

DE MEDICOTEM ILLUMINATING WHITE CELL AMPOULE- niacinamide liquid BELLANEZ

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](#).

ACTIVE INGREDIENT

Active ingredients: Niacinamide 2.2%

INACTIVE INGREDIENT

Inactive ingredients:

Water, Dipropylene Glycol, Glycereth-26, Human Stem Cell Conditioned Media, Glycerin, Butylene Glycol, 1,2-HEXANEDIOL, C12-13 Pareth-9, Synthetic Fluorophlogopite, Titanium Dioxide(CI77891), TinOxide, Camellia sinensis Callus Culture extracts, Bambusa Vulgaris Callus Culture Extract, Hydroxyethylacrylate/ Sodium Acryloyldimethyltaurate Copolymer, Butyrospermum Parkii(Shea Butter), Xanthan Gum, Betaine, Ferulic Acid, Pentylene Glycol, Beta-Glucan, Sclerotium gum, Fullerenes, Astaxanthin, Polyglyceryl-10 DIIsostearate, Caprylic/Capric Triglyceride, Sodium Hyaluronate, Hydrogenated Lecithin, Pvp

PURPOSE

Purpose: Skin Brightening

WARNINGS

Warnings:

For external use only

Avoid contact with eyes.

Discontinue use if signs of irritation or rashes appear.

Do not use on the wounded areas.

Keep out of reach of children.

Keep away from direct sunlight

KEEP OUT OF REACH OF CHILDREN

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Uses

Uses:

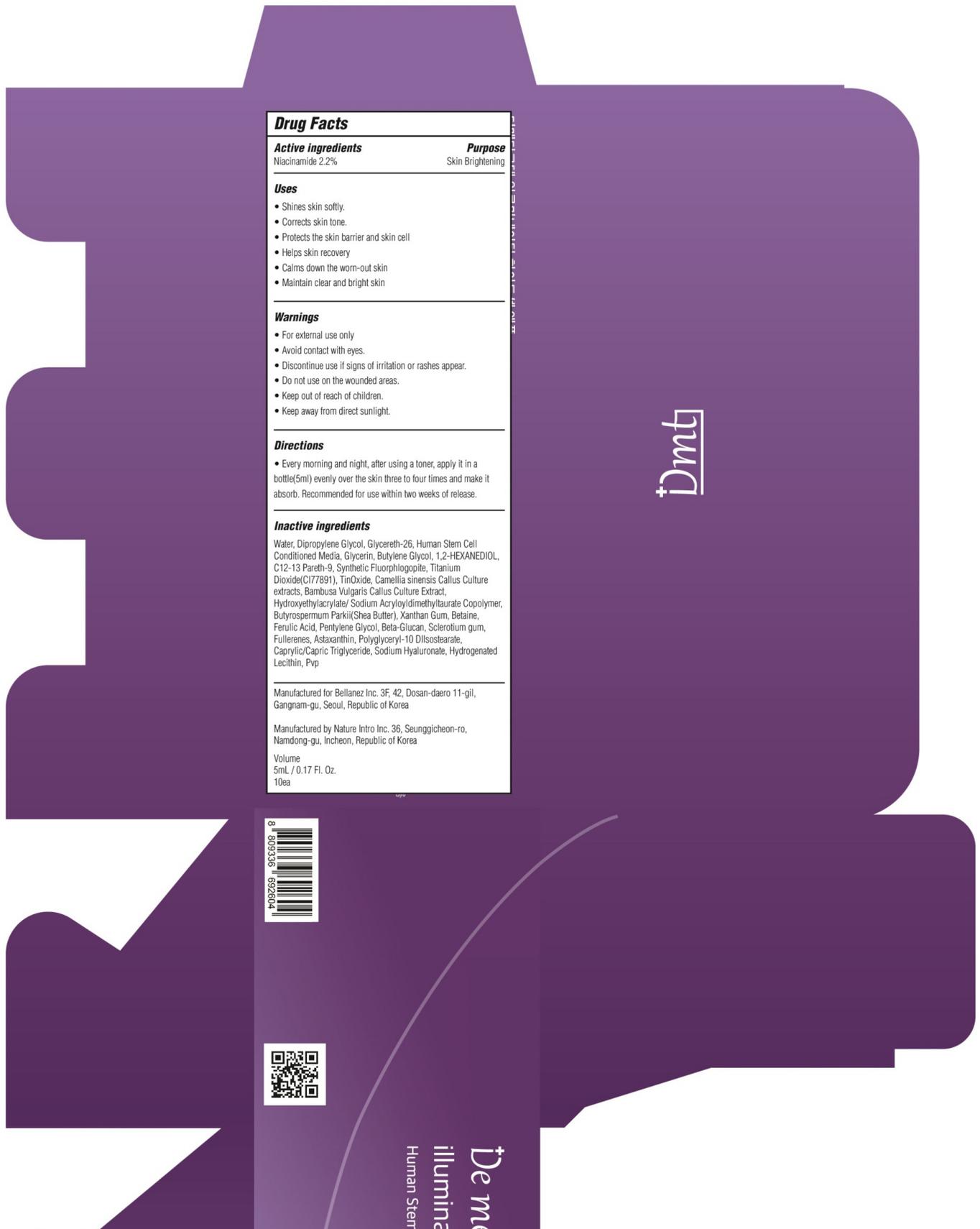
- Shines skin softly.
- Corrects skin tone.
- Protects the skin barrier and skin cell
- Helps skin recovery
- Calms down the worn-out skin
- Maintain clear and bright skin

Directions

Directions:

- Every morning and night, after using a toner, apply it in a bottle(5ml) evenly over the skin three to four times and make it absorb. Recommended for use within two weeks of release.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





DE MEDICOTEM ILLUMINATING WHITE CELL AMPOULE

niacinamide liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72799-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NIACINAMIDE (UNII: 25X5118 RD4) (NIACINAMIDE - UNII:25X5118 RD4)	NIACINAMIDE	0.11 g in 5 mL

Inactive Ingredients

Ingredient Name		Strength		
Water (UNII: 059QF0KO0R)				
Dipropylene Glycol (UNII: E107L85C40)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72799-010-02	10 in 1 CARTON	12/01/2018	
1	NDC:72799-010-01	5 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		12/01/2018		

Labeler - BELLANEZ (694798857)

Registrant - BELLANEZ (694798857)

Establishment

Name	Address	ID/FEI	Business Operations
Nature Intro Co., Ltd.		688282814	manufacture(72799-010)