THE FIRST SUN SPF50 PLUS PA PLUS PLUS PLUS- octinoxate, zinc oxide, octisalate, titanium dioxide liquid
NATURE REPUBLIC 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%, ZINC OXIDE 6.5%, ETHYLHEXYL SALICYLATE 5%, TITANIUM DIOXIDE 2.5%

Inactive Ingredient

Inactive Ingredients:
WATER, CYCLOPENTASILOXANE, ALCOHOL, DICAPRYLYL CARBONATE, BIOSACCHARIDE GUM-1, BUTYLENE GLYCOL, PEG-10 DIMETHICONE, MAGNESIUM SULFATE, METHYL METHACRYLATE CROSSPOLYMER, PHENOXYETHANOL, METHICONE, ALUMINUM HYDROXIDE, ALUMINUM STEARATE, CAPRYLYL GLYCOL, GLYCERYL CAPRYLATE, CITRUS GRANDIS (GRAPEFRUIT) PEEL OIL, POLYGLYCERYL-6 POLYRICINOLEATE, SACCHAROMYCES FERMENT, OLEA EUROPAEA (OLIVE) FRUIT OIL, ACACIA SENEGAL FLOWER/STEM EXTRACT, CITRUS AURANTIFOLIA (LIME) OIL, RIBES NIGRUM (BLACK CURRANT) SEED OIL, ROSE FLOWER OIL, ILLICIUM VERUM (ANISE) FRUIT/SEED OIL, EUTERPE OLERACEA FRUIT EXTRACT, HYDROLYZED COLLAGEN, SODIUM LAUROYL SARCOSINATE, SOLUBLE COLLAGEN

Purpose

Purpose: This sun liquid, with its active ingredients of fermented enzyme, makes your skin look clear and transparent.

WARNINGS

Cautions:
For external use only.
Avoid contact with eyes and mouth.
Discontinue use if signs of irritation or rash appear
Replace the cap after use.

Keep out of reach of children

Keep out of reach of children:

Keep out of reach of children

INDICATION AND USAGE

How to Use:
Gently shake the bottle before use.
Apply an appropriate amount to your face, following the skin's texture and softly pat your face to enhance its absorption into your skin.

DOSAGE AND ADMINISTRATION

How to Use:
Gently shake the bottle before use.
Apply an appropriate amount to your face, following the skin's texture and softly pat your face to enhance its absorption into your skin.

PACKAGE LABEL. PRINCIPAL DISPLAY PANEL

THE FIRST SUN SPF50 PLUS PA PLUS PLUS PLUS

octinoxate, zinc oxide, octisalate, titanium dioxide liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51346-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	3 g in 40 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	2.61 g in 40 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	2 g in 40 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	1 g in 40 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
ALCOHOL (UNII: 3K9958V90M)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51346-001-01	40 mL in 1 CARTON		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/01/2012	

Labeler - NATURE REPUBLIC CO., LTD. (631172020)

Registrant - NATURE REPUBLIC CO., LTD. (631172020)

Establishment

Name	Address	ID/FEI	Business Operations
NATURE REPUBLIC CO., LTD.		631172020	manufacture(51346-001)

Revised: 8/2012 NATURE REPUBLIC CO., LTD.