MEDI HYDRO DP B TOX AMPOULE- adenosine liquid
Inc MBG

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.

Open the lid, press the lid to fill the dropper and apply evenly 3-4 drops to the desired parts, gently tap with the fingers to allow it to absorb

Keep out of reach of children.

for external use only

Discontinue use of the product if any of the following symptoms occurred. Consult a dermatologist if the symptoms got worse.
a) If redness, Swelling, Itchiness, Irritation occurred during using.
b) If the above symptoms started to occur when exposed to direct sunlight.
Do not use on skins with wounds, eczema or dermatitis.

Storage instructions
a) Always close the lid after finish using.
b) Store in a safe place away and out of reach of children.
c) Store in a place with not too high or low temperature, keep awayfrom direct sunlight exposure.

Ampoule that provides skin elasticity with active ingredients such as atelocollagen and 3D hyaluronic acid.

water, butylene glycol, hyaluronic acid, etc.

ADENOSINE

For All Skin Types

Main Ingredients

Common ingredients – Hydrogen water, Dendropanax Morbifera, Sparassis Crispa Fermented Extract (Patent Number 10-1275971).
Other ingredients – Marked separately on products label.

Caution

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   b) If the above symptoms started to occur when exposed to direct sunlight.
2. Do not use on skins with wounds, eczema or dermatitis.
3. Storage instructions
   a) Always close the lid after finish using.
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MEDI HYDRO DP B TOX AMPOULE

adenosine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70694-0010
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**
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70694-0010-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/24/2019	

Labeler - Inc MBG (688436167)

Establishment

Name	Address	ID/FEI	Business Operations
Inc MBG		688436167	manufacture(70694-0010)

Revised: 1/2019 Inc MBG