BATH AND BODY WORKS SPF TAHITI ISLEavobenzone, homosalate, octisalate, octocrylene gel Bath & Body Works, Inc.

Bath & Body Works SPF 30 Gel To Oil Sunscreen Tahiti Isle

Active Ingredients

Avobenzone 3%, Homosalate 10%, Octisalate 5%, Octocrylene 10%

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 immediately.

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-sleeved shirts, pants, hats, and sunglasses • children under 6 months: Ask a doctor

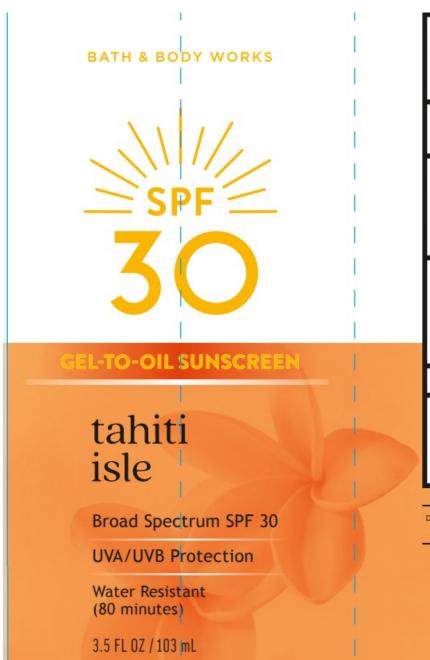
Inactive ingredients

INACTIVE INGREDIENTS: C12-15 Alkyl Benzoate, Caprylic/Capric Triglyceride, Polyamide-3, Isodecyl Neopentanoate, Polyester-8, Butyloctyl Salicylate, Diethylhexyl 2,6-Naphthalate, Diisopropyl Sebacate, Lauryl Lactate, Squalane, Simmondsia Chinensis (Jojoba) Seed Oil, Fragrance (Parfum), Diethylhexyl Syringylidenemalonate, Pentaerythrityl Tetra-di-t-butyl Hydroxyhydrocinnamate, Tocopheryl Acetate, Benzyl Salicylate, Hexyl Cinnamal, Hydroxycitronellal.

Other information

protect this product from excessive heat and direct sun Bath & Body Works, Distr.
95 West Main Street New Albany, OH 43054
1-800-395-1001
shop bathandbodyworks.com

Principal Display Panel



Drug Facts	
Active Ingredients	Purpose
Avobenzone (3%)	
Octisalate (596))	Sunscreen
Octocrylene (10%))	J
llsps	

• Helos prevent sunburn • If used as directed with other sun protection measures (see Directions), decreased the risk of skin cancer and early skin aging caused by the sun.

Warnings

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Do not use on damaged or broken skin.

When using this product • keep but of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash occurs.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

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Other Information

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

Inactive Ingredients

Macure myreoments.
12-15 alxyl benzoate, caprylic/capric triglyceride, polyamide-3, isodecyl nedpentanoate, polyaster B, butyloctyl salicylate, diethylhexyl 26-naphithalate, disdpropyl sebicate, lauryl lactate, squalane, simmondsia chinensis (jojoba) seed dil, fragranice (PARFUM), DIETHYLHEXYL SYRINGYLIDENEMALONATE, PENTAERYTHRITYL TETRA-DI-T-BUTYL HYDROXYHYDROCINNAMATE, TOCOPHERYL ACETATE, BENZYL SALICYLATE, HEXYL CINNAMAL, HYDROXYCITRONELIAL

DERMATOLOGIST TESTED

NOT TESTED & VEGAN

MADE WITHOUT ARTIFICIAL DYES

Bath & Body Works, Distr. 95 West Main Street New Albany, OH 43054 1-800-395-1001

BATH AND BODY WORKS SPF TAHITI ISLE

avobenzone,homosalate,octisalate,octocrylene gel

Product Information

NDC:62670-6864 **Product Type** HUMAN OTC DRUG **Item Code (Source)**

Route of Administration TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10 mg in 100 mg
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 mg in 100 mg
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII: V06SV4M95S)	HOMOSALATE	10 mg in 100 mg

OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y059

OCTISALATE

5 mg in 100 mg

Strength
S

WATER (UNII: 059QF0KO0R)

Pacl	kaging

7	tem Code	Package Description	Marketing Start Date	Marketing End Date
:	NDC:62670- 6864-0	103 mg in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2025	

Marketing Information

Marketing in	ioi iidcioii		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	05/05/2025	

Labeler - Bath & Body Works, Inc. (878952845)

Establishment			
Name	Address	ID/FEI	Business Operations
KDC/ONE Chatsworth, Inc.		118542196	manufacture(62670-6864)

Revised: 6/2025 Bath & Body Works, Inc.