

**CAVIALL WRINKLE-FREE POWERTOX- niacinamide 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](#).*

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**Drug Facts**

Niacinamide

Water, Glycerin, Butylene Glycol, Caviar Extract, etc.

Skin Protectant -Anti-Wrinkle

keep out or reach of the children

1. Take 1ea of powertox on the palm entirely.
2. Mixing well with fingertips and gently patting along the skin texture for absorption.
3. Even if you do not apply emulsion of cream, you can feel the moistness until next morning.
- 4.

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

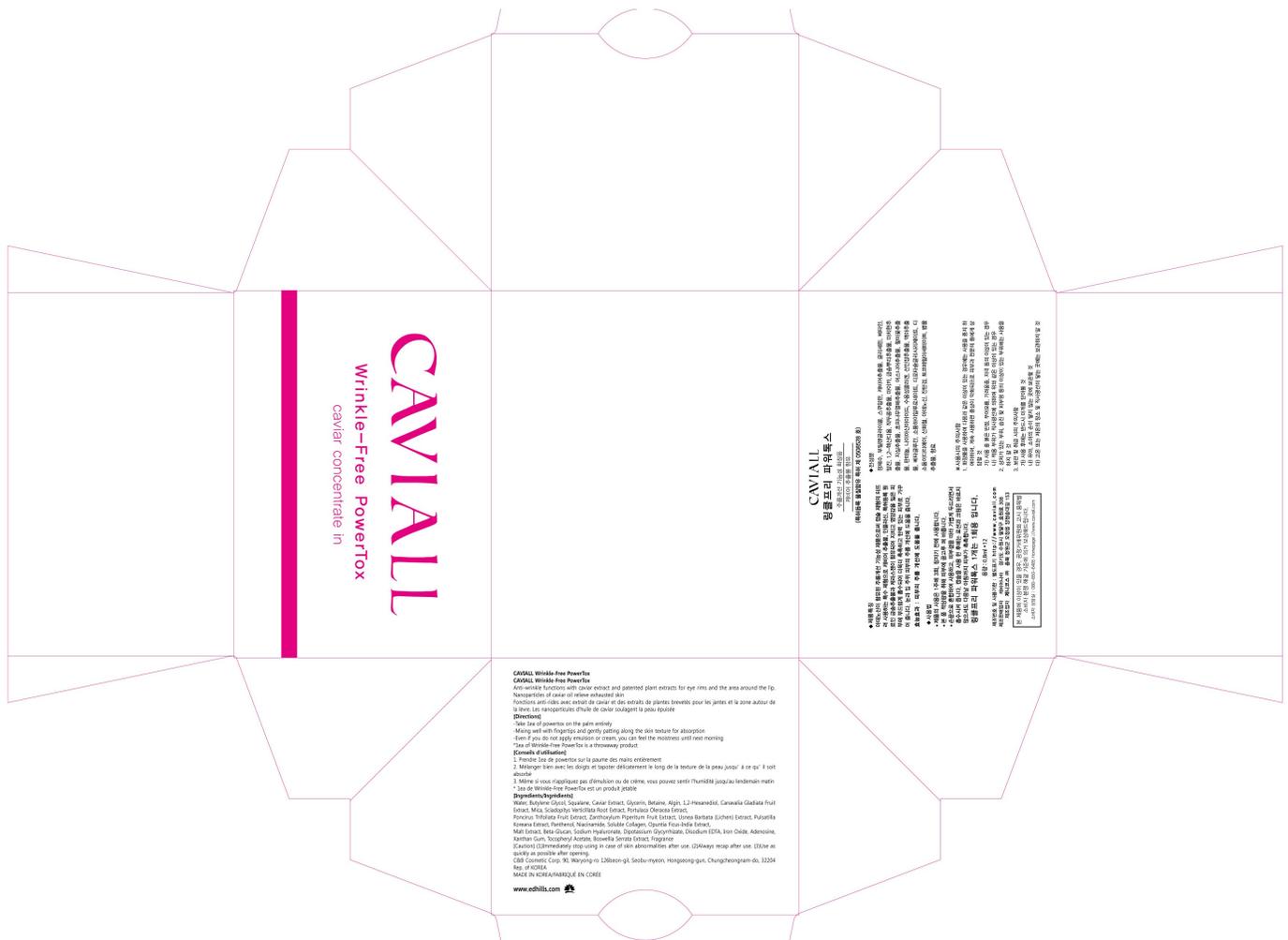
4.Storage and handling precautions

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



# CAVIALL WRINKLE-FREE POWERTOx

niacinamide liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:60611-0003
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X5118 RD4) (NIACINAMIDE - UNII:25X5118 RD4)	NIACINAMIDE	0.3 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CAVIAR, UNSPECIFIED (UNII: 020K6HLU0O)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60611-0003-1	12 in 1 PACKAGE	02/16/2017	
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		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	manufacture(60611-0003) , label(60611-0003) , pack(60611-0003)

Revised: 2/2017

C&amp;BCOSMETIC Co.,Ltd.