BIOTOC REGEN PEEL- adenosine gel
Dermafirm INC.

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.

Drug Facts

Adenosine

Water, Glycerin

Anti-Wrinkle

keep out of reach of the children

Apply an appropriate amount of BIOTOC Regen Peel to pores and wrinkles especially around the lips and under the eyes.

Then massage roundly with your fingertips pressure for 2 or 3 minutes, and press firmly with your fingers.

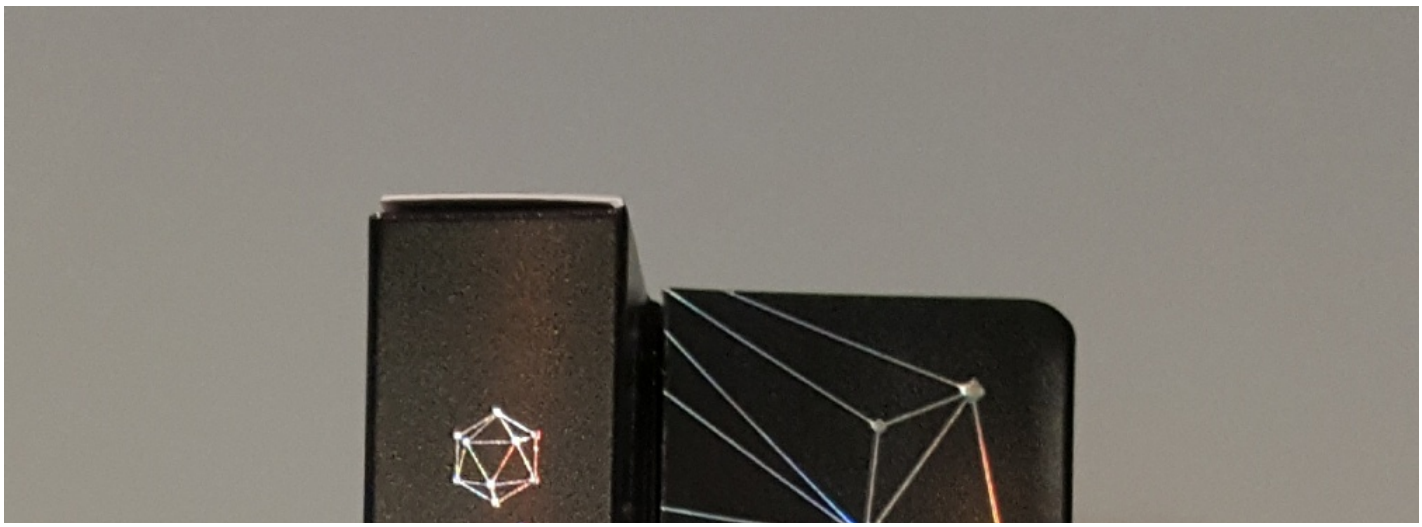
Apply BIOTOC Serum and Cream thickly after using BIOTOC Regen Peel.

It is effective to use mask pack together.

For the best results, apply BIOTOC Regen Peel twice a week and BIOTOC Regen Ampoule, Serum, and Cream every morning and evening.

1. Do not use in the following cases(Eczema and scalp wounds)
2.Side Effects
1)Due to the use of this druf if rash, irritation, itching and symptopms of hypersnesitivity occur dicontinue use and consult your phamacisr or doctor
3.General Precautions
1)If in contact with the eyes, wash out thoroughty with water If the symptoms are servere, seek medical advice immediately
2)This product is for exeternal use only. Do not use for internal use
4.Storage and handling precautions
1)If possible, avoid direct sunlight and store in cool and area of low humidity
2)In order to maintain the quality of the product and avoid misuse
3)Avoid placing the product near fire and store out in reach of children

for external use only



BIOTOC REGEN PEEL

adenosine gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71638-0004
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 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71638-0004-1	13.5 g in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	09/04/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/04/2017	

Labeler - Dermafirm INC. (690171603)

Registrant - Dermafirm INC. (690171603)

Establishment

Name	Address	ID/FEI	Business Operations
Dermafirm INC.		690171603	label(71638-0004) , pack(71638-0004) , manufacture(71638-0004)

Revised: 9/2017 Dermafirm INC.