AAPE SKIN AMPOULE- niacinamide 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

Niacinamide 2.00%

INACTIVE INGREDIENT

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

[Solvent] Water, Butylene Glycol, Glycerin, Sorbitol, Hydroxyacetophenone, Propanediol, Xanthan Gum, Allantoin, Adenosine, Caprylhydroxamic Acid, Glycyrrhiza Glabra (Licorice) Root Extract, Camellia Sinensis Leaf Extract, Disodium EDTA, Sophora Angustifolia Root Extract, Angelica Gigas Root Extract, Panax Ginseng Root Extract, Cnidium Officinale Root Extract, Glycine Max (Soybean) Seed Extract, Polygonum Multiflorum Root Extract, 1,2-Hexanediol, Caprylyl Glycol, Palmitoyl Tripeptide-5

PURPOSE

Skin Brightening

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

Helps brighten skin tone.

Directions

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

QUESTIONS

www.prostemics.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

AAPE SKIN AMPOULE

niacinamide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62041-320
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Niacinamide (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	Niacinamide	2.0 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Butylene Glycol (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62041-320-02	6 in 1 CARTON	05/01/2021	
1	NDC:62041-320-01	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(62041-320)

Revised: 5/2021 PROSTEMICS Co., Ltd.