# SPF 50 STAY PUT SUNSCREEN- octinoxate, octisalate, oxybenzone, titanium dioxide lotion Sawyer Products

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.

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#### SPF 50 Stay Put Sunscreen

#### **Drug Facts**

#### **Active ingredients:**

Octinoxate 7.5%, Octisalate 5.0%, Oxybenzone 6.0%, Titanium Dioxide 2.6%

#### Purpose:

Sunscreen

#### Uses:

· Helps prevent sunburn.

#### Warnings:

#### **Skin Cancer/Skin Aging Alert**

• ||For external use only

#### Do not use

on broken or damaged skin.

#### Keep out of reach of children.

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

#### When using this product

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

#### Stop use and ask a doctor

if rash occurs.

#### **Directions:**

- · Apply liberally to cool, dry skin 15 minutes before sun exposure.
- · Use a water resistant sunscreen if swimming or sweating.
- · Reapply at least every 2 hours.
- · For children under 6 months of age: ask a doctor before using.

#### Other information:

· Protect this product container from excessive heat and direct sun.

#### **Inactive ingredients:**

Aloe Barbadensis Leaf Juice, Dicaprylyl Maleate, Neopentyl Glycol Diethylhexanoate (and) Neopentyl Glycol Diisostearate, Octyldodecyl Neopentanoate, PEG-30 Dipolyhydroxystearate, Phenoxyethanol (and) Methyl paraben (and) Ethylparaben (and) Butylparaben (and) Propyl paraben (and) Isobutylparaben, Poly ethylene, Poly quaternium 37 (and) Propylene Glycol Dicaprylate/Dicaprate (and) PPG-1 Trideceth-6, Water

#### Package Labeling:



SAWYER® STAY PUT® SUNSCREEN—SPF 50

# **Drug Facts**

Active ingredients: (Purpose: Sunscreen)

Octinoxate 7.5%, Octisalate 5.0%, Oxybenzone 6.0%, Titanium Dioxide 2.6%

## Uses:

· Helps prevent sunburn.

## Warnings:

Skin Cancer/Skin Aging Alert

Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to prevent sunburn, **not** skin cancer or early skin aging.

- ·For external use only
- **.∙Do not use** on broken or damaged skin.
- Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
  - When using this product keep out of eyes. Rinse with warm water to remove.
    - Stop use and ask a doctor if rash occurs.

### Directions:

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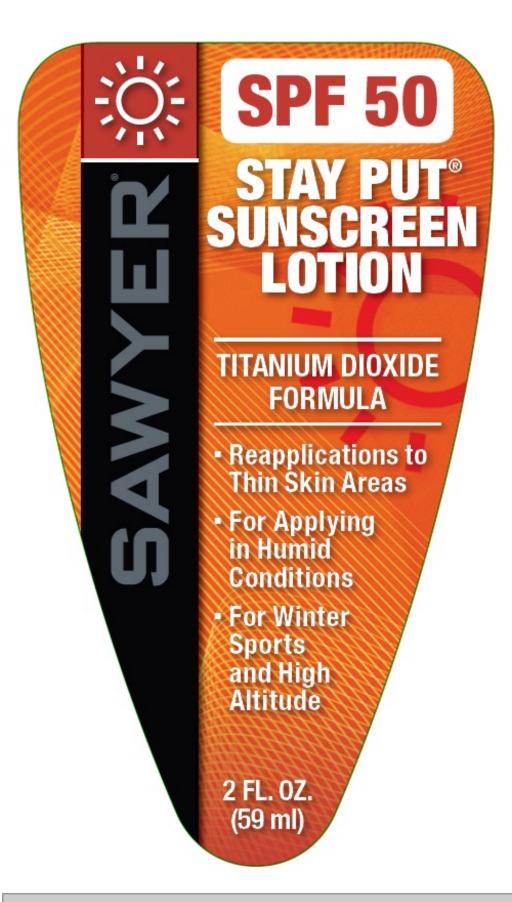
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> Item # SP1502 Distributed by: Sawyer Products, Inc. P.O. Box 188, Safety Harbor, FL 34695 www.sawyer.com 800-356-7811



#### **SPF 50 STAY PUT SUNSCREEN**

octinoxate, octisalate, oxybenzone, titanium dioxide lotion

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70392-014

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL		
OCTISALATE (UNII: 4X49 Y0 59 6 W) (OCTIS ALATE - UNII:4X49 Y0 59 6 W)	OCTISALATE	50 mg in 1 mL		
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	60 mg in 1 mL		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	26 mg in 1 mL		

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
DIO CTYL MALEATE (UNII: OD88G8439L)		
NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C)		
NEOPENTYL GLYCOL DIISOSTEARATE (UNII: 4M6 OQ34JWW)		
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)		
PEG-30 DIPOLYHYDRO XYSTEARATE (UNII: 9713Q0S7FO)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
METHYLPARABEN (UNII: A218 C7H19 T)		
ETHYLPARABEN (UNII: 14255EXE39)		
BUTYLPARABEN (UNII: 3QPI1U3FV8)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
ISOBUTYLPARABEN (UNII: 0 QQJ25X58G)		
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)		
PROPYLENE GLYCOL DICAPRYLATE (UNII: 581437HWX2)		
PROPYLENE GLYCOL DICAPRATE (UNII: U783H9JHWY)		
PPG-1 TRIDECETH-6 (UNII: 1K7417JX6Q)		
WATER (UNII: 059QF0KO0R)		

]	Packaging					
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date		
1	NDC:70392-014-01	1 in 1 CARTON				
1		59 mL in 1 BOTTLE; Type 0: Not a Combination Product				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part352	0 1/14/20 16		

## Labeler - Sawyer Products (118285923)

Revised: 1/2016 Sawyer Products