

**SISEUNDEUSI LUMINANT MASK PACK STEP2(WHITENING)- niacinamide cream
GK COSMETIC 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.

Drug Facts

Niacinamide

Aloe Barbadensis Leaf Water, Butylene Glycol, Etc

Skin Protectant - Whitening

1. After wash your face, take our STEP01 Luminant Umbrella Facial Mask and put it on your face.
2. Take off the sheet after 15 to 20 minutes when it gets absorbed into your skin
3. Tap your skin softly for your skin to absorb anything left on the skin.
4. Put STEP02 Luminant Whitening Aqua Creme(2 ml) evenly on your face. When we consider skin turn over period, using the product consistently for 4 weeks is the most effective way to use the product.

keep out of reach of the children

for external use only

1. Do not use in the following cases(Eczema and scalp wounds)

2.Side Effects

1)Due to the use of this drug if rash, irritation, itching and symptoms of hypersensitivity occur discontinue use and consult your pharmacist or doctor

3.General Precautions

1)If in contact with the eyes, wash out thoroughly with water If the symptoms are severe, seek medical advice immediately

2)This product is for external use only. Do not use for internal use

4.Storage and handling precautions

1)If possible, avoid direct sunlight and store in cool and area of low humidity

2)In order to maintain the quality of the product and avoid misuse

3)Avoid placing the product near fire and store out in reach of children



SISEUNDEUSI LUMINANT MASK PACK STEP2(WHITENING)

niacinamide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70098-0010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X5118 RD4) (NIACINAMIDE - UNII:25X5118 RD4)	NIACINAMIDE	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70098-0010-1	2 mL in 1 PACKAGE; Type 0: Not a Combination Product	03/09/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/09/2017	

Labeler - GK COSMETIC CO., LTD. (688477309)

Registrant - GK COSMETIC CO., LTD. (688477309)

Establishment

Name	Address	ID/FEI	Business Operations
GK COSMETIC CO., LTD.		688477309	manufacture(70098-0010) , label(70098-0010) , pack(70098-0010)

Revised: 3/2017

GK COSMETIC CO., LTD.