

BL VITAL ESSENCE - allantoin liquid
RECELL U KOREA

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

allantoin

water, butylene glycol, etc.

skin protectant

keep out of reach of the children

apply proper amount to the skin

1. Do not use in the following cases(Eczema and scalp wounds)

2.Side Effects

1)Due to the use of this drug if rash, irritation, itching and symptoms of hypersensitivity occur discontinue use and consult your pharmacist or doctor

3.General Precautions

1)If in contact with the eyes, wash out thoroughly with water. If the symptoms are severe, seek medical advice immediately

2)This product is for external 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 of reach of children

for external use only

Natural Cosmetic

Deabus

여신의미소 

BL Vital Essence

Revitalizing, moisturizing and
balancing treatment for all skin types
Ultra-moisturizing soothes skin
with its patented Bladderwrack and
Prunellae Spica. White Egg Plant.

BL VITAL ESSENCE

allantoin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70047-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70047-0003-1	50 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	08/31/2015	

Labeler - RECELL U KOREA (689515043)

Registrant - RECELL U KOREA (689515043)

Establishment

Name	Address	ID/FEI	Business Operations
RECELL U KOREA		689515043	manufacture(70047-0003)

Revised: 8/2015

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