MOISTIE PURE ESSENCE- niacinamide liquid
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

Water, Glycerin, Butylene Glycol

Skin Protectant - Whitening

keep out or reach of the children

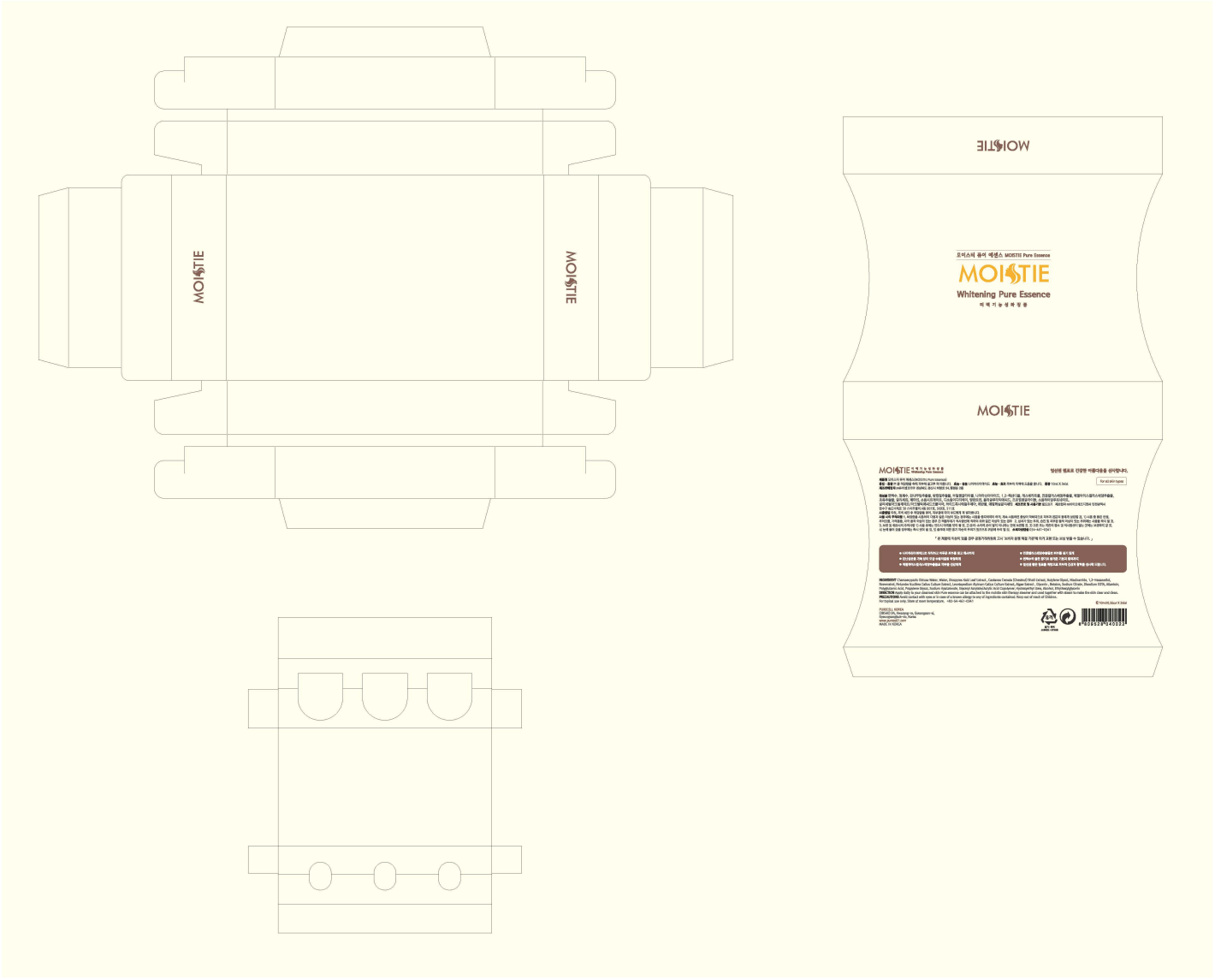
Apply daily to your cleansed skin.

Pure Essence can be attached to the Moistie Skin Therapy Steamer and used together with steam to make the skin clear and clean.

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

pantone 175U　홀로그램 금박 유광　Y:17(연한 아이보리)　무광코팅



MOISTIE PURE ESSENCE

niacinamide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71609-0004
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71609-0004-1	3 in 1 PACKAGE	06/01/2017	
1		10 mL in 1 VIAL; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/01/2017	

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(71609-0004)

Establishment

Name	Address	ID/FEI	Business Operations
Purecell Korea Co., Ltd.		694667185	label(71609-0004)

Revised: 8/2017 Purecell Korea Co., Ltd.