# PLAGENTRA BABY SUNCITY- octinoxate, octisalate, titanium dioxide cream C.A Pharm Co., Ltd.

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 Ingredients: ETHYLHEXYL METHOXYCINNAMATE 7.5%, ETHYLHEXYL SALICYLATE 5.0%, TITANIUM DIOXIDE 3.0%

### **INACTIVE INGREDIENT**

**Inactive Ingredients:** 

Water, Cyclopentasiloxane, Isohexadecane, Butylene Glycol, Isoamyl p-Methoxycinnamate, Glycerin, Polysorbate 60, C12-15 Alkyl Benzoate, Dimethicone, Glyceryl Stearate, Dimethicone Crosspolymer, Sorbitan Stearate, Stearic Acid, VP/Eicosene Copolymer, PEG-100 Stearate, Cetearyl Alcohol, Sodium Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Polysorbate 80, PEG-10 Dimethicone, Hexyl Laurate, Panthenol, Caprylyl glycol, Fragrance, Ethylhexylglycerin, Tropolone, Tocopheryl Acetate, Alumina, Magnesium Aluminum Silicate, Polyglutamic Acid, Leuconostoc/Radish Root Ferment Filtrate, Arnica Montana Flower Extract, Achillea Millefolium Extract, Gentiana Lutea Root Extract, Artemisia Absinthium Extract, Xanthan Gum, Disodium EDTA, Sodium Hyaluronate, Ethyl Ascorbyl Ether, Human Oligopeptide-1, Oligopeptide-3

#### **PURPOSE**

Purpose: UVA/UVB Protection

#### WARNINGS

Warnings:

- 1. Stop using the product and go to a doctor immediately if one of the following symptoms occurs. If immediate care is not sought, the symptoms may worsen:
- 1) Itching, redness, swelling, rash, etc. 2) If one of the symptoms above occurs due to direct sunlight.
- 2. Do not apply the product to wounds or skin with dermatitis such as eczema.
- 3. Storage and Handling, 1) Keep the lid closed after use. 2) Keep the product out of children's reach. 3) Keep away from direct sunlight, do not store at high or low temperature.
- 4. It contains AHA. If it is your first time to use this product, take a small amount and gently apply to the skin to check if it cause andy trouble to your skin.

### KEEP OUT OF REACH OF CHILDREN

Keep out of reach of babies and children.

### **INDICATIONS & USAGE**

Indication and usage:

- 1) Tighten the lid after using it.
- 2) Don't keep it in the place where the temperature is extremely hot or low and exposed to the direct sunlight.
- 3) Use it on the day you will be in the sun, 30 minutes before you go outside, apply an ounce of

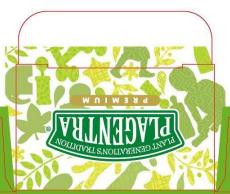
sunscreen.

### **DOSAGE & ADMINISTRATION**

Dosage and administration:

To apply sunscreen, squeeze a dollop of sun cream onto your hand and rub it all the skin that will be exposed to UV.

### PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





Active Ingredients: %(W/W)
Ethylhexyl Methoxycinnamate 7.500
Ethylhexyl Salicylate 5.000
Titanium Dioxide 3.000

Inactive Ingredients: 
Water, Oxfopertasilozone, Isohexadecane 
Burlene Glycal, Isoamilp Methoxychnanoste. 
Glycerin, Polysorbate 60, C12-15 Alkyl 
Benzoate, Dimethicone, Glyceryl Steanate 
Dimethicone Crosspolymer, Sorbitar 
Stearite, Stearit Add, VP/Ecosene Copolyme 
FDG-100 Stearate, Cetearyl Alcohol, Sodium 
Acrylate/Sodium Acryloydilmethyl Taurate 
Acrylate/Sodium Acryloydilmethyl Taurate 
Copolymer, Polysorbate 80, PEG-10 
Dimethicone, Heavyl Laurate, Parthenol 
Capnyl gylox, Fingarune, Ethyhdwyllycerin 
Tiropolone, Tocopheryl Acetate, Allumina 
Magnesium Aluminum Silicate, Dydygutamik 
Add, Leuconostov, Taddish Root Ferment 
Filfrate, Amria Montana Flower Extract 
Adrillea Millefolium Ethract, Gentiana Lutes 
Kanthan Gum, Disodium EDTA, Sodium 
EValukonostov, Ethyl Accorbel Ether, Human 

