

ULTRA WHITENING- niacinamide cream
MIGUHARA

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 Ingredient: Niacinamide 2.0%

INACTIVE INGREDIENT

Inactive Ingredients: Camellia Sinensis Leaf Extract, Glycerin, Ethylhexyl Palmitate Ethylhexyl Palmitate, Dimethicone/Vinyl Dimethicone Crosspolymer /Cyclomethicone/Hydrogenated Lecithin/Alcohol/Aqua, Helianthus Annuus (Sunflower) Seed Oil, Allium Cepa (Onion) Bulb Extract, Beta-Glucan, Ceramide/Phospholipids/Phytosterols/Soybean Glycerides/ Stearyl Alcohol /Behenyl Alcohol, Cetearyl Alcohol, Cyclomethicone, 1,2-Hexanediol, Panthenol, Urtica Dioica (Nettle) Extract, Argania Spinosa Kernel Oil, Sodium Hyaluronate, Beeswax, Butyrospermum Parkii (Shea) Butter, Stearic Acid, Dimethicone, Ecklonia Cava Extract, Tocopheryl Acetate, Arginine, Carbomer, Palmitoyl Dipeptide-7, Xanthan Gum, Allantoin, Disodium EDTA, Citrus Paradisi (Grapefruit) Peel Oil

PURPOSE

Purpose: Skin Protectant

WARNINGS

Warnings:

1. Stop usage immediately if any of the below symptoms occur. Continued use could aggravate symptoms, so it is advisable to consult with a dermatologist immediately. 1) Symptoms include but not limited to: red spots, swelling, itchiness. 2) When having the same symptoms as above due to direct sunlight. 2. Do not apply to areas affected by dermatitis, eczema or wounds. 3. Storage and handling: 1) Tightly close lid after each use. 2 Keep out of reach of children 3) Store in a cool dry area, away from sunlight 4. According to supporting evidence that prove its effect. It causes slight papular and uredo, when applying the same contained ingredient medicine.

KEEP OUT OF REACH OF CHILDREN

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Usage

Usage: Take an appropriate amount and gently apply it on the face

Usage

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PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

MIGUHARA

Ultra whitening cream

W-Ctrl

Ultra Whitening Cream helps getting rid of oxygen radicals and free radicals generated by ultraviolet ray exposure from everyday activities and inhibits the activation of melanin pigment to keep your skin whiter and cleaner. Extracts from onion, nettle and ecklonia are especially effective in protecting your skin from harmful ultraviolet exposure from everyday activities.

50 ml

[Active Ingredient] Niacinamide 2.0%

[Inactive Ingredients] Camellia Sinesis Leaf Extract, Glycerin, Ethylhexyl Palmitate/Ethylhexyl Palmitate, Dimethicone/Vinyl Dimethicone Crosspolymer /Cyclomethicone/Hydrogenated Lecithin/Alcohol/Aqua, Helianthus Annuus (Sunflower) Seed Oil, Allium Cepa (Onion) Bulb Extract, Beta -Glucan, Ceramide/Phospholipids/Phytosterols/Soybean Glycerides / Stearyl Alcohol /Behenyl Alcohol, Cetearyl Alcohol, Cyclomethicone, 1,2-Hexanediol, Panthenol, Urtica Dioica (Nettle) Extract, Argania Spinosa Kernel Oil, Sodium Hyaluronate, Beeswax, Butyrospermum Parkii (Shea) Butter, Stearic Acid, Dimethicone, Ecklonia Cava Extract, Tocopheryl Acetate, Arginine, Carbomer, Palmitoyl Dipeptide-7, Xanthan Gum, Alanitol, Disodium EDTA, Citrus Paradisi (Grapefruit) Peel Oil

[Purpose] Skin Protectant

[Warnings]

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Ultra Whitening Ample

[Usage]

Take an appropriate amount and gently apply it on the face.

- **Manufacturer and Distributor** : Miguhara Inc.
- **Manufacturer** : Beautist Inc.

If the product has any defects,
we will compensate with you
by following all regulations conducted
by fair trade commission.
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MIGUHARA COSMETICS
MADE IN KOREA



ULTRA WHITENING

niacinamide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70380-340
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Niacinamide (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	Niacinamide	1.0 g in 50 mL

Inactive Ingredients

Ingredient Name	Strength
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
Glycerin (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70380-340-01	50 mL in 1 CARTON; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/02/2016	

Labeler - MIGUHARA (689204213)**Registrant** - MIGUHARA (689204213)**Establishment**

Name	Address	ID/FEI	Business Operations
MIGUHARA		689204213	manufacture(70380-340)