APPLICELL ADVANCED- dimethicone lotion Anc HnB Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ACTIVE INGREDIENT

Active Ingredient: Dimethicone 3.5%

INACTIVE INGREDIENT

Inactive Ingredients: Water, Hydrogenated Polydecene, Triethylhexanoin, Butylene Glycol, Glycerin, Citrus Paradisi (Grapefruit) Fruit Extract, PEG-5 Glyceryl Stearate, PEG-100 Stearate, Glyceryl Stearate, Isostearic Acid, Stearic Acid, Macadamia Ternifolia Seed Oil, Betaine, Trehalose, Glyceryl Stearate SE, Behenic Acid, Behenyl Alcohol, Human fibroblast Conditioned Media, Neofinetia Falcata Callus Culture Extract, Phenoxyethanol, Fragrance, Carbomer, Portulaca Oleracea Extract, Hydrolyzed Keratin, Hydrolyzed Collagen, Hydrolyzed Elastin, Coix Lacryma-Jobi Ma-yuen Seed Extract, Potassium Hydroxide, Trisodium EDTA, Caprylhydroxamic Acid, Caprylyl Glycol

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 Advanced Lotion is available to nourish to the skin and brings a soft, moist feel.

DOSAGE & ADMINISTRATION

How to use: After serum, take proper amount and apply over the face along skin texture.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



APPLICELL ADVANCED

dimethicone lotion

IAN OTC DRUG	Item Code (Source)	NDC:71098-010
PICAL		
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Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	

Dimethicone (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	Dimethicone	4.55 g in 130 mL
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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:71098-010- 02	1 in 1 CARTON	10/10/2016		
1	NDC:71098-010- 01	130 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	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(71098-010)		

Revised: 11/2016 Anc HnB Inc.