DR. GLODERM TIME TO WHITENING MASK- niacinamide 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

Water, Glycerin, Sodium Hyaluronate, Butylene Glycol, Pearl Extract, Hydrolyzed Hyaluronic Acid, Hyaluronic Acid, Glutathione, etc.

Skin Protectant - Whitening

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

제품명 타임 투 화이트닝 마스크_단상자 10매입 **날짜** 2016.11.01

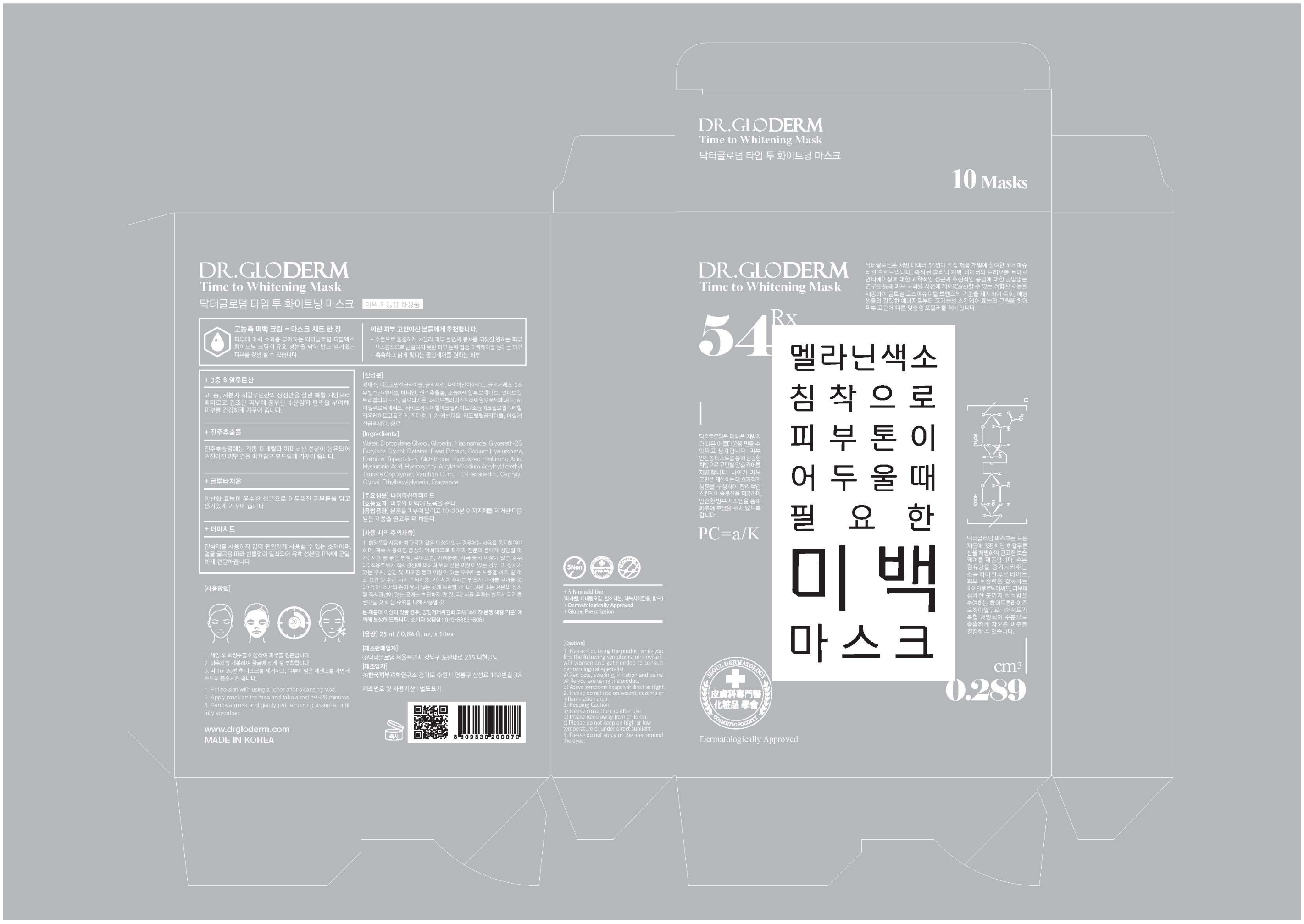
사양
- 규격: 122*40*183(mm)
- 지류: 로얄아이보리 350g + 은헤어라인 무광
- 코팅: 유광라미네이팅
- 인쇄: 별색 3도
- 수량: 8,000

인쇄 컬러
연그레이
백1도
먹1도

특이사항 양산 시 감리_배경컬러 최대한 연하게 (투명그레이)

상품기획 / 디자인			대 표
담당	과장	팀장	

DR.GLODERM
Better Prescription, Better Beauty



DR. GLODERM TIME TO WHITENING MASK

niacinamide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71342-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71342-0003-1	10 in 1 PACKAGE	03/01/2017	
1		25 mL in 1 POUCH; Type 0: Not a Combination Product		

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
KOREA DERMAL RESEARCH CENTER CO., LTD.		694640098	manufacture(71342-0003)

Establishment

Name	Address	ID/FEI	Business Operations
DR.GLODERM		694773267	label(71342-0003)

Revised: 3/2017 DR.GLODERM