HELLO CARE MOISTURIZING AMPOULE MASK- adenosine, niacinamide patch
WORLD PACK 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, Adenosine

Water, Butylene Glycol, Glycerin, etc

Skin Protectant - Whitening, Anti-Wrinkle, Pores Tightening & Sebum Formation Control

keep out of reach of the children

1. After washing your face, drain off the water and adjust lotion on your face skin.
2. Open the product and attach a mask sheet on your around eyes and full face.
3. Relax for 15-20 minutes and remove it.
4. After getting rid of it, help the rest of liquid on your face to absorb as patting lightly.

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

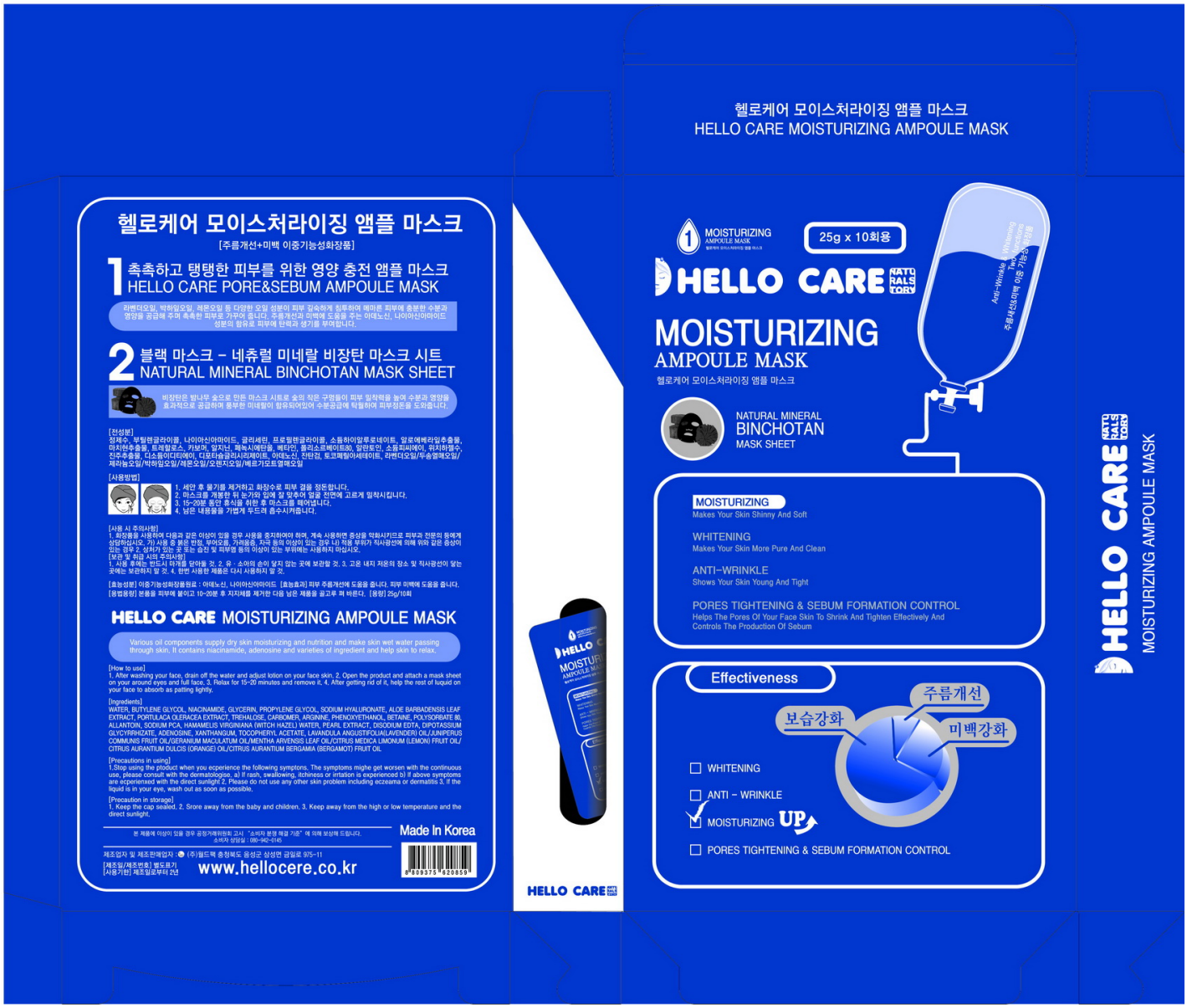
1) Close the lid after use

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



HELLO CARE MOISTURIZING AMPOULE MASK

adenosine, niacinamide patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70959-0001
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 g
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.04 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**

BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70959-0001-1	10 in 1 PACKAGE	08/01/2016	
1		25 g 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		08/01/2016	

Labeler - WORLD PACK CO., LTD. (689054355)

Registrant - WORLD PACK CO., LTD. (689054355)

Establishment

Name	Address	ID/FEI	Business Operations
WORLD PACK CO., LTD.		689054355	manufacture(70959-0001)

Revised: 9/2016 WORLD PACK CO., LTD.