DR. GLODERM TIME TO LIFTUP MASK- niacinamide, adenosine liquid DR.GLODERM

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, Adenosine

Water, Glycerin, Butylene Glycol, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Sodium Hyaluronate, Centella Asiatica Extract, Sodium DNA, etc.

Skin Protectant - Whitening, Anti-Wrinkle

keep out of reach of the children

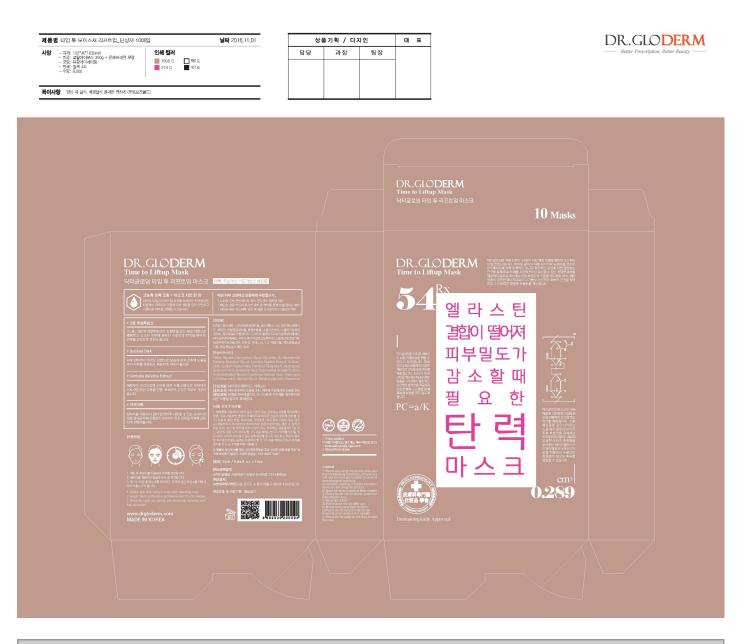
- 1. Refine skin with using a toner after cleansing face.
- 2. Apply mask on the face and take a rest 10-20 minutes.
- 3. Remove mask and gently pat remaining essence until fully absorbed.

1. If the following symptoms occur after product use, stop using the product immediately and consult a dermatologist (continuous use can exacerbate the symptoms).

1) Occurrence of red spots, swelling, itchiness, and other skin irritation

- 2) If the symptoms above occur after the application area is exposed to direct sunlight
- 2. Do not use on open wounds, eczema, and other skin irritations
- 3. Precaution for Storage and Handling
- 1) Close the lid after use
- 2) Keep out of reach of infants and children
- 3) Do not to store in a place with high/low temperature and exposed to direct sunlight
- 4. Use as avoiding eye areas.

for external use only



DR. GLODERM TIME TO LIFTUP MASK

niacinamide, adenosine liquid

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Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:71342-0001		
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ing	Basis of Streng	th Strength				
ADENO SINE (UNII: K72T3FS567) (ADENO SINE - UNII:K72T3FS567) ADENO SINE				$0.04\;g$ in 100 mL		
NIACINAMIDE (UNII: 25X5118RD4) (NIACINAMIDE - UNII:25X5118RD4) NIACINAMIDE				2 g in 100 mL		
Inactive Ingredients						
Ingredient Name			Strength			

BU	TYLENE GLYCOL	(UNII: 3XUS85K0RA)				
W	ATER (UNII: 059QF	OKOOR)				
Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:71342-0001-1		10 in 1 PACKAGE	0 3/0 1/20 17			
1		25 mL in 1 POUCH; Type 0: Not a Combination Product				
Marketing Information						
Μ	arketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
una	approved drug other		0 3/0 1/20 17			

Labeler - DR.GLODERM (694773267)

Registrant - DR.GLODERM (694773267)

Establishment

Name	Address	ID/FEI	Business Operations
KOREA DERMAL RESEARCH CENTER CO., LTD.		694640098	manufacture(71342-0001)

Establishment

Name	Address	ID/FEI	Business Operations
DR.GLODERM		694773267	label(71342-0001)

Revised: 3/2017

DR.GLODERM