AAPE HAIR AMPOULE- panthenol liquid PROSTEMICS 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.

ACTIVE INGREDIENT

Panthenol 0.5%

INACTIVE INGREDIENT

[Powder] MANNITOL, Human Adipose Derived Mesenchymal Cell Exosomes.

[Solvent] Water, Alcohol, Propylene Glycol, Glycerin, Lactic Acid, Hydroxyacetophenone, Propanediol, Polyquaternium-7, Niacinamide, Polysorbate 20, Butylene Glycol, Caprylhydroxamic Acid, Disodium EDTA, Glycyrrhiza Glabra (Licorice) Root Extract, Camellia Sinensis Leaf Extract, Panax Ginseng Root Extract, Sophora Angustifolia Root Extract, Angelica Gigas Root Extract, Cnidium Officinale Root Extract, Glycine Max (Soybean) Seed Extract, Polygonum Multiflorum Root Extract, 1,2-Hexanediol, Sodium Benzoate

PURPOSE

Hair elasticity

WARNINGS

For external use only

- 1. Discontinue use if signs of irritation or rashes appear. If symptoms get worse, consult with a dermatologist. 1) In case of swelling, itching, or other side effects while or after using this product
- 2. Do not apply to open wounds.
- 3. Avoid contact with eyes.

Storage and handling

- 4. Keep in the refrigerator at 2-6°C.
- 5. Keep out of reach of children.
- 6. Avoid direct sunlight.

KEEP OUT OF REACH OF CHILDREN

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Uses

■ Improves effects of hair elasticity.

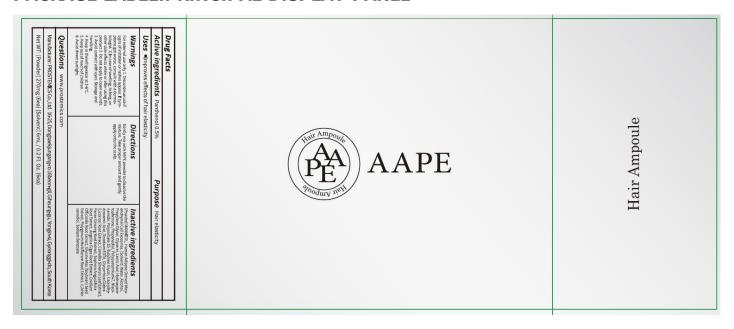
Directions

Gently mix with AAPE powder to dissolve the mixture. Take proper amount and gently apply onto the scalp

QUESTIONS

www.prostemics.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



AAPE HAIR AMPOULE panthenol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62041-330	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
Panthenol (UNII: WW9CM0O67Z) (PANTHENOL - UNII:WW9CM0O67Z)	Panthenol	0.5 g in 100 mL

Inactive Ingredients				
Ingredient Name	Strength			
Water (UNII: 059QF0KO0R)				
Alcohol (UNII: 3K9958V90M)				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62041- 330-02	6 in 1 CARTON	05/01/2021	
1	NDC:62041- 330-01	6 mL in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information				
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date	
unapproved drug other		05/01/2021		

Labeler - PROSTEMICS Co., Ltd. (689605919)

Registrant - PROSTEMICS Co., Ltd. (689605919)

Establishment				
Name	Address	ID/FEI	Business Operations	
Prostemics Co., Ltd. Factory		695687674	manufacture(62041-330)	

Revised: 5/2021 PROSTEMICS Co., Ltd.