

PETER ISLAND BABY SUNSCREEN SPF 50- homosalate, oxybenzone, octisalate, avobenzone, octocrylene lotion
AMWAY CORP.

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 Ingredients

Homosalate 13.0%
Octocrylene 7.0%
Octisalate 5.0%
Oxybenzone 4.0%
Avobenzone 3.0%

Warnings:

FOR EXTERNAL USE ONLY.

Avoid contact with eyes. Rinse with water if contact occurs.

Discontinue use if signs of rash or irritation develop.

For use on children under the age of 6 months consult a physician.

Keep out of reach of children.

Directions

Apply generously and evenly 30 minutes before sun exposure. Reapply frequently and after swimming, excessive perspiration and towel drying.

Other Information

- May stain some fabrics
- Sun Alert: Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

This formula provides Broad Spectrum UVA/UVB protection from the sun's damaging rays. It is PABA free, vitamin enriched and pediatrician tested.

Principal Display Panel

PETER ISLAND
baby Sunscreen lotion
spf 50
Pediatrician Tested
Photostable
Broad Spectrum
UVA/UVB Protection
Very Water Resistant
8 FL.OZ. (237 mL)

PETER
ISLAND

baby
sunscreen lotion



spf

50

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ACTIVE INGREDIENTS: Homosalate 13.0%, Octocrylene 7.0%, Octisalate 5.0%, Oxybenzone 4.0%, Avobenzone 3.0%.

INACTIVE INGREDIENTS: Water, Sorbitol, Stearic Acid, Triethanolamine, Aluminum Starch Octenylsuccinate, Benzyl Alcohol, Sorbitan Isostearate, VP/Eicosene Copolymer, Dimethicone, Polyglyceryl-3 Distearate, Carbomer, Tocopherol, Disodium EDTA, Methylparaben, Propylparaben, Fragrance.



Amway.

Dist. by Amway Corp.
Ada, MI 49355.

For Questions: 1-800-253-6500
amway.com

U0006-A

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50390-706
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	13 g in 100 g
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:950OS7VE0Y)	OXYBENZONE	4 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g

OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	7 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TOCOPHEROL (UNII: R0ZB2556P8)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50390-706-56	226 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	11/26/2012	

Labeler - AMWAY CORP. (083416854)

Revised: 11/2012

AMWAY CORP.