MORINGA PURIFY WATER- niacinamide, adenosine cream
Oneface 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 2.0%, Adenosine 0.04%

INACTIVE INGREDIENT

Inactive ingredients:

Moringa Oleifera Seed Extract, Glycerin, Cyclopentasiloxane, Cyclohexasiloxane, Butylene Glycol, Methyl Hydrogenated Rosinate, Cetyl PEG/PPG-10/1 Dimethicone, Sodium Hyaluronate, Beta-Glucan, 1,2-Hexanediol, Caprylhydroxamic Acid, Propanediol, Sodium Chloride, Titanium Dioxide, Aluminum Hydroxide, Triethoxycaprylylsilan, Cyclomethicone, PEG-10 Dimethicone, Dimethicone, Dimethicone/Vinyl Dimethicone, Crosspolymer, Fragrance, Water, Caulerpa Lentillifera Extract, Codium Fragile Extract, Donkey Milk, Honey Extract, Goat Milk Extract, Phenoxyethanol, Ethylhexyglycerin, Saccharide Isomerate, Citric Acid, Sodium Citrate, Potassium PCA, Opuntia Ficus Indica Stem Extract

PURPOSE

Purpose: Skin Brightening, Anti-wrinkle

WARNINGS

Warnings:

For external use only

Avoid contact with eyes.

Discontinue use if signs of irritation or rashes appear.

Replace the cap after use.

Keep out of reach of children

Keep out of High, Low temperature.

KEEP OUT OF REACH OF CHILDREN

KEEP OUT OF REACH OF CHILDREN

Uses

Uses:

Helps brighten skin. Helps anti wrinkle.

Directions

Directions:

1. Place desired amount into palm, and rub softly on your face skin
2. Massage gently over skin

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

MORINGA PURIFY WATER

niacinamide, adenosine cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72255-010
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
Adenosine (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	Adenosine	0.04 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Glycerin (UNII: PDC6A3C0OX)	
Butylene Glycol (UNII: 3XUS85K0RA)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72255-010-02	1 in 1 CARTON	04/01/2018	
1	NDC:72255-010-01	100 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		04/01/2018	

Labeler - Oneface co., Ltd. (688219085)

Registrant - Oneface co., Ltd. (688219085)

Establishment

Name	Address	ID/FEI	Business Operations
Oneface co., Ltd.		688219085	manufacture(72255-010)

Revised: 5/2018 Oneface co., Ltd.