DR. GLODERM TABRX WHITENING- niacinamide cream
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

Water, Butylene Glycol, Glycerin etc.

Skin Protectant - Whitening

keep out of reach of the children

It's melting on the pam taken two capsules with spatula at the last stage of skin care, and apply on the entire face.

Gently pat to help absorption into the skin.

* Do not eat capsules.

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

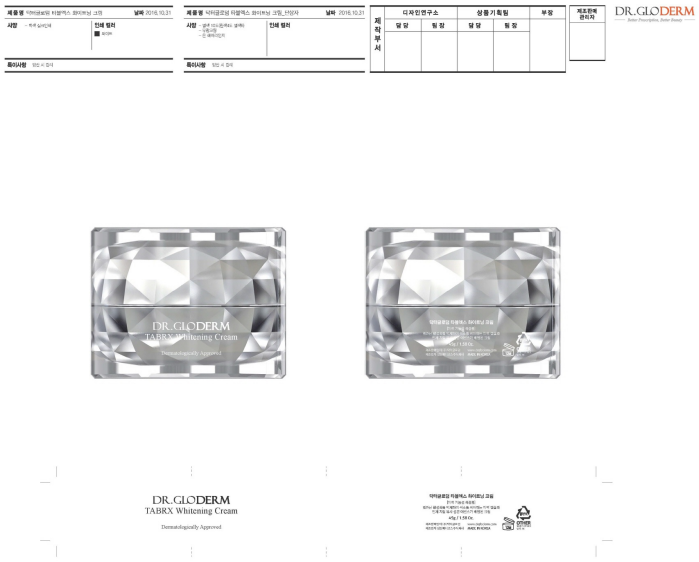
1) Close the lid after use

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 TABRX WHITENING

niacinamide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71342-0012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2.14 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71342-0012-1	45 g in 1 JAR; Type 0: Not a Combination Product	03/28/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/01/2017	

Labeler - DR.GLODERM (694773267)

Registrant - DR.GLODERM (694773267)

Establishment

Name	Address	ID/FEI	Business Operations
SAMSUNG MEDICOS. CO., LTD. Hyangnam Factory		689851701	manufacture(71342-0012)

Establishment

Name	Address	ID/FEI	Business Operations
DR.GLODERM		694773267	label(71342-0012)

Revised: 3/2017 DR.GLODERM