Cetyl Alcohol, Stearyl Alcohol

Package Label - Principal Display Panel

SUBIR EYELASH

setting cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81716-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
STRONTIUM BROMATE (UNII: 1T54WJB20V) (STRONTIUM BROMATE -	STRONTIUM	0.084 g

UNII:1T54WJB20V)	BROMATE	in 0.7 g

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Cetyl Alcohol (UNII: 936JST6JCN)	
Stearyl Alcohol (UNII: 2KR89I4H1Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81716-003-01	0.7 g in 1 POUCH; Type 0: Not a Combination Product	03/31/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/31/2021	

Labeler - Dream Polymer (695501990)

Registrant - Dream Polymer (695501990)

Establishment

Name	Address	ID/FEI	Business Operations
Dream Polymer		695501990	manufacture(81716-003)

Revised: 3/2021 Dream Polymer