

## **ULTA SUNSCREEN SPF 30 - avobenzone, homosalate, octocrylene 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.*

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### **Drug Facts**

#### **Active ingredients**

Avobenzone 1.8%, Homosalate 7.0%, Octocrylene 5.0%

#### **Purpose**

Sunscreen

#### **Uses**

- helps prevent sunburn
- if used as directed with other sun protection measures (see **Directions**), decreases the risk of skin cancer and early skin aging caused by the sun

#### **Warnings**

##### **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 children**

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

- May stain some fabrics.

#### **Directions**

- apply liberally 15 minutes before sun exposure
- reapply: after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours
- **Sun Protection Measures**. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a broad spectrum SPF of 15 or higher and other sun protection measures including:
- limit time in the sun, especially from 10 a.m. - 2 p.m.

- wear long-sleeve shirts, pants, hats, and sunglasses
- children under 6 months of age: Ask a doctor

### **Inactive Ingredients**

Water, Cetearyl Alcohol, Stearyl Alcohol, Glycerin, Phenoxyethanol, Acrylates/C12-22 Alkyl Methacrylate Copolymer, Caprylyl Glycol, Cetyl Alcohol, Carbomer, Ceteth-10 Phosphate, Dicaprylyl Phosphate, Coco-Glucoside, Methylparaben, Xanthan Gum, Propylparaben, Sodium Hydroxide, Disodium EDTA, Hydrogenated Methyl Abietate, Lauryl PEG-8 Dimethicone, Phenylisopropyl Dimethicone, Polyglyceryl-3 Stearate/Isostearate/Dimer Dilinoleate Copolymer, Sodium Ascorbyl Phosphate, Tocopheryl Acetate, Aloe Barbadensis Leaf Juice.

### **Other information**

- protect this product from excessive heat and direct sun

### **Principal Display Panel**

**62296-2273-2**

SUNCARE

**BROAD SPECTRUM SPF 30**

WATER RESISTANT

(80 MINUTES) OIL-FREE

**SUNSCREEN LOTION**

ULTA

3 fl oz/88 mL

**Moisturizing • Oil-Free • UVA/UVB Protection**

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MADE IN THE USA  
 FROM US AND IMPORTED INGREDIENTS  
 DISTRIBUTED BY ULTA  
 BOLINGBROOK, IL 60440  
 www.ulta.com

62296-2273-4

SUNCARE

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(80 MINUTES) OIL FREE

**SUNSCREEN LOTION**

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6 flo oz/176 mL

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**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:62296-2273
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	1.8 g in 100 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	5 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CETETH-10 PHOSPHATE (UNII: 4E05O5N49G)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
COCO GLUCOSIDE (UNII: ICS790225B)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYDROGENATED METHYL ABIETATE (UNII: A23O709X8O)	
SODIUM ASCORBYL PHOSPHATE (UNII: 836SJG51DR)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62296-2273-2	88 g in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:62296-2273-4	176 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	08/11/2015	

**Labeler** - Ulta (608168597)

Revised: 8/2015

Ulta