

APPLICELL PERFECT EYE- adenosine cream
Anc HnB 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](#).

ACTIVE INGREDIENT

Active Ingredient: Adenosine 0.04%

INACTIVE INGREDIENT

Inactive Ingredients: Water, Glycerin, Dipropylene Glycol, Hydrogenated Polydecene, Cyclopentasiloxane, Cyclohexasiloxane, Hydrolyzed Keratin, Hydrolyzed Collagen, Hydrolyzed Elastin, Cetearyl Alcohol, Polyglyceryl-3 Methylglucose Distearate, Betaine, Butyrospermum Parkii(Shea Butter), Neopentyl Glycol Dicaprate, Dimethicone, Triethylhexanoin, Glyceryl Stearate SE, Sodium Hyaluronate, Human fibroblast Conditioned Media, Stearic Acid, Beeswax, PEG-100 Stearate, Citrus Paradisi (Grapefruit) Fruit Extract, Glyceryl Stearate, Tocopheryl Acetate, Xanthan Gum, Ceramide 3, Hydrogenated Lecithin, Neofinetia Falcata Callus Culture Extract, Caprylhydroxamic Acid, Caprylyl Glycol, Phenoxyethanol, Carbomer, Potassium Hydroxide, Trisodium EDTA, Fragrance

PURPOSE

Purpose: Skin Protectant

WARNINGS

Warnings: For external use only When using this product do not get into eyes, Stop use and ask a doctor if rash occurs Do not use on - deep puncture wounds - animal bites - serious burns Keep out of reach of children

KEEP OUT OF REACH OF CHILDREN

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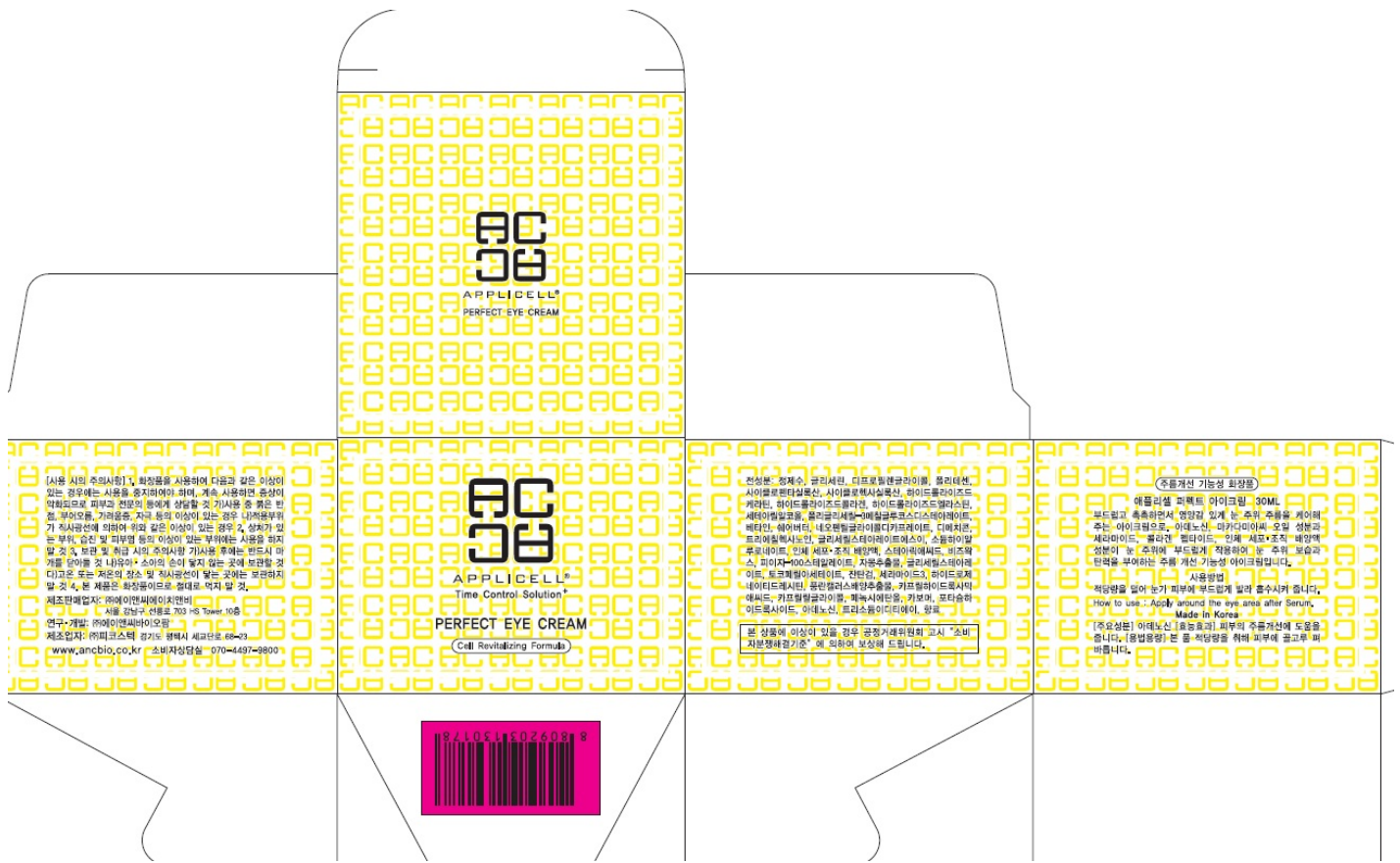
INDICATIONS & USAGE

Indications & usage: APPLICELL Perfect Eye Cream will moist and elastically care for wrinkle around the eyes.

DOSAGE & ADMINISTRATION

How to use: Apply around the eye area after Serum

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



APPLICELL PERFECT EYE

adenosine cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71098-050
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Adenosine (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	Adenosine	0.01 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71098-050-02	1 in 1 CARTON	10/10/2016	
	NDC:71098-050	30 mL in 1 BOTTLE, PLASTIC, Type 0; Net Combination		

1	NDC: 71098-050-01	30 mL in 1 BOTTLE, PLASTIC; Type of: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		10/10/2016		

Labeler - Anc HnB Inc. (689031836)

Registrant - Anc HnB Inc. (689031836)

Establishment			
Name	Address	ID/FEI	Business Operations
Anc HnB Inc.		689031836	manufacture(71098-050)

Revised: 11/2016

Anc HnB Inc.