

**LEBODY LAB RENEWAL DUAL EFFECT IDEBENONE SERUM- niacinamide,  
adenosine liquid  
GTG Wellness 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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**ACTIVE INGREDIENTS**

Niacinamide 2%

Adenosine 0.04%

**INACTIVE INGREDIENTS**

Water, Butylene Glycol, Glycereth-26, Salicornia Herbacea Extract, Phragmites Communis Extract, Dipropylene Glycol, Ammonium Acryloyldimethyltaurate/VP Copolymer, Hydroxyacetophenone, Glycerin, Cellulose Gum, PEG-40 Hydrogenated Castor Oil, Caprylyl Glycol, Xanthan Gum, 1,2-Hexanediol, Hydrogenated Lecithin, Caprylic/Capric Triglyceride, Disodium EDTA, Panthenol, Hydroxydecyl Ubiquinone, Glycine Soja (Soybean) Sterols, Sodium Hyaluronate, Dipotassium Glycyrrhizate, Honey Extract, Beta-Glucan

**PURPOSE**

Skin Brightening

Anti wrinkle

**Cautions**

1. In case of using cosmetics or after use, please consult a specialist if there is any abnormal symptom or side effect such as red spot, swelling or itching in direct sunlight.
2. Do not use on wounded areas.
3. Precautions for storage and handling
  - a) Keep out of reach of children.
  - b) Avoid direct sunlight.

**KEEP OUT OF REACH OF CHILDREN**

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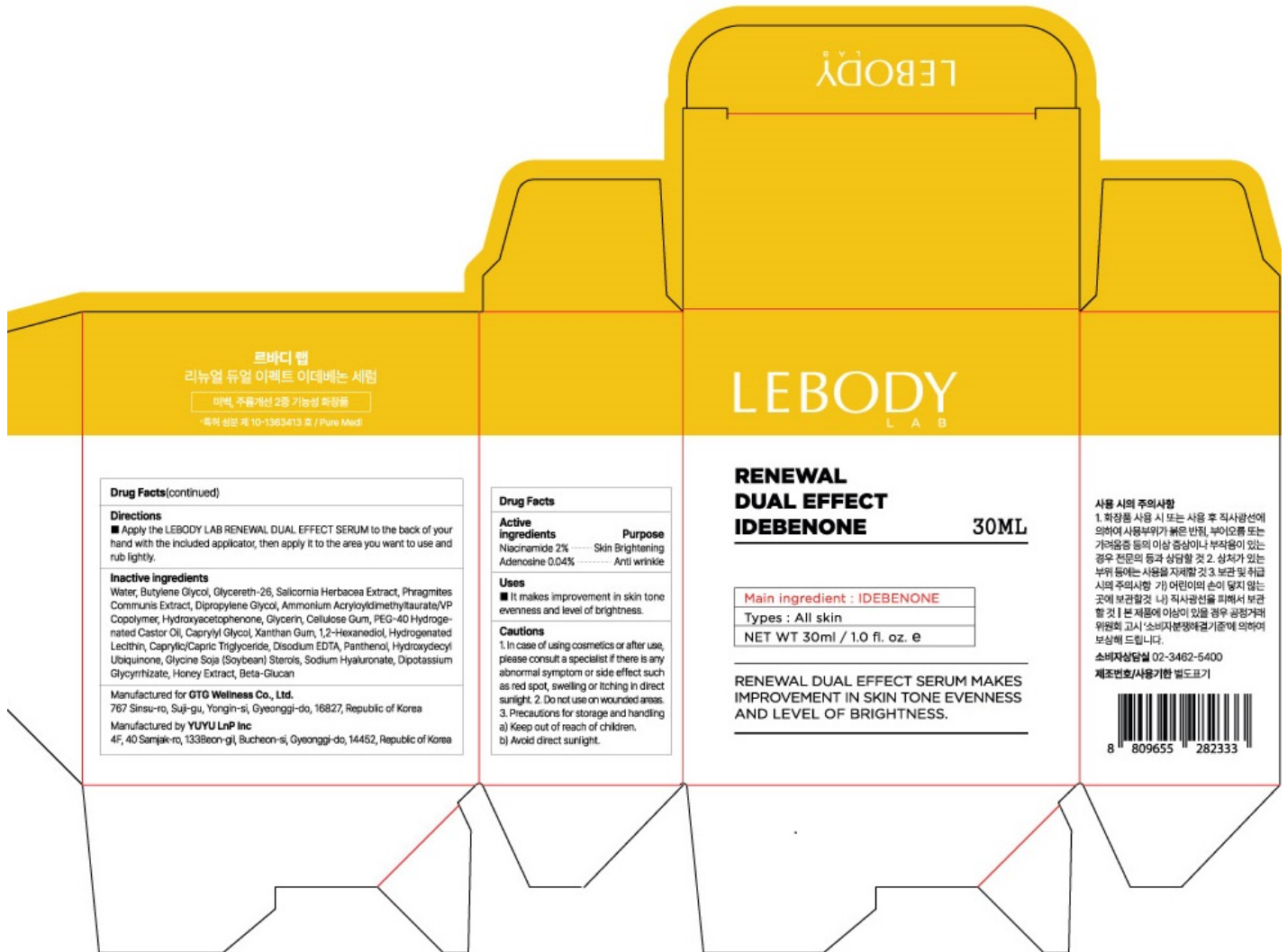
**Uses**

- It makes improvement in skin tone evenness and level of brightness.

## Directions

■ Apply the LEBODY LAB RENEWAL DUAL EFFECT SERUM to the back of your hand with the included applicator, then apply it to the area you want to use and rub lightly.

## PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



LEBODY LAB RENEWAL DUAL EFFECT IDEBENONE SERUM			
niacinamide, adenosine liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71080-0008
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Niacinamide (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	Niacinamide	0.6 g in 30 mL	

## Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Propanediol (UNII: 5965N8W85T)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71080-0008-2	1 in 1 CARTON	03/01/2019	
1	NDC:71080-0008-1	30 mL in 1 CONTAINER; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/01/2019	

**Labeler** - GTG Wellness Co., Ltd. (689458057)

**Registrant** - GTG Wellness Co., Ltd. (689458057)

## Establishment

Name	Address	ID/FEI	Business Operations
YUYU LnP Inc		694752447	manufacture(71080-0008)

Revised: 2/2023

GTG Wellness Co., Ltd.