

**ANY MI NATURAL MAGIC BB NO.7 - allantoin cream**

**K.N.Life Co., Ltd.**

*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.*

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**Drug Facts**

allantoin

water, cyclomethicone, titanium dioxide, zinc oxide, ethylhexyl methoxycinnamate, panax ginseng root ext, peg-7 dimethicone, propylene glycol, arbutin, bentonite, cetyl peg/ppg-10/1 dimethicone, hexyl laurate, betaine, polymethyl methacrylate, sodium chloride, etc

whitening  
anti-wrinkle  
sun block

keep out of reach of the children

after foundation, apply small amounts to whole face by tapping until it is absorbed to skin completely

1. if you have any abnormal symptoms as followings, you should discontinue to use this cream. In case of using continuously, you have to consult to dermatologist because it may make symptoms worse.
  - a. red macule, swelling, urtication, and irritation during use
  - b. symptoms like above by direct ray
2. do not use if you have a wound, eczema, and dermatitis to the area where you apply this cream
3. caution during storing and handling
  - a. close the cap after use
  - b. keep it to the area where babies and infants cannot reach to
  - c. keep it to the area where has not high or low temperature and has no direct ray

for external use only

BB 7호 케이스



# ANY MI NATURAL MAGIC BB NO.7

allantoin cream

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:62695-2001
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.0005 g in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CYCLOMETHICONE (UNII: NMQ347994Z)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ARBUTIN (UNII: C51NA23HXF)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62695-2001-1	40 mL in 1 TUBE		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	03/15/2014	

**Labeler** - K.N.Life Co., Ltd. (688270562)

**Registrant** - K.N.Life Co., Ltd. (688270562)

## Establishment

Name	Address	ID/FEI	Business Operations
K.N.Life Co., Ltd.		688270562	manufacture(62695-2001)