REORGANIC WOMANS INNER CARE ESSENCE GEL- adenosine gel
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.

Adenosine

Water
Butylene Glycol
Glycereth-26
Hydroxyethylcellulose
Panthenol
Lactobacillus Ferment
Glyceryl Caprylate
Carbomer
Allantoin
Decylene Glycol
Hydroxyacetophenone
Caprylyl Glycol
Dipropylene Glycol
Dipotassium Glycyrrhizate
Xanthan Gum
Arginine
Salix Alba (Willow) Bark Extract
Tremella Fuciformis (Mushroom) Extract
Propanediol
Polyglyceryl-4 Oleate
Melaleuca Alternifolia (Tea Tree) Extract
Collagen Extract
Olea Europaea (Olive) Fruit Extract
Nelumbo Nucifera Flower Extract
Isopentyldiol
Asiatic Acid
Asiaticoside
Portulaca Oleracea Extract
Sodium Stearoyl Glutamate
Sodium Surfactin
Salvia Officinalis (Sage) Leaf Extract
Centella Asiatica Extract
Chamomilla Recutita (Matricaria) Flower Extract
Madecassic Acid
Madecassoside
Rosmarinus Officinalis (Rosemary) Leaf Extract
Glycerin
1,2-Hexanediol

skin protect

KEEP OUT OF REACH OF THE CHILDREN

Take an appropriate amount of this product and spread it evenly on the skin

1. When using cosmetics or after use, consult with a sp ecialist if there are any abnormal symptoms or side effects such as red spots, swelli ng, or itching in the area of use due to direct sunlight. 3. Precautions for storage and handling A) Keep out of reach of children B) Keep away from direct sunlight

topical use only

REORGANIC WOMANS INNER CARE ESSENCE GEL

adenosine gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82083-0018
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

Inactive Ingredients

Ingredient Name	Strength
PANTHENOL (UNII: WV9CM0O67Z)	
ALLANTOIN (UNII: 344S277G0Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82083-0018-1	10 in 1 BOX	05/01/2023	
1		1.7 mL in 1 SYRINGE, PLASTIC; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2023	

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(82083-0018) , label(82083-0018)

Revised: 5/2023 LAON COMMERCE co ltd