BL VITAL - allantoin emulsion
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 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

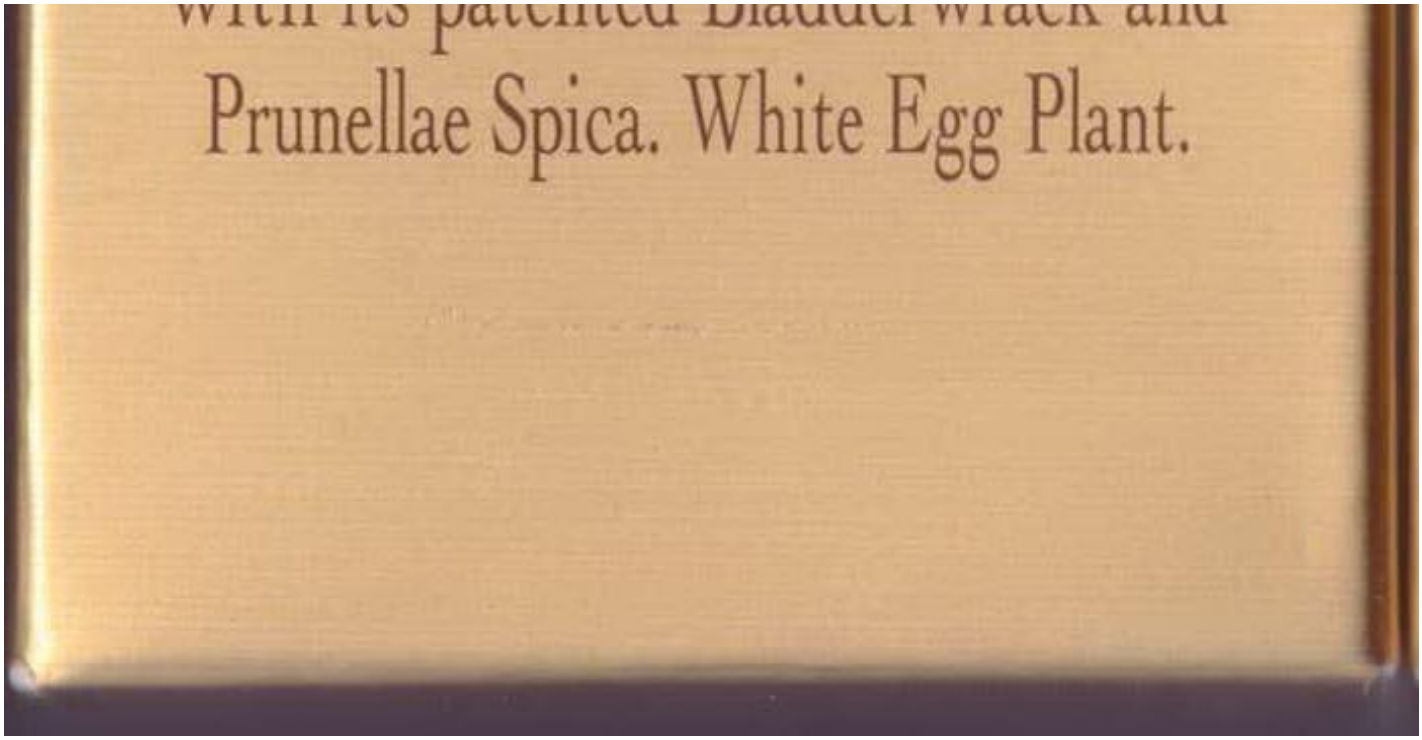
Natural Cosmetic

Deabus

여신의미소

BL Vital Emulsion

Revitalizing, moisturzing and balancing treatment for all skin types
Ultra-moisturizing soothes skin with its patented Bladderwrack and



BL VITAL

allantoin emulsion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70047-0002
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:70047-0002-1	120 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(70047-0002)

Revised: 8/2015 RECELL U KOREA