GOLDKIWI SUN SPF 50 PLUS PA PLUS PLUS PLUS- octinoxate stick
SKINFOOD 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.

Drug Facts

Active ingredients: OCTINOXATE 7%, ZINC OXIDE 4.9%, TITANIUM DIOXIDE 4.15%

Inactive ingredients:
Dimethicone, Triethylhexanoin, Silica, Talc, Phenyl Trimethicone, Isoamyl p-Methoxycinnamate, Polyethylene, Ceresin, Sorbitan Isostearate, Microcrystalline Wax, Vinyl Dimethicone/Methicone Silsesquiox ane Crosspolymer, Aluminum Hydroxide, Stearic Acid, Triethoxycaprylylsilane, Fragrance(Parfum), Polyglyceryl-2 Triisostearate, Iron Oxides (CI 77492), Glyceryl Caprylate, Caprylyl Glycol, Iron Oxides (CI 77491), Actinidia Chinensis (Kiwi) Fruit Extract, Argania Spinosa Kernel Oil, Hydroxyapatite, Helianthus Annuus (Sunflower) Seed Oil

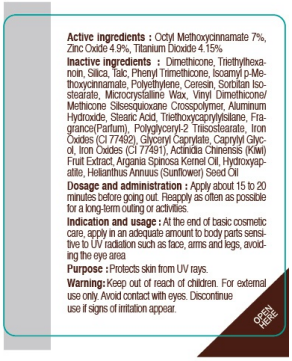
Purpose: Protects skin from UV rays.

Warnings:
For external use only.
Avoid contact with eyes.
Discontinue use if signs of irritation appear.

Keep out of reach of children:
Keep out of reach of children.

Indication and usage:
At the end of basic cosmetic care, apply in an adequate amount to body parts sensitive to UV radiation such as face, arms and legs, avoiding the eye area

Dosage and administration:
Apply about 15 to 20 minutes before going out.
Reapply as often as possible for a long-term outing or activities.



GOLDKIWI SUN SPF 50 PLUS PA PLUS PLUS PLUS

octinoxate stick

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76214-039
Route of Administration	CUTANEOUS		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	1.47 g in 21 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC OXIDE	1.03 g in 21 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM - UNII:D1JT611TNE)	TITANIUM DIOXIDE	0.87 g in 21 g

Inactive Ingredients	
Ingredient Name	**Strength**
DIMETHICONE (UNII: 92RU3N3Y1O)	
TRIETHYLHEXANOIN (UNII: 7K3W1BIU6K)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
CERESIN (UNII: Q1LS2UJO3A)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
GLYCERYL CAPRYLATE (UNII: TM2TZD4G4A)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
KIWI FRUIT (UNII: 71ES77LGJC)	
ARGAN OIL (UNII: 4V59G5UW9X)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:76214-039-01	21 g in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/01/2011	

Labeler - SKINFOOD CO., LTD. (690324173)

Registrant - SKINFOOD CO., LTD. (690324173)

Establishment

Name	Address	ID/FEI	Business Operations
SKINFOOD CO., LTD.		690324173	manufacture

Revised: 10/2011 SKINFOOD CO., LTD.