ULTA TANNING SPF 4 - octinoxate, padimate o lotion Ulta

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.

Drug Facts

Active ingredients

Octinoxate 2.0%, Padimate O 1.4%

Purpose

Sunscreen

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 aging.

For external use only

Do not use

on damaged or broken skin

When using this product

keep out of eyes. Rinse with water to remove

Stop use and ask a doctor

if rash occurs

Keep out of reach of childrem

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

Directions

- apply liberally 15 minutes before sun exposure
- reapply after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every2 hours
- children under 6 months of age: Ask a doctor

Inactive ingredients

Mineral Oil, Cocos Nucifera (Coconut) Oil, Theobroma Cacao (Cocoa) Seed Butter, Aloe Barbadensis Leaf Extract, Musa Sapientum (Banana) Fruit Extract, Retinyl Palmitate (and) Zea Mays (Corn) Oil (and) BHA (and) BHT, Tocopheryl Acetate, Daucus Carota Sativa (Carrot) Root Extract, Fragrance: Alpine Aromatics Coco Rum Banana #122-778, Phenoxyethanol (and) Methylparaben (and) Isopropylparaben (and) Butylparaben.

Other information

• protect this product from excessive heat and direct sun

Principal Display panel

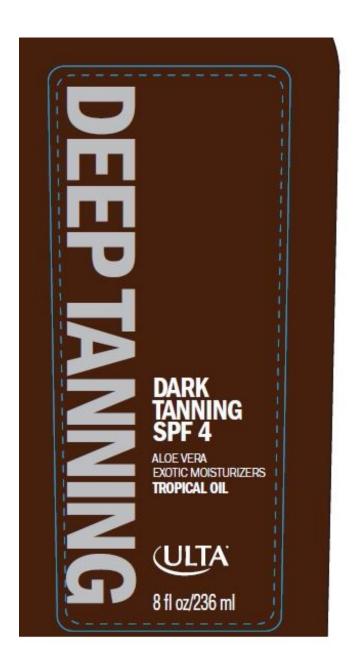
62296-7031-6

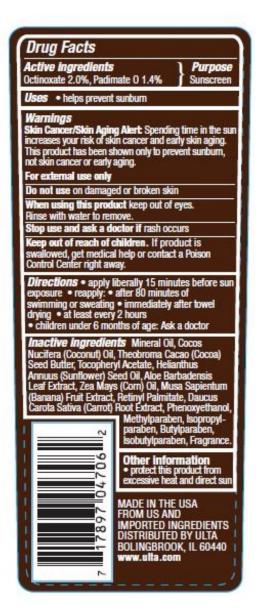
DEEP TANNING

DARK TANNING

SPF 4

ALOE VERA
EXOTIC MOISTURIZERS
TROPICAL OIL
ULTA
8 fl oz/ 236 ml





ULTA TANNING SPF 4

octinoxate, padimate o lotion

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:62296-7031

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	2 g in 100 g
PADIMATE O (UNII: Z11006CMUZ) (PADIMATE O - UNII:Z11006CMUZ)	PADIMATE O	1.4 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
MINERAL OIL (UNII: T5L8T28FGP)		
COCONUT OIL (UNII: Q9L0O73W7L)		
COCOA BUTTER (UNII: 512OYT1CRR)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
BANANA (UNII: 4AJZ4765R9)		
VITAMIN A PALMITATE (UNII: 1D1K0 N0 VVC)		
.ALPHATO COPHERO L ACETATE (UNII: 9E8X80D2L0)		
CARROT (UNII: L56Z1JK48B)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
METHYLPARABEN (UNII: A218 C7HI9T)		
ISOPROPYLPARABEN (UNII: A6EOX47QK0)		
ISOBUTYLPARABEN (UNII: 0QQJ25X58G)		
BUTYLPARABEN (UNII: 3QPI1U3FV8)		

l	Packaging					
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
l	1 NDC:62296-7031-6	236 g 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	08/03/2015		

Labeler - Ulta (608168597)

Revised: 8/2015 Ulta