ABALONE CRYSTALDOUBLE EX- niacinamide, adenosine cream
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

Niacinamide, Adenosine

Water, Lactobacillus/Leuconostoc/Saccharomyces Abalone Fermented Extract, Glycerin, Butylene Glycol, Etc.

Skin Protectant - Moisturizing, Anti-Wrinkle, Whitening

keep out or reach of the children

At the step of skin care, use twice in a day at the morning and evening.

Take an adequate amount of cream amd apply on skin and neck.

Pat lightly for well absorption.

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

아발론 크림 77.5*77.5*53 13/04/08

ABALONE CRYSTALDOUBLE EX

niacinamide, adenosine cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60611-0007
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
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.04 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:60611-0007-1	50 mL in 1 JAR; Type 0: Not a Combination Product	02/20/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/16/2017	

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

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

Establishment

Name	Address	ID/FEI	Business Operations
C&BCOSMETIC Co.,Ltd.		689909208	label(60611-0007) , manufacture(60611-0007) , pack(60611-0007)

Revised: 2/2017 C&BCOSMETIC Co.,Ltd.