**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.*

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## ACTIVE INGREDIENT

Active ingredients: Panthenol 0.5%

## INACTIVE INGREDIENT

Inactive ingredients:
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

Purpose: Hair elasticity

## WARNINGS

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

KEEP OUT OF REACH OF CHILDREN

### Uses

Uses:

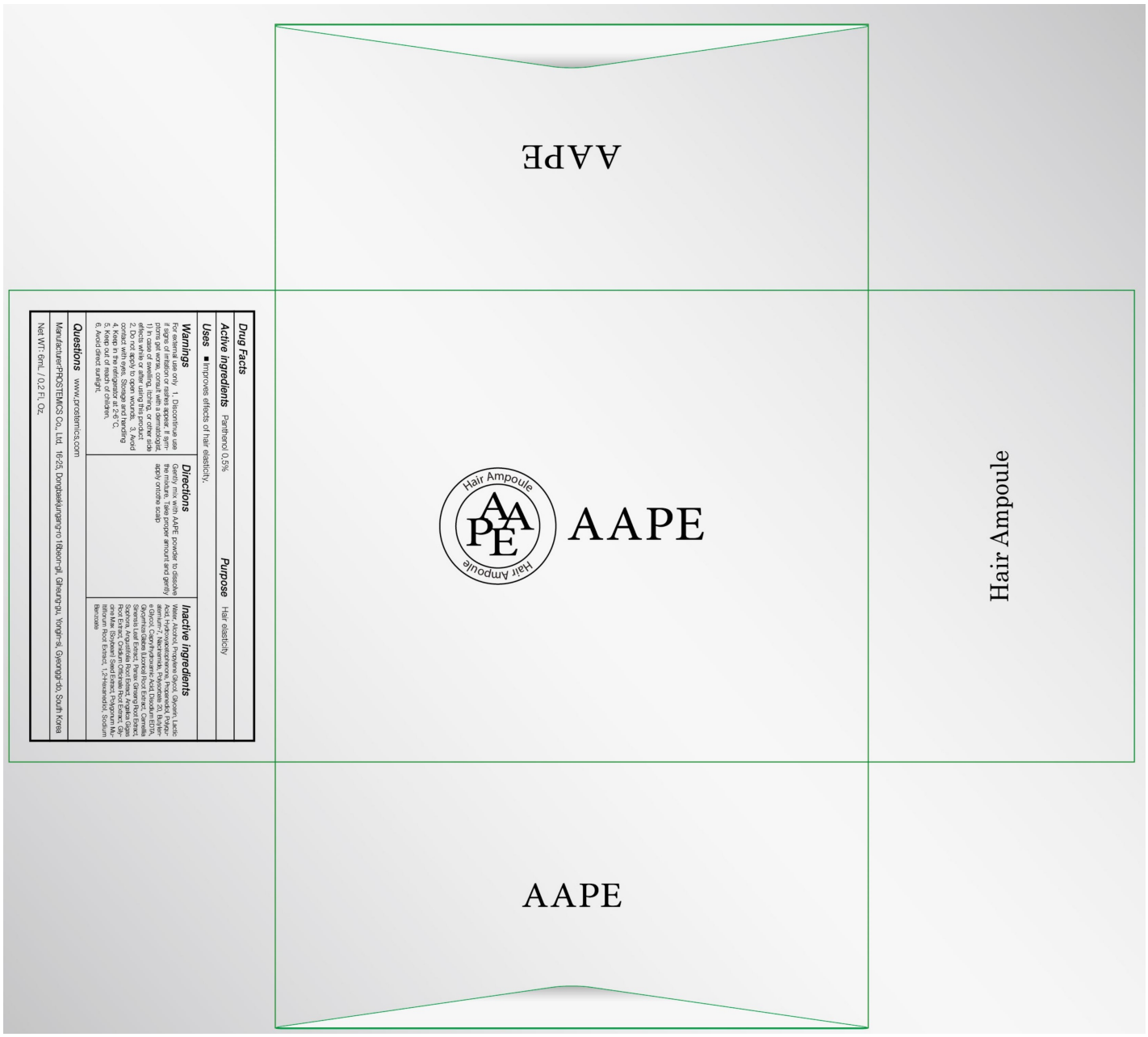
Improves effects of hair elasticity.

### Directions

Directions:

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

## PACKAGE LABEL.PRINCIPAL DISPLAY PANEL




# AAPE HAIR AMPOULE

panthenol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:62041-260 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **Panthenol** (UNII: WV9CM0O67Z) (PANTHENOL - UNII:WV9CM0O67Z) | Panthenol | 0.03 g in 6 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:62041-260-01 | 6 mL in 1 CONTAINER; Type 0: Not a Combination Product | 08/01/2019 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 08/01/2019 | |

## Labeler - PROSTEMICS Co., Ltd. (689605919)

## Registrant - PROSTEMICS Co., Ltd. (689605919)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Prostemics Co., Ltd. Factory | | 695687674 | manufacture(62041-260) |

Revised: 8/2019 PROSTEMICS Co., Ltd.