

MOISTIE FILL PLUS- niacinamide, adenosine cream
Purecell Korea 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, Glycerin, Butylene Glycol

Skin Protectant - Whitening & Anti-wrinkle

keep out or reach of the children

After cleansing, apply your skin softner.

Apply an appropriate amount FILL PLUS along your face skin's texture.

After 10 to 15 minutes with the FILL PLUS on your face and Finish with your lotion or cream.

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

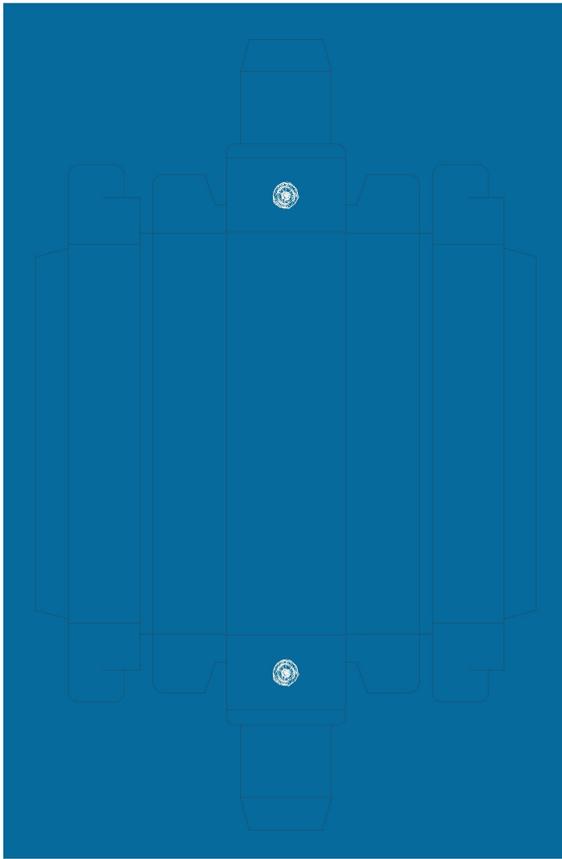
4.Storage and handling precautions

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



■ P 7691C □
 지질 : CCP 350
 무광 코팅

MOISTIE FILL PLUS

niacinamide, adenosine cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71609-0002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X51I8 RD4) (NIACINAMIDE - UNII:25X51I8 RD4)	NIACINAMIDE	2 g in 100 mL
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.04 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71609-0002-1	10 mL in 1 SYRINGE, PLASTIC; Type 0: Not a Combination Product	10/02/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/02/2018	

Labeler - Purecell Korea Co., Ltd. (694667185)**Registrant** - Purecell Korea Co., Ltd. (694667185)**Establishment**

Name	Address	ID/FEI	Business Operations
BIO-FD&C. Co., Ltd.		688203268	manufacture(71609-0002)

Establishment

Name	Address	ID/FEI	Business Operations
Purecell Korea Co., Ltd.		694667185	label(71609-0002)

Revised: 9/2018

Purecell Korea Co., Ltd.