

**DROPS- avobenzone octisalate octocrylene liquid
Vacation Inc.**

Vacation Studio Tone Bronzing Drops

Dosage and administration

Shake well

Apply liberally 15 minutes before sun exposure

Reapply at least every 2 hours

Use a water resistant sunscreen if swimming or sweating

Warning

For external use only

Do not use on damaged or broken skin

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

Stop use and ask a doctor if rash occurs

Inactives

Butyloctyl Salicylate, C13-14 Alkane, Calcium Sodium Borosilicate, Caprylhydroxamic Acid, Caprylyl Glycol, Carbomer, Cellulose Gum, Ceramide AP, Ceramide EOP, Ceramide NP, Cetearyl Alcohol, Chlorella Vulgaris Extract, Cholesterol, Coco-Glucoside, Coffea Arabica (Coffee) Seed Extract , Corylus Avellana (Hazel) Seed Oil, Ethyl Ferulate, Ferulic Acid, Fragrance, Glycerin, Glyceryl Stearate, Helianthus Annuus (Sunflower) Seed Oil, Hyaluronic Acid, Iron Oxides, Mica, Microcrystalline Cellulose, Niacinamide

Octyldodecyl Oleate, Phytosphingosine, Propanediol, Sodium Benzoate, Sodium Gluconate, Sodium Lauroyl Lactylate

Sodium Stearoyl Glutamate, sr-Hydrozoan Polypeptide-1, Theobroma Cacao (Cocoa) Extract, Titanium Dioxide, Tocopherol, Tridecane

Trimethylpentanediol/Adipic Acid/Glycerin Crosspolymer, Undecane, Water, Xanthan Gum

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

Purpose

Sunscreen

Vacation Studio Tone Bronzing Drops



DROPS

avobenzone octisalate octocrylene liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80641-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2.9 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.9 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	7.6 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:80641-013-94	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/07/2024	
2	NDC:80641-013-80	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/07/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/07/2024	

Labeler - Vacation Inc. (117644631)

Revised: 3/2024

Vacation Inc.