

PETER ISLAND CONTINUOUS SPORT SUNSCREEN SPF 50 - homosalate, oxybenzone, octisalate, avobenzone, octocrylene spray

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%

Oxybenzone 6.0%

Octisalate 5.0%

Avobenzone 3.0%

Octocrylene 2.0%

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

FLAMMABLE

Do not spray near heat, flame or while smoking. Store at room temperature only. Use in well ventilated areas. Avoid long term storage above 40° C(104° F).

CONTENTS UNDER PRESSURE: Do not puncture or incinerate. Do not store at temperatures above 120° F.

DIRECTIONS

Do not spray directly on face. 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 light, dry formula was created for active adults. It provides water resistant, sweat resistant protection from the sun's UVA/UVB rays. Dries quickly and can be applied from any angle. PABA free.

Principal Display Panel

PETER ISLAND

Continuous Spray SPORT

Sunscreen

spf 50

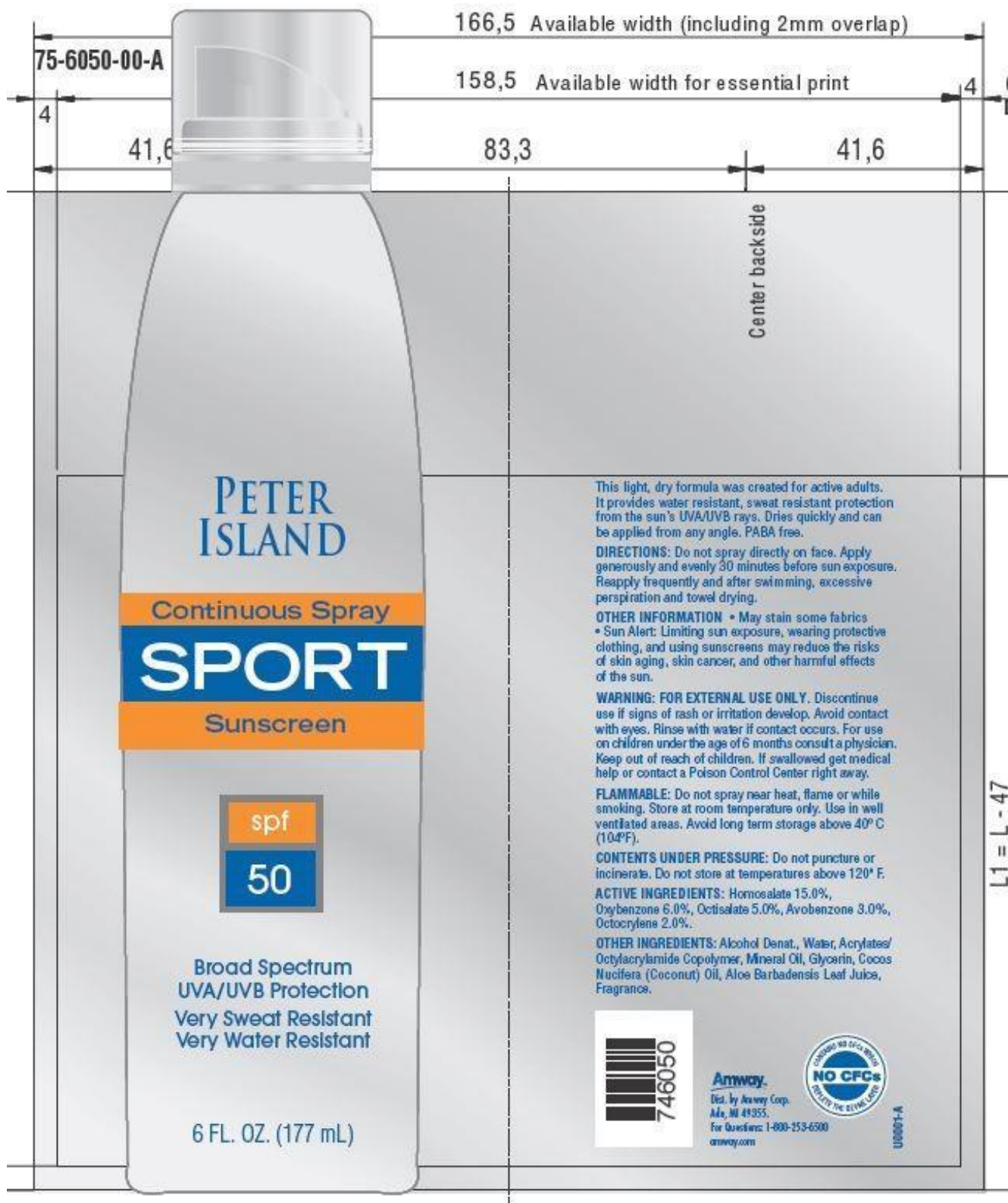
Broad Spectrum

UVA/UVB Protection

Very Sweat resistant

Very Water Resistant

6 FL.OZ. (177 mL)



OTHER INGREDIENTS:

Alcohol Denat., Water, Acrylates/Octylacrylamide Copolymer, Mineral Oil, Glycerin, Cocos Nucifera (Coconut) Oil, Aloe Barbadensis Leaf Extract, Fragrance.

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10056-701
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 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
MINERAL OIL (UNII: T5L8T28FGP)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCONUT OIL (UNII: Q9L0O73W7L)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10056-701-34	170 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	06/27/2013	

Labeler - Access Business Group LLC (839830713)

Revised: 6/2013

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