PETER ISLAND ULTIMATE SHEER DRY TOUCH SUNSCREEN SPF 70- 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 15.0%
Oxybenzone 6.0%
Octisalate 5.0%
Avobenzone 3.0%
Octocrylene 2.8%

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.

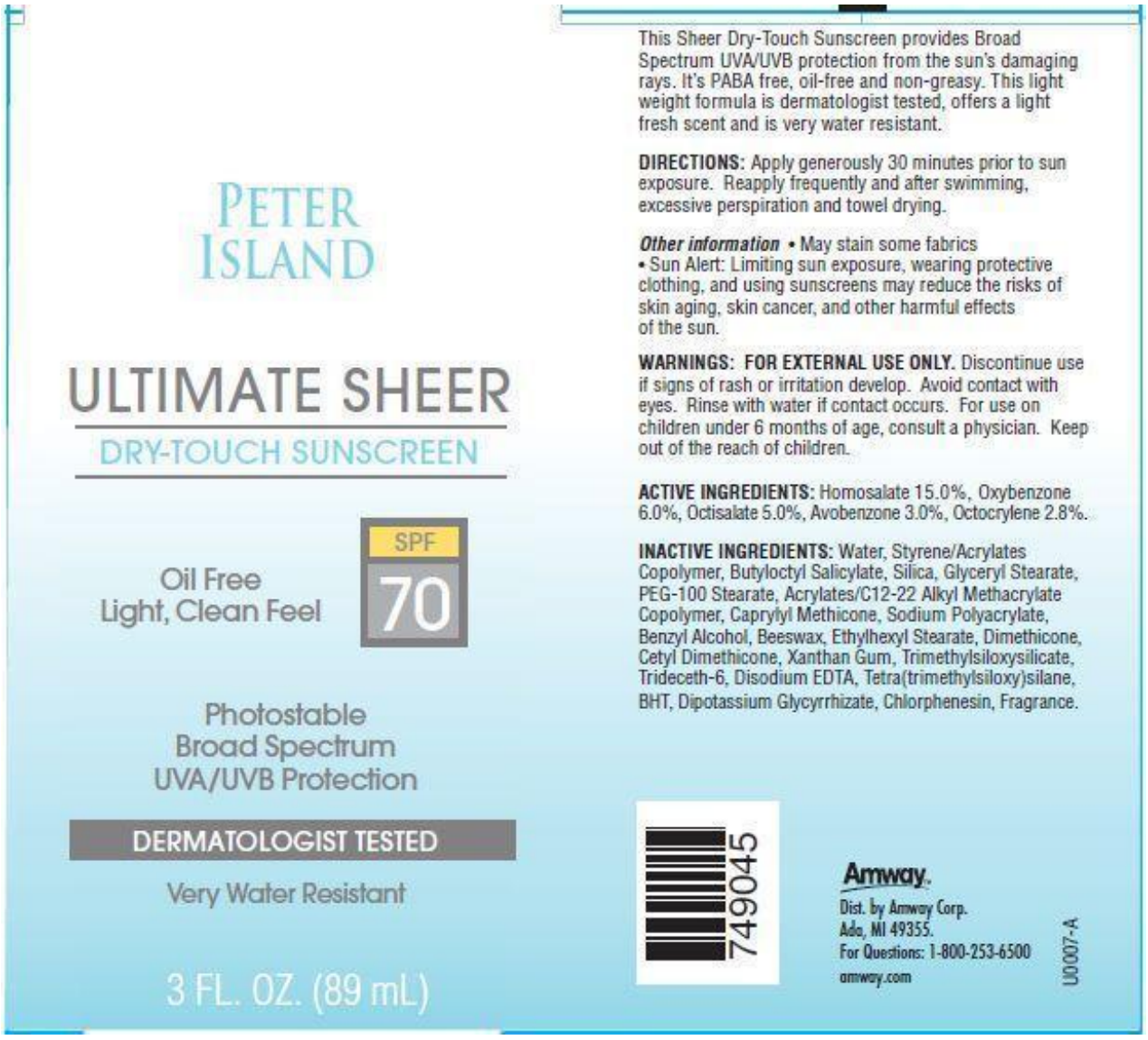
Directions
Apply generously and evenly 30 minutes before sun exposure. Reapply frequently and after swimming, excessive perspiration and towel drying.

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 Sheer Dry-Touch Sunscreen provides Broad Spectrum UVA/UVB protection from the sun's damaging rays. It's PABA free,oil-free and non-greasy. This light weight formula is dermatologist tested, offers a light fresh scent and is very water resistant.

Principal Display Panel
PETER ISLAND
ULTIMATE SHEER
DRY-TOUCH SUNSCREEN
Oil Free Light, Clean Feel
spf 70
Photostable
Broad Spectrum
UVA/UVB Protection
DERMATOLOGIST TESTED
Very Water Resistant

3 FL.OZ. (89 mL)



Inactive Ingredients
Water, Styrene/Acrylates Copolymer, Butyloctyl Salicylate, Glyceryl Stearate, Silica, PEG-100 Stearate, Acrylates/C12-22 Alkyl Methacrylate Copolymer, Caprylyl Methicone, Sodium Polyacrylate, Benzyl Alcohol, Beeswax, Ethylhexyl Stearate, Dimethicone, Cetyl Dimethicone, Xanthan Gum, Trimethylsiloxysilicate, Trideceth-6, Disodium EDTA, Dipotassium Glycyrrhizate, BHT, Chlorphenesin

PETER ISLAND ULTIMATE SHEER DRY TOUCH SUNSCREEN SPF 70
homosalate, oxybenzone, octisalate, avobenzone, octocrylene lotion

Product Information	

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50390-707
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	2.8 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
XANTHAN GUM (UNII: TTV12P4NEE)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TETRASILANE (UNII: OI9DXJ0BL4)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
GLYCYRRHIZINATE DIPOTASSIUM (UNII: CA2Y0FE3FX)	
CHLORPHENESIN (UNII: I670DAL4SZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50390-707-12	85 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.