

**CALIFORNIA ALOE FRESH POW DERY SUN- homosalate, octocrylene, ethylhexyl salicylate, butyl methoxydibenzoylmethane stick
NATURE REPUBLIC CO., LTD.**

California Aloe Fresh Powdery Sun Stick SPF 50

Drug Facts

Active Ingredients

HOMOSALATE 9.0%
OCTOCRYLENE 9.0%
ETHYLHEXYL SALICYLATE 4.5%
BUTYL METHOXYDIBENZOYLMETHANE 2.9%

Purpose

Sunscreen

Uses

Protects skin from direct sunlight and UV rays.

WARNINGS

For external use only. Avoid contact with eyes. In the event of contact with eyes, rinse thoroughly with water. Discontinue use if signs of irritation or rashes appear. Keep out of reach of children. Replace cap after use.

Keep out of reach of children.

Directions

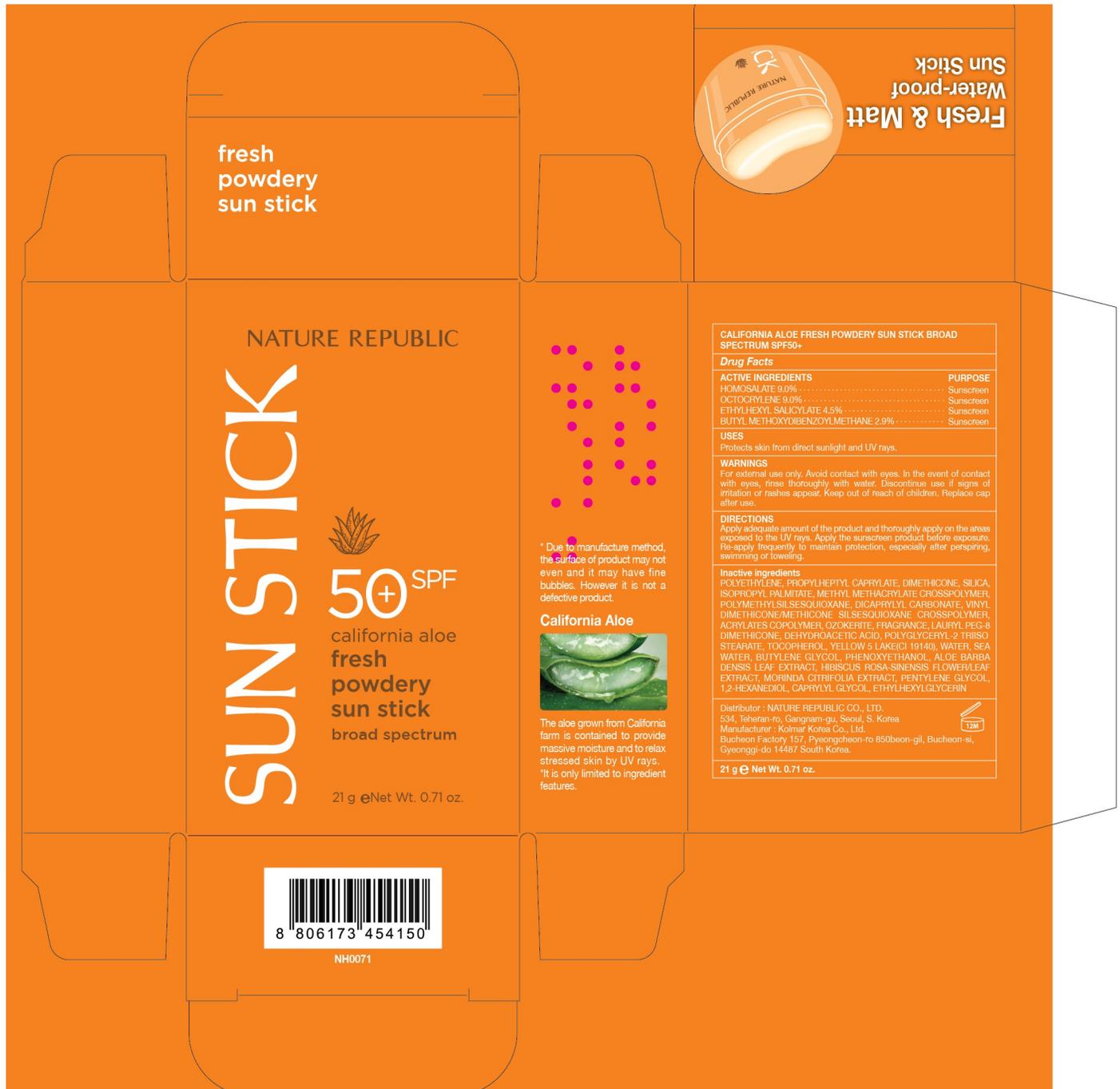
Apply adequate amount of the product and thoroughly apply on the areas exposed to the UV rays. Apply the sunscreen product before exposure. Re-apply frequently to maintain protection, especially after perspiring, swimming or toweling.

Inactive Ingredients

POLYETHYLENE, PROPYLHEPTYL CAPRYLATE, DIMETHICONE, SILICA, ISOPROPYL PALMITATE, METHYL METHACRYLATE CROSSPOLYMER, POLYMETHYLSILSESQUIOXANE, DICAPRYLYL CARBONATE, VINYL DIMETHICONE/METHICONE SILSESQUIOXANE CROSSPOLYMER, ACRYLATES COPOLYMER, OZOKERITE, FRAGRANCE, LAURYL PEG-8 DIMETHICONE, DEHYDROACETIC ACID, POLYGLYCERYL-2 TRIISOSTEARATE, TOCOPHEROL, YELLOW 5 LAKE(CI 19140), WATER, SEA WATER, BUTYLENE GLYCOL, PHENOXYETHANOL, ALOE BARBADENSIS LEAF EXTRACT, HIBISCUS ROSA-SINENSIS

FLOWER/LEAF EXTRACT, MORINDA CITRIFOLIA EXTRACT, PENTYLENE GLYCOL, 1,2-HEXANEDIOL, CAPRYLYL GLYCOL, ETHYLHEXYLGLYCERIN, LINALOOL, LIMONENE

PRINCIPAL DISPLAY PANEL



CALIFORNIA ALOE FRESH POW DERY SUN			
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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51346-532
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	1.89 g in 21 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	1.89 g in 21 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	0.94 g in 21 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.6 g in 21 g

Inactive Ingredients

Ingredient Name	Strength
PROPYLHEPTYL CAPRYLATE (UNII: 991Z19V2OD)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51346-532-01	21 g in 1 CARTON; Type 0: Not a Combination Product	08/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/01/2020	

Labeler - NATURE REPUBLIC CO., LTD. (631172020)**Registrant** - NATURE REPUBLIC CO., LTD. (631172020)**Establishment**

Name	Address	ID/FEI	Business Operations
Kolmar Korea Co.,Ltd. Bucheon Factory		963271763	manufacture(51346-532)

Revised: 1/2026

NATURE REPUBLIC CO., LTD.