AAPE NUTRIENT FACIAL TONER- 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

Active ingredients: Niacinamide 2.00%

INACTIVE INGREDIENT

Inactive ingredients:
Water, Butylene Glycol, PEG/PPG-17/6 Copolymer, Trehalose, Human Adipocyte Conditioned Media Extract, 1,2-Hexanediol, Caprylyl Glycol, Bis-PEG-18 Methyl Ether Dimethyl Silane, Ethylhexylglycerin, Salix Alba (Willow) Bark Extract, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Tromethamine, Glycerin, Cellulose Gum, Panthenol, PEG-60 Hydrogenated Castor Oil, Hamamelis Virginiana (Witch Hazel) Leaf Water, Disodium EDTA, Glyceryl Citrate/Lactate/Linoleate/Oleate, Sodium Hyaluronate, Olea Europaea (Olive) Fruit Oil, Polysorbate 60, Lilium Candidum, Flower Extract, Leontopodium Alpinum Flower/Leaf Extract, Convallaria Majalis Bulb/Root Extract, Magnolia Liliflora Flower Extract, Paeonia Lactiflora Extract, Palmitoyl Tripeptide-5, Fragrance

PURPOSE

Purpose: Skin Brightening

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. Replace the cap after use
5. Keep out of reach of children.
6. Avoid direct sunlight.

KEEP OUT OF REACH OF CHILDREN

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Uses

Uses:

Helps brighten skin tone.

Directions

Directions:

Take proper amount and gently apply onto the face.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

AAPE®

Nutrient
Facial Toner

ANTI-AGING
CONDITIONED MEDIA

8 809246 215016

Drug Facts

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Questions www.prostemics.com

Manufacturer: PROSTEMICS Co., Ltd.
16-25, Dongbaekjungang-ro 16beon-gil, Giheung-gu, Yongin-si, Gyeonggi-do, South Korea

Net WT:
130mL / 4.39 Fl.Oz

AAPE NUTRIENT FACIAL TONER

niacinamide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62041-200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Niacinamide (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	Niacinamide	2.60 g in 130 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:62041-200-02	1 in 1 CARTON	08/01/2019	
1	NDC:62041-200-01	130 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		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(62041-200)

Revised: 8/2019 PROSTEMICS Co., Ltd.