# DERLADIE LABORATOIRE PORE TIGHTENING AMPOULE- niacinamide liquid LAON COMMERCE co ltd

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.

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#### Niacinamide

Water

**Butylene Glycol** 

1,2-Hexanediol

Zinc PCA

Olea Europaea (Olive) Fruit Extract

Centella Asiatica Extract

Portulaca Oleracea Extract

Chamomilla Recutita (Matricaria) Flower Extract

Salvia Officinalis (Sage) Leaf Extract

Rosmarinus Officinalis (Rosemary) Leaf Extract

Salix Alba (Willow) Bark Extract

Melaleuca Alternifolia (Tea Tree) Extract

Tremella Fuciformis (Mushroom) Extract

Nelumbo Nucifera Flower Extract

Hamamelis Virginiana (Witch Hazel) Water

Glycerin

Sodium Hyaluronate

Caprylyl Glycol

Allantoin

Disodium EDTA

Ethylhexylglycerin

skin protect

#### KEEP OUT OF REACH OF THE CHILDREN

After preparing the skin texture with toner, gently spread it over the enlarged pores and let it absorb.

- 1. During using product or after use, if there are any abnormal symptoms or side effects such as red spots, swelling or itching in the area of use due to direct sunlight, consult with a specialist.
- 2. Refrain from using on areas with scars, etc.
- 3. Precautions for storage and handling
- A) Keep out of reach of chil

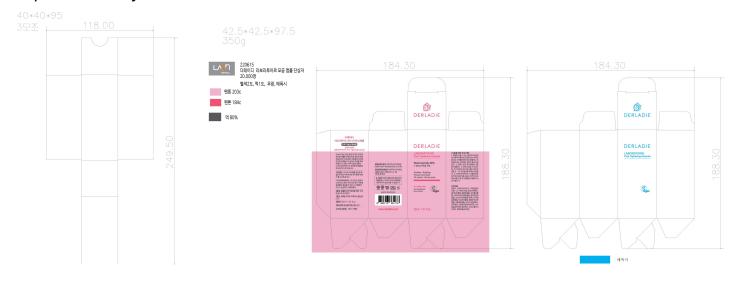
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B) Store away from direct sunlight

4.

Since a high dose of niacinamide is prescribed, please adjust the amount after performing a skin sensitivity test when using for the first time

### topical use only



#### **DERLADIE LABORATOIRE PORE TIGHTENING AMPOULE**

niacinamide liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82083-0028	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
NIACINAMIDE (UNII: 25X5118RD4) (NIACINAMIDE - UNII:25X5118RD4)	NIACINAMIDE	20 g in 100 mL	

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
CENTELLA ASIATICA (UNII: 7M867G6T1U)				

Packaging					
	#	Item Code Package Description		Marketing Start Date	Marketing End Date
	1	NDC:82083- 0028-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	12/01/2023	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		12/01/2023		

### Labeler - LAON COMMERCE co ltd (557839830)

## Registrant - LAON COMMERCE co ltd (557839830)

Establishment				
Name	Address	ID/FEI	Business Operations	
LAON COMMERCE CO Ltd		557839830	label(82083-0028) , manufacture(82083-0028)	

Revised: 12/2023 LAON COMMERCE co ltd