

KARATICA I M CURE- niacinamide patch

Karatica 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](#).

niacinamide

SodiumHyaluronate, Niacinamide, CamelliaJaponicaFlowerExtract, CentellaAsiaticaExtract, Tranexamicacid, SalicylicAcid, TetraPeptide-44, Sh-Oligopeptide-1, Sh-Polypeptide-1

skin soothing and care

keep out or reach of the children

Clean and dry the face. Apply the patch on the troubled area and press it gently for 10 seconds. Leave the patch on for 2-3 hours.

* The Essence Chip (micro projections) must be kept dry.

- 1) Discontinue use or consult dermatologist if irritation, itching or swelling appears after applying the product.
- 2) Avoid applying patches on the wounded, inflamed skin parts.
- 3) Cautions for the storage and handling.
 - Keep out of children.
 - Avoid applying around eyes.

for external use only

Karatica I'm Cure Patch

► How to Use

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► Ingredients:

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Tranexamicacid, SalicylicAcid, TetraPeptide-44, Sh-Oligopeptide-1, Sh-Polypeptide-1

► Cautions

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2) Avoid applying patches on the wounded, inflamed skin parts.

3) Cautions for the storage and handling.

- Keep out of children.

- Avoid applying around eyes.

► Content(Nedle patch)5mg X 6ea / (Hydrocolloid patch)6ea

► Manufacturer:EldisysCo., Ltd

119, Yulbuk-ro, Cheongbuk-myeon, Pyeongtaek-si, Gyeonggi-do, Republic of Korea

► Distributor:Karatica Co., Ltd

3F, 6-14, Bongeunsa-ro 68-gil, Gangnam-gu, Seoul, Republic of Korea

► **MFD & EXP:** Printed Separately

KARATICA I M CURE

niacinamide patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70514-0012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)		NIACINAMIDE	2.7 mg in 100 mg	
Inactive Ingredients				
Ingredient Name		Strength		
HYALURONATE SODIUM (UNII: YSE9PPT4TH)				
WATER (UNII: 059QF0K00R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70514-0012-1	6 in 1 PACKAGE	12/05/2018	
1		5 mg in 1 PATCH; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		12/05/2018		

Labeler - Karatica Co., Ltd (689605545)

Registrant - Karatica Co., Ltd (689605545)

Establishment

Name	Address	ID/FEI	Business Operations
Karatica Co., Ltd		689605545	manufacture(70514-0012) , label(70514-0012) , pack(70514-0012)

Revised: 12/2018

Karatica Co., Ltd