Kanthan Gum, Disodium EDTA, Sodium 

Ethylaurante, Ethyl Accorbel Ether, Human 

Hylaurante, Ethyl Accorbel Ethyle, Human 

Hylaurante, Ethyl Accorbel Ethyl 

Hylaurante, Ethyl Accorbel Ethyle, Human 

Hylaurante, Ethylaurante, Ethyle, Human 

Hylaurante, Ethylaura

#### Purpose: UVA/UVB Protection

#### Warnings

L. Stop using the product and go to doctor immediately if one of the following improms occurs. If immediate care is not lought, the symptoms may worsen: I) Itching, redness, swelling, rash, etc. I) If one of the symptoms above occurs due to direct sunlight.

 Do not apply the product to wounds or skin with dermatitis such as eczema.
 Storage and Handling, 1) Keep the lid closed after use.
 Keep the product out of children's reach

3) Keep away from direct sunlight, do not store at high or low temperature.

4. This product contains AHA. If it is your first time to use this product, take a small amount and gently apply to a small patch of skin to check for any irritation.

Keep out of reach of babies and children.



Sun&Air pollution block



**Baby Suncity Cream** 



"Everyone's goose is a gander We make products with

### Baby Suncity Cream SPF36/PA++

Baby Suncity Cream which maximizes SPF efficiency and at the same time, minimizes skin irritation by distributing powder particles evenly based on the advanced UV dispersing technology "Powder in Gel System".

Efficacy: Protects the skin from UV rays

#### ndication and usage :

Tighten the lid after using it. 2) Don't keep it in the place where the temperature is extremely hot or low and exposed to the direct sunlight. 3) Use it on the day you will be in the sun, 30 minutes before you go outside, apply an ounce of sunscreer Dosage and administration:

To apply sunscreen, squeeze a dollop of sun cream onto your hand and rubing all the skin that will be exposed to UV. Consumer Compensation Policy: fany defect is found, compensation, refund or exchange will be offered in accordance with the Consumer Dispute Resolution Standard of Fair Trade Commission.

\* Please refer the user guide and warnings

#### Manufactured for WINNOVA CO.,LTD 14-6, Gajang Industry dong-ro, Osan,

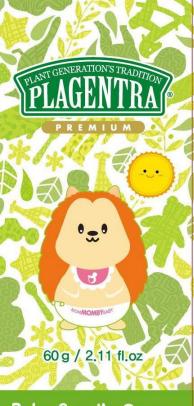
Distributed by C.A PHARM USA, INC 3435 Wilshine Blvd #2300 Los Angeles, CA 90010 Distributed by C.A PHARM CO.,LTD #1004. Gasan Digital Empire. 1130 Beoman -ro. Geumcheon-gu. Seoul. Korea.

+82-2-532-3478 www.capharm.net www.capharm.co.kr





Sun&Air pollution block



**Baby Suncity Cream** 



### PLAGENTRA BABY SUNCITY

octinoxate, octisalate, titanium dioxide cream

### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68988-090
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	4.5 g in 60 g	
OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	3 g in 60 g	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	1.8 g in 60 g	

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:68988-090-01	60 g in 1 CARTON		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	0 4/0 1/20 14	

# Labeler - C.A Pharm Co., Ltd. (688198385)

## **Registrant -** C.A Pharm Co., Ltd. (688198385)

Establishment				
Name	Address	ID/FEI	Business Operations	
C.A Pharm Co., Ltd.		688198385	manufacture(68988-090)	

Revised: 12/2014 C.A Pharm Co., Ltd.