

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

<p>Drug Facts</p> <p>Active Ingredients Panthenol 0.5%</p> <p>Uses Improves effects of hair elasticity.</p> <p>Warnings</p> <p>For external use only. 1. Discontinue use if you experience irritation. 2. Do not use if you are allergic to any of the ingredients. 3. Do not use if you are pregnant or breastfeeding. 4. Do not use if you have a known hypersensitivity to any of the ingredients. 5. Do not use if you have a known hypersensitivity to any of the ingredients. 6. Do not use if you have a known hypersensitivity to any of the ingredients. 7. Do not use if you have a known hypersensitivity to any of the ingredients. 8. Do not use if you have a known hypersensitivity to any of the ingredients. 9. Do not use if you have a known hypersensitivity to any of the ingredients. 10. Do not use if you have a known hypersensitivity to any of the ingredients.</p> <p>Directions</p> <p>Gently mix with AAPE powder to dissolve the mixture. Take proper amount and gently apply onto the scalp.</p> <p>Inactive Ingredients</p> <p>Propylene Glycol, Glycerin, Sodium Lauryl Sulfate, Hydroxyethyl Cellulose, Polysorbate 20, Polysorbate 80, Polysorbate 40, Polysorbate 60, Polysorbate 120, Polysorbate 140, Polysorbate 160, Polysorbate 180, Polysorbate 200, Polysorbate 220, Polysorbate 240, Polysorbate 260, Polysorbate 280, Polysorbate 300, Polysorbate 320, Polysorbate 340, Polysorbate 360, Polysorbate 380, Polysorbate 400, Polysorbate 420, Polysorbate 440, Polysorbate 460, Polysorbate 480, Polysorbate 500, Polysorbate 520, Polysorbate 540, Polysorbate 560, Polysorbate 580, Polysorbate 600, Polysorbate 620, Polysorbate 640, Polysorbate 660, Polysorbate 680, Polysorbate 700, Polysorbate 720, Polysorbate 740, Polysorbate 760, Polysorbate 780, Polysorbate 800, Polysorbate 820, Polysorbate 840, Polysorbate 860, Polysorbate 880, Polysorbate 900, Polysorbate 920, Polysorbate 940, Polysorbate 960, Polysorbate 980, Polysorbate 1000.</p> <p>Questions www.prostemics.com</p> <p>Manufacturer: PROSTEMICS Co., Ltd. 15-25, Donggok-ro, Gyeonggi-do, Gyeonggi-do, South Korea</p> <p>Net Wt. (Powder) 270mg (5ea) (Solvent) 6ml / 0.21 Oz. (5ea)</p>	 <p>AAPE</p>	<p>Hair Ampoule</p>
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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: W9CM0067Z) (PANTHENOL - UNII:W9CM0067Z)	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 Category	Application Number or Monograph 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.