ABALONE CRYSTALDOUBLE EX AMPOULE- arbutin, adenosine liquid
C&BCOSMETIC 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

Arbutin, Adenosine

Water, Glycerin, Butylene Glycol, Etc.

Skin Protectant - Anti-Wrinkle & Whitening

keep out or reach of the children

Take 1EA of ampoule on the palm entirely

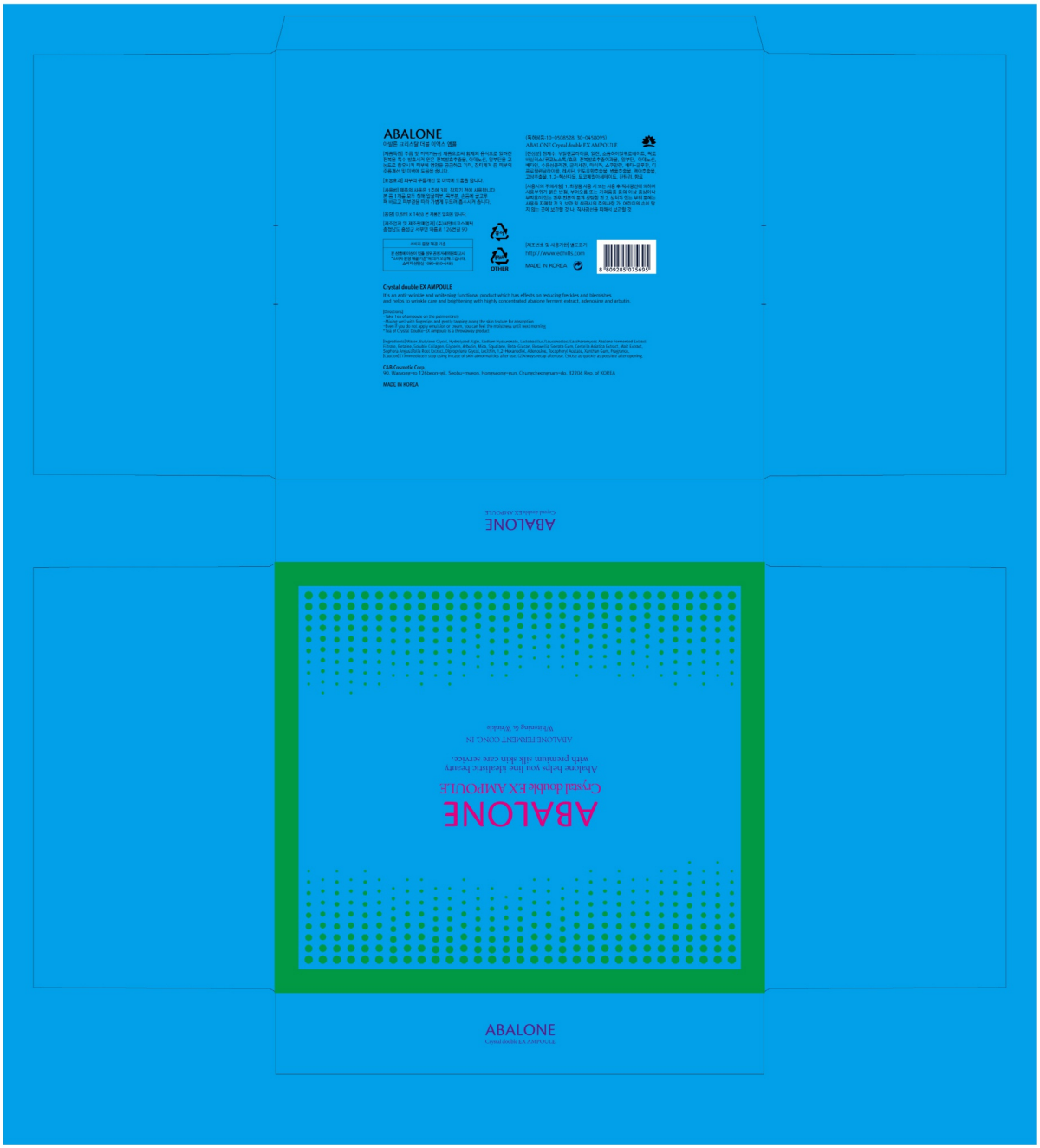
Mixing well with fingertips and gently tapping along the skin texture for absorption

Even if you do not apply emulsion or cream, you can feel the moistness until next morning

*1EA of crystal Double-EX Ampoule is a throwaway 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



ABALONE CRYSTALDOUBLE EX AMPOULE

arbutin, adenosine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60611-0006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ARBUTIN (UNII: C5INA23HXF) (ARBUTIN - UNII:C5INA23HXF)	ARBUTIN	2.2 g in 100 mL
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.044 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60611-0006-1	14 in 1 PACKAGE	03/10/2017	
1		0.8 mL in 1 SYRINGE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/10/2017	

Labeler - C&BCOSMETIC Co.,Ltd. (689909208)

Registrant - C&BCOSMETIC Co.,Ltd. (689909208)

Establishment

Name	Address	ID/FEI	Business Operations
C&BCOSMETIC Co.,Ltd.		689909208	manufacture(60611-0006) , label(60611-0006) , pack(60611-0006)

Revised: 3/2017 C&BCOSMETIC Co.,Ltd.