DERMAHAN UV SUN- adenosine, niacinamide cream
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.

Niacinamide, Adenosine

Water
Zinc Oxide
Cyclopentasiloxane
Titanium Dioxide
Butyloctyl Salicylate
Glycerin
Cetyl PEG/PPG-10/1 Dimethicone
Niacinamide
Dicaprylyl Carbonate
Sodium Chloride
Triethoxycaprylylsilane
Aluminum Hydroxide Oxide
PEG-10 Dimethicone
Panthenol
Stearic Acid
Disteardimonium Hectorite
Benzyl Glycol
Water
Styrene/Acrylates Copolymer
Butylene Glycol
Allantoin
Propylene Carbonate
Ethylhexylglycerin
Adenosine
Disodium EDTA
1,2-Hexanediol
Borago Officinalis Extract
Corchorus Olitorius Leaf Extract
Ilex Paraguariensis Leaf Extract
Daucus Carota Sativa (Carrot) Root Extract
Sodium Hyaluronate
Fragrance

skin protect

KEEP OUT OF REACH OF THE CHILDREN

In the last step of basic care, apply an appropriate amount to the face, neck, arms, legs, etc. that are prone to UV exposure.

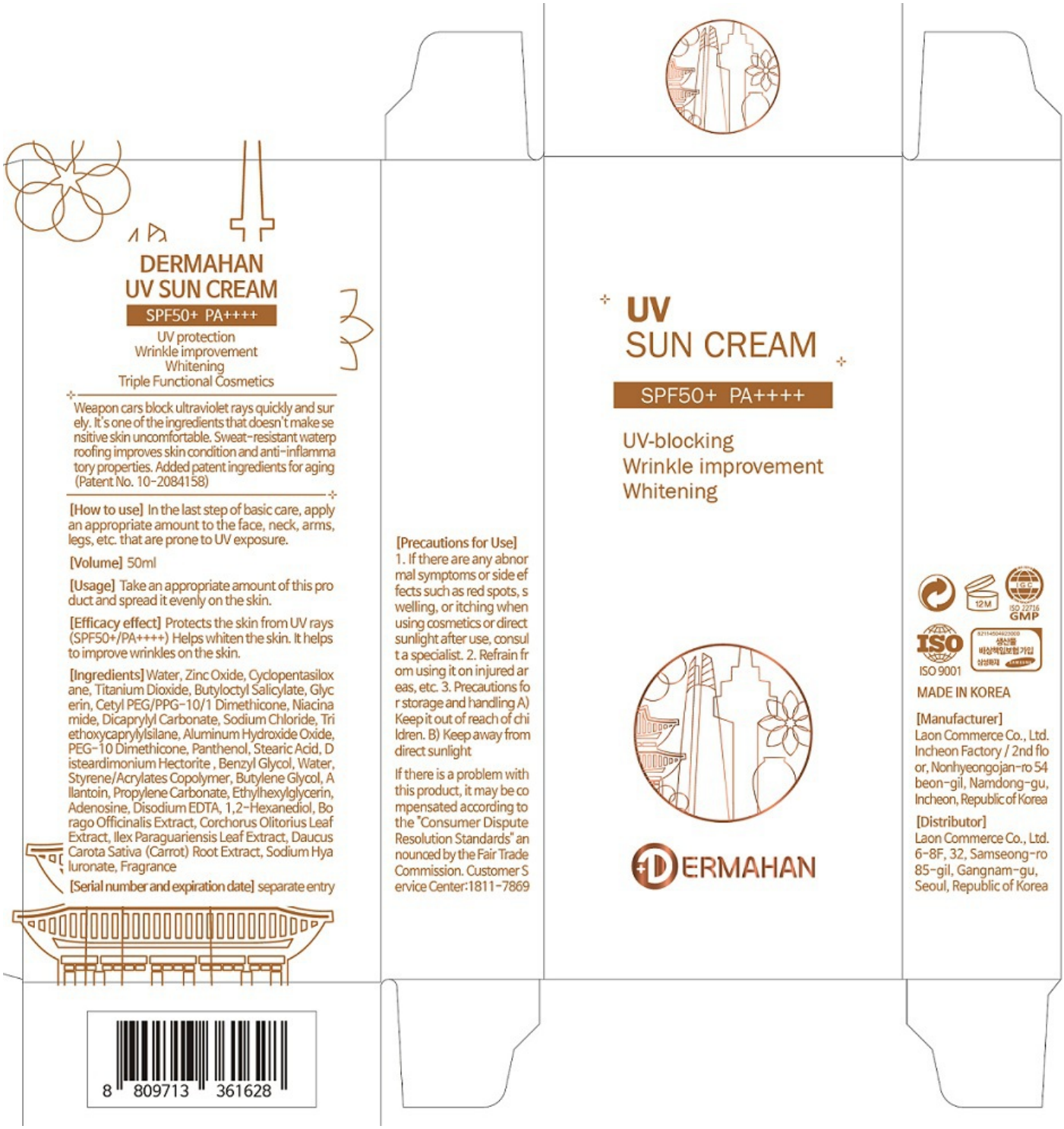
1. If there are any abnormal symptoms or side effects such as red spots, swelling, or itching when using cosmetics or direct sunlight after use, consult a specialist.

2. Refrain from using it on injured areas, etc.

3. Precautions for storage and handling A) Keep it out of reach of children. B) Keep away from direct sunlight

3. Protects the skin from UV rays(SPF50+/PA++++) Helps whiten the skin. It helps to improve wrinkles on the skin.

topical use only



DERMAHAN UV SUN

adenosine, niacinamide cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82083-0016
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ADENOSINE (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	ADENOSINE	0.04 g in 100 mL
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82083-0016-1	50 mL in 1 TUBE; Type 0: Not a Combination Product	04/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/01/2023	

Labeler - LAON COMMERCE co ltd (557839830)

Registrant - LAON COMMERCE co ltd (557839830)

Establishment

Name	Address	ID/FEI	Business Operations
LAON COMMERCE CO Ltd		557839830	manufacture(82083-0016) , label(82083-0016)

Revised: 4/2023 LAON COMMERCE co ltd