

**VACATION INSTANT BROWNING- avobenzone octisalate octocrylene lotion**  
**Vacation Inc.**

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**Vacation Instant Browning Lotion**

**Dosage and administration**

Apply liberally 15 minutes before sun exposure.

Reapply at least every 2 hours.

Use water-resistant sunscreen if swimming or sweating

**Warnings**

For external use only

Do not use on damaged or broken skin

When using this product, keep out of reach of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash occurs

In case of ingestion, get medical help or contact a Poison Control Center right away

**Inactives**

OCTISALATE, OCTOCRYLENE, ACETYL L-TYROSINE, LEVOMENOL, MAGNESIUM STEARATE, OCTYLDODECYL OLEATE, WATER, C13-15 ALKANE, PROPANEDIOL, COCOA BUTTER, TOCOPHEROL, VANILLA PLANIFOLIA OIL, DOCOSANOL, DECYL GLUCOSIDE, SODIUM METABISULFITE, PORPHYRIDIUM PURPUREUM, ALCOHOL, BUTYLOCTYL SALICYLATE, BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES, CAPRYLHYDROXAMIC ACID, CAPRYLYL GLYCOL, CETOSTEARYL ALCOHOL, CETYL ALCOHOL, CITRIC ACID MONOHYDRATE, COCO GLUCOSIDE, ARABICA COFFEE OIL, EUROPEAN HAZELNUT OIL, DIHYDROXYACETONE, DIMETHICONE, ETHYLENE BRASSYLATE, ETHYL FERULATE, GLYCERIN, FERRIC OXIDE RED, LAURYL GLUCOSIDE, PASSIFLORA EDULIS SEED OIL, CHASTE TREE

**Indications and Usage**

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.

**Keep out of reach of children**

Keep out of reach of children

In case of ingestion, get medical help or contact a Poison Control Center right away

# Purpose

Sunscreen

# Vacation Browning Lotion



## VACATION INSTANT BROWNING

avobenzene octisalate octocrylene lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:80641-521
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2.9 g in 100 mL
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	9.5 g in 100 mL
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.9 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
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<b>OCTYLDODECYL OLEATE</b> (UNII: MCA43PK7MH)
<b>WATER</b> (UNII: 059QF0KO0R)
<b>C13-15 ALKANE</b> (UNII: 114P5I43UJ)
<b>COCOA BUTTER</b> (UNII: 512OYT1CRR)
<b>TOCOPHEROL</b> (UNII: ROZB2556P8)
<b>VANILLA PLANIFOLIA OIL</b> (UNII: 0A3F415158)
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)
<b>BUTYLOCTYL SALICYLATE</b> (UNII: 2EH13UN8D3)
<b>BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES</b> (UNII: 0C9AC7D6XU)
<b>CAPRYLHYDROXAMIC ACID</b> (UNII: UPY805K99W)
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)
<b>COCO GLUCOSIDE</b> (UNII: ICS790225B)
<b>ARABICA COFFEE OIL</b> (UNII: IK55HKE887)
<b>EUROPEAN HAZELNUT OIL</b> (UNII: 8RQ2839AVG)
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)
<b>ETHYL FERULATE</b> (UNII: 5B8915UELW)
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)
<b>LAURYL GLUCOSIDE</b> (UNII: 76LN7P7UCU)
<b>PASSIFLORA EDULIS SEED OIL</b> (UNII: F3VOA31UHQ)
<b>TRihePTANOIN</b> (UNII: 2P6O7CFW5K)
<b>SODIUM GLUCONATE</b> (UNII: R6Q3791S76)
<b>APRICOT SEED OIL</b> (UNII: 54JB35T06A)
<b>SODIUM STEAROYL GLUTAMATE</b> (UNII: 65A9F4P024)
<b>HYALURONIC ACID</b> (UNII: S270N0TRQY)
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)
<b>ARACHIDYL ALCOHOL</b> (UNII: 1QR1QRA9BU)
<b>CARROT SEED OIL</b> (UNII: 595AO13F11)
<b>BISABOLOL</b> (UNII: 24WE03BX2T)
<b>HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL</b> (UNII: 3W1JG795YI)
<b>TRIMETHYLPENTANEDIOL/ADIPIC ACID/GLYCERIN CROSSPOLYMER (25000 MPA.S)</b> (UNII: 587WKM3S9Q)
<b>BETA-SITOSTEROL</b> (UNII: S347WMO6M4)
<b>SQUALENE</b> (UNII: 7QWM220FJH)
<b>BEHENYL ALCOHOL</b> (UNII: 9G1OE216XY)

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80641-521-10	148 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/06/2024	
2	NDC:80641-521-80	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/06/2024	

## Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M020	03/06/2024	

**Labeler** - Vacation Inc. (117644631)

Revised: 4/2025

Vacation Inc.