PETER ISLAND SUNSCREEN SPF 70- homosalate, oxybenzone, octisalate, avobenzone, octocrylene lotion
Access Business Group LLC

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 15.0%

Octocrylene 10.0%

Oxybenzone 6.0%

Octisalate 5.0%

Avobenzone 3.0%

WARNING:

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 vitamin enriched formula is very water resistant, fragrance free, PABA free and provides Broad Spectrum UVA/UVB protection.

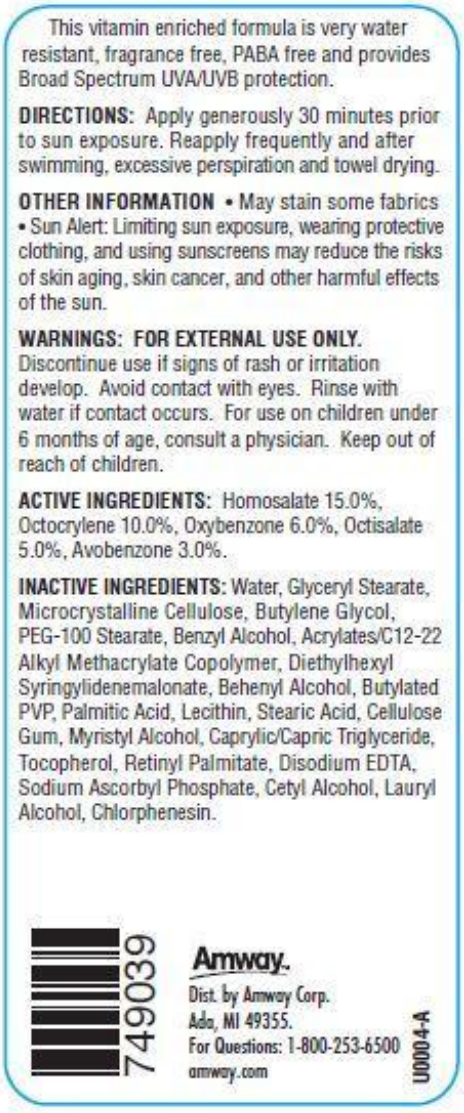
Principal Display Panel

PETER ISLAND

Sunscreen lotion
spf 70
Photostable
Broad Spectrum
UVA/UVB Protection
Very Water Resistant


PETER
ISLAND
Sunscreen
lotion
spf
70
Photostable
Broad Spectrum
UVA/UVB Protection
Very Water Resistant
8 FL. OZ. (237 mL)

8 FL.OZ. (237 mL)



INACTIVE INGREDIENTS:

Water, Glyceryl Stearate, Microcrystalline Cellulose, Butylene Glycol, PEG-100 Stearate, Acrylates/C12-22 Alkyl Methacrylate Copolymer, Benzyl Alcohol, Diethylhexyl Syringylidenemalonate, Behenyl Alcohol, Butylated PVP, Palmitic Acid, Stearic Acid, C12-16 Alcohols, Cellulose Gum, Caprylic/Capric Triglyceride, Lecithin, Retinyl Palmitate, Tocopherol, Sodium Ascorbyl Phosphate, Disodium EDTA, Chlorphenesin.

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10056-704
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	15 g in 100 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	6 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	10 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
DIETHYLHEXYL SYRINGYLIDENEMALONATE (UNII: 3V5U97P248)	
DOCOSANOL (UNII: 9G1OE216XY)	
PALMITIC ACID (UNII: 2V16EO95H1)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
LAURYL ALCOHOL (UNII: 178A96NLP2)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
PEG-100 STEARATE (UNII: YD01N1999R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRICAPRIN (UNII: O1PB8EU98M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:10056-704-56	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	06/28/2013	

Labeler - Access Business Group LLC (839830713)

Revised: 6/2013 Access Business Group LLC