

NAAMAN- allantoin cream
LEADER GREEN 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.

ACTIVE INGREDIENT

Active ingredient: Allantoin 0.5%

INACTIVE INGREDIENT

Inactive ingredients: PONCIRUS TRIFOLIATA FRUIT, RED CLAY POWDER, NARINGINASE, LOBELIA SPICATA LEAF, MINERAL OIL, LIMONENE, (+)-, VITAMINE C, VINCAMINE

PURPOSE

Purpose: Skin Protectant

WARNINGS

Warnings:

1. In case of having following symptoms after using this, you're advised to stop using it immediately. If you keep using it, the symptoms will get worse and need to consult a dermatologist.
 - 1) In case of having problems such as red rash, swollenness, itching, stimulation during usage.
 - 2) In case of having the same symptoms above on the part you put this product on by direct sunlight.
2. You are banned to use it on the part where you have a scar, eczema, or dermatitis.
3. In case of getting it into your eyes, you have to wash it immediately.

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children:

Keep out of reach of babies and children

Indications and usage

Indications and usage:

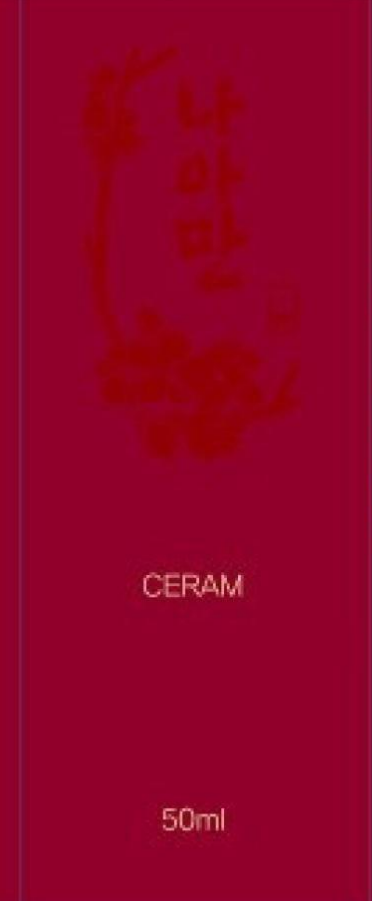
After basic skin care, apply adequate amount following the skin texture and gently tap to enhance absorption.

Dosage and administration

Dosage and administration:

Take an adequate amount of this product.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



CERAM

50ml

NAAMAN CREAM

[Drug Fact]

Active ingredient: Allantoin 0.5%

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Purpose: Skin Protectant

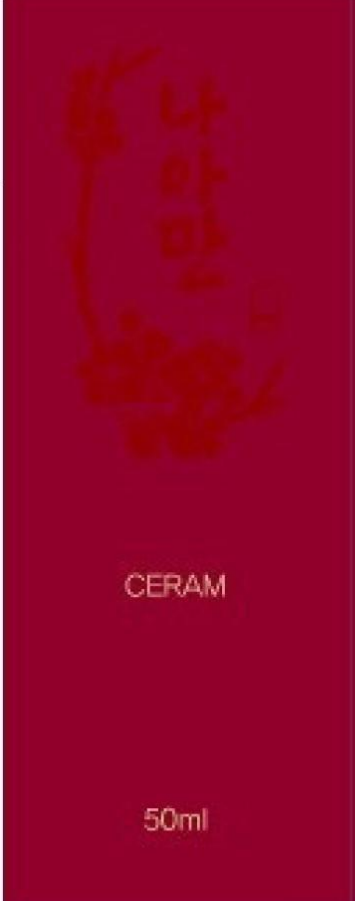
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CERAM


50ml

• **사용법**
비이스다(비이소)를 적당량 피부에 바르고 미지 히트 충분히 문질러서 잘스며들게 하고난 후 나만안(Allaman)을 피부에 골고루 바른 후 10-15분뒤 물로 씻어 냅니다.

• **주의사항**
1) 본 제품을 사용하여 다음과 같이 이상이 있을 경우에는 사용을 중지 하여야 하며 피부과 전문의에게 상담을 받아야 합니다.
가시용증 붉은 반점이 생기거나 부어 오를 경우,
내적증부위기 침사공선에 의하여 의하여 위와 같은 증상이 있을 경우.

• **보관 및 취급상 주의사항**
1) 본 제품은 빛을 차단하여서 사용이 가능하며
2) 고온이나 직사광선에 닿지 않는 곳에 보관하세요.

• **사용기간:제조일로부터 3년**


 용기 : 유리
 포장 : PP

NAAMAN allantoin cream				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:6 1221-0 10	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
Allantoin (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)		Allanto in	0.25 mg in 50 mL	
Inactive Ingredients				
Ingredient Name			Strength	
PONCIRUS TRIFOLIATA FRUIT (UNII: 44445L49J2)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:6 1221-0 10-01	50 mL in 1 CARTON		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	11/01/2013	

Labeler - LEADER GREEN CO., LTD. (688735493)

Registrant - LEADER GREEN CO., LTD. (688735493)

Establishment

Name	Address	ID/FEI	Business Operations
LEADER GREEN CO., LTD.		688735493	manufacture(61221-010)

Revised: 8/2013

LEADER GREEN CO., LTD.