ALL NATIONS A D C SPECIAL PROGRAM- adenosine solution AN 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.

ACTIVE INGREDIENT

Active ingredients: Adenosine 0.04%

INACTIVE INGREDIENT

Inactive ingredients:

SP F I) Water, illite Extract, Carbomer, Butylene Glycol, Caffeine, Phenoxyethanol, Carprylyl Glycol, Glycerin, 1,2-Hexanediol, Avena Sativa (Oat) Kernel Extract, Beta-Glucan, Acetyl Hexapeptide-8, Copper Tripeptide-1, Diospyros Kaki Leaf Extract, Centella Asiatica Extract, Theobroma Cacao Extract, Chamomilla Recutita (Matricaria) Extract, Wine Extract, Ethylhexylglycerin, Hovenia Dulcis Fruit Extract, Camellia Sinensis Leaf Extract, Glycine Soja (Soybean) Seed Extract, Origanum Vulgare Flower Extract, Scutellaria Baicalensis Root Extract, Sophora Flavescens Root Extract

SP F II) Water, illite Extract, Sodium Silicate, Butylene Glycol, Hydroxypropyl Chitosan, Glycerin, 1,2-Hexanediol, Phenoxyethanol, Citric acid, Polyepsilon-Lysine, Alcohol, Paeonia Albiflora Flower Extract, Ipomoea Purpurea Extract, Magnolia Liliflora Flower Extract, Lilium Candidum Flower Extract, Portulaca Oleracea Extract, Camellia Sinensis Leaf Extract

Volume) SP F I: 6mL, SP F II: 1mL

PURPOSE

Purpose: SKIN PROTECTANT

Precautions on use

Precautions on use: 1. Stop using the product if there are any of the following abnormal symptoms appearing after use and consult with your dermatologist as continued use can worsen the symptoms. A) If there are red spots, swelling, itchiness, or irritation B) If the above symptoms appear around the skin to which the product has been applied after being exposed to direct sunlight. 2. Precautions on storage and handling. A) Make sure to close the cap after use. B) Keep out of the reach of children. C) Do not keep the product in a hot or cold place or a place getting direct sunlight. 3. Do not apply the product to any parts of the skin with wound, eczema, or dermatitis.

KEEP OUT OF REACH OF CHILDREN

KEEP OUT OF REACH OF CHILDREN

INDICATIONS & USAGE

Suggested use and application order:

1. Open SP F I until the dotted line. (try not to spill the liquid content inside) 2. Open SP F I and mix with SP F I. (try not to spill the liquid content inside) 3. Seal the package of SP F I and strongly shake the package more than 10 times to mix two agents well. (turning into gel when mixed enough) 4. Apply the content over your face enough with a brush contained inside the package. 5. Regardless of individual

difference, wait for 15 ~ 20 minutes. (gently fan yourself while waiting) Try not to move or speak. 6. For the first few times, your skin may feel strongly stretched and there is a difference in color change depending on area. (discharge of toxin out of the skin) Feeling of stretching may differ by individual skin condition. 7. After using wash your face with warm water and then use soap to cleanse your face once again. 8. Dry your face with a dry towel.

*Use up a single package at a time.

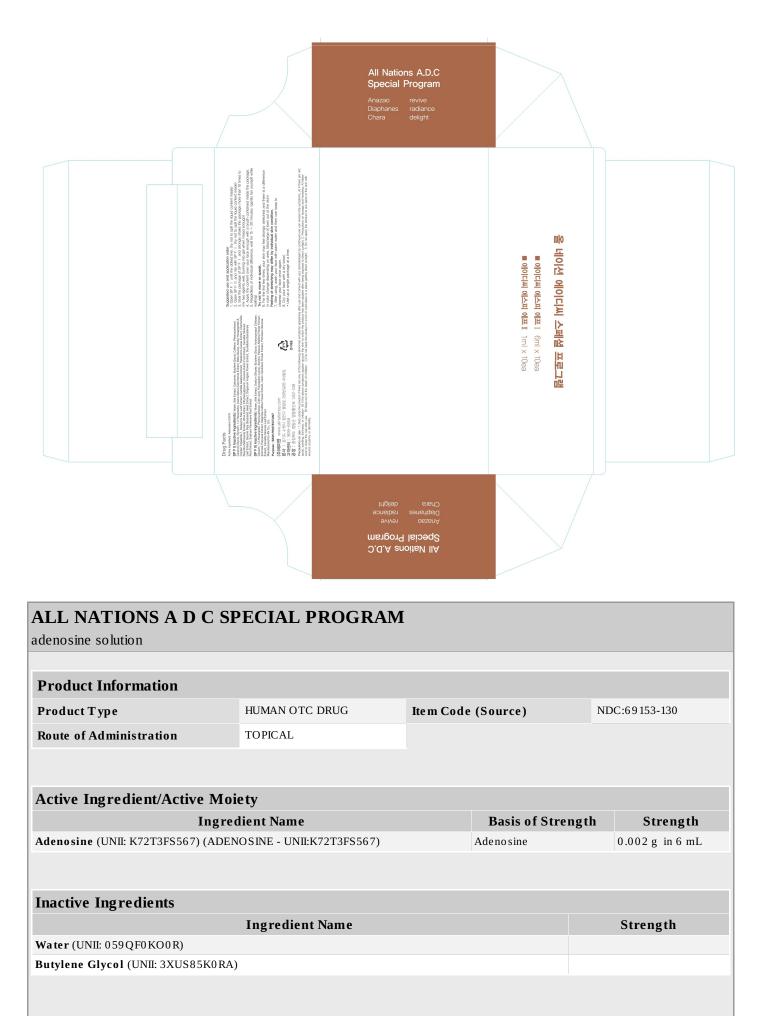
DOSAGE & ADMINISTRATION

Suggested use and application order:

1. Open SP F I until the dotted line. (try not to spill the liquid content inside) 2. Open SP F I and mix with SP F I. (try not to spill the liquid content inside) 3. Seal the package of SP F I and strongly shake the package more than 10 times to mix two agents well. (turning into gel when mixed enough) 4. Apply the content over your face enough with a brush contained inside the package. 5. Regardless of individual difference, wait for 15 ~ 20 minutes. (gently fan yourself while waiting) Try not to move or speak. 6. For the first few times, your skin may feel strongly stretched and there is a difference in color change depending on area. (discharge of toxin out of the skin) Feeling of stretching may differ by individual skin condition. 7. After using wash your face with warm water and then use soap to cleanse your face once again. 8. Dry your face with a dry towel.

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PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69153-130-02	10 in 1 CARTON	06/01/2017			
1	NDC:69153-130-01	6 mL in 1 CONTAINER; Type 0: Not a Combination Product				
Marketing Information						
N	Aarketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
ur	approved drug other		06/01/2017			

Labeler - AN Co Ltd. (688448454)

Registrant - AN Co Ltd. (688448454)

Establishment						
Name	Address	ID/FEI	Business Operations			
AN Co Ltd.		688448454	manufacture(69153-130)			

Revised: 6/2017

AN Co Ltd.