

PURITO SUNDAY ADVENTURE SUNSCREEN SPF 50, 50 ML- avobenzone, homosalate, octisalate, octocrylene cream
Hinature

Purito Sun Day Adventure Sunscreen SPF 50

Drug Facts

Active ingredients

Avobenzone 3%
Homosalate 9%
Octisalate 4.5%
Octocrylene 5%

Purpose

Sunscreen

Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see Directions), decreases the risk of skin cancer and early skin aging caused by the sun

Warnings

For external use only.

Do not use

- on damaged or broken skin.

When using this product

- keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if

- rash occurs.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating
- **Sun Protection Measures.** Spending time in the sun increases your risk of skin

cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:

- limit time in the sun, especially from 10 a.m.-2 p.m.
- wear long-sleeved shirts, pants, hats, and sunglasses.
- Children under 6 months of age: Ask a doctor

Other information

- protect the product and this container from excessive heat and direct sun

Inactive ingredients

Water (Aqua), Dipropylene Glycol, Methyl Methacrylate Crosspolymer, Glyceryl Polymethacrylate, Propylene Glycol, Acrylates Copolymer, Butyloctyl Salicylate, Cetareth-20, Ethylhexyl Methoxycrylene, Polysorbate 60, Sorbitan Stearate, Caprylyl Glycol, Cetearyl Oliviate, Panthenol, Sorbitan Oliviate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Ethylhexylglycerin, Tromethamine, Sodium Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, 1,2-Hexanediol, Isohexadecane, Ammonium Acryloyldimethyltaurate/VP Copolymer, BHT, Butylene Glycol, Polysorbate 80, Trisodium Ethylenediamine Disuccinate, Sorbitan Oleate, Xanthan Gum, Java Waterdropwort (Oenanthe javanica) Extract

Questions or Comments?

research@purito.co.kr

Package Labeling:



Rooted
in
Wonder

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Questions or comments?

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MADE IN KOREA

TO REPORT A SERIOUS
ADVERSE EVENT, CONTACT
HINATURE INC.
C/O 144 RESEARCH DRIVE,
HAMPTON, VA, 23666

EXP YYYYMMDD



Purito

SEOUL



SUN DAY
ADVENTURE
SUNSCREEN

BROAD SPECTRUM
SPF 50

DAILY DEFENSE
& SOFT TOUCH

1.69 fl oz (50 mL)

Purito.com

Purito
SEOUL



Preview



PURITO SUNDAY ADVENTURE SUNSCREEN SPF 50, 50 ML

avobenzone, homosalate, octisalate, octocrylene cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73251-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	90 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (ETHYLHEXYL SALICYLATE - UNII:4X49Y0596W)	OCTISALATE	45 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
SORBITAN OLEATE (UNII: 06XEA2VD56)	

XANTHAN GUM (UNII: TTV12P4NEE)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
CETEARETH-20 (UNII: YRC528SWJY)	
ETHYLHEXYL METHOXYCRYLENE (UNII: S3KFG6Q5X8)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
PANTHENOL (UNII: WW9CM0O67Z)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
TROMETHAMINE (UNII: 023C2WHX2V)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
ISOHEXADECANE (UNII: 918X1OUF1E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73251-010-00	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/15/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/15/2025	

Labeler - Hinature (694862999)

Revised: 2/2025

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