LUMINANT UMBRELLA MASK PACK- allantoin liquid
GK COSMETIC CO., LTD.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

allantoin

glycerin, butylene glycol, etc.

skin protectant

keep out of reach of the children

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.

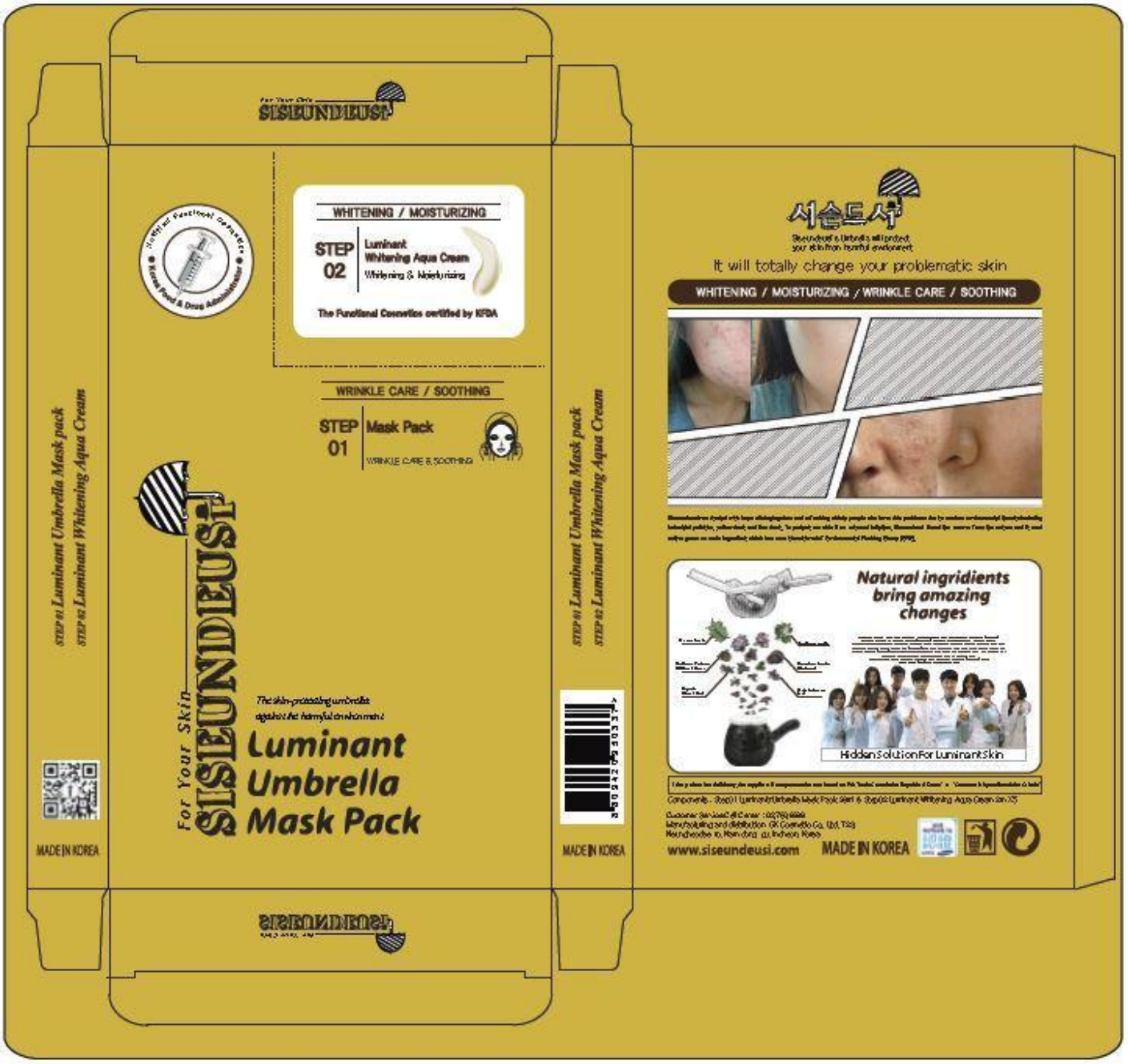
4. Put STEP02 Luminant Whitening Aqua Creme(2 ml) evenly on your face.

When we consider skin turn over period, using the productconsistently for

4 weeks is the most effective way to use the product.

1. Do not use in the following cases(Eczema and scalp wounds)
2.Side Effects
1)Due to the use of this druf if rash, irritation, itching and symptopms of hypersnesitivity occur dicontinue use and consult your phamacisr or doctor
3.General Precautions
1)If in contact with the eyes, wash out thoroughty with water If the symptoms are servere, seek medical advice immediately
2)This product is for exeternal 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

for external use only



LUMINANT UMBRELLA MASK PACK

allantoin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70098-0002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.1 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70098-0002-1	38 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/06/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	01/01/2016	

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(70098-0002)

Revised: 12/2019 GK COSMETIC CO., LTD.