BONDI SANDS BROAD SPECTRUM SPF 60 SUNNY CREAM BODY- avobenzone, homosalate, octisalate, octocrylene lotion
Baxter Laboratories Pty. Ltd.

Bondi Sands Broad Spectrum SPF 60 Sunny Cream Body Lotion

Active Ingredients

Avobenzone 3%

Homosalate 15%

Octisalate 5%

Octocrylene 10%

Purpose

Sunscreen

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

Stop use and ask a doctor if rash or irritation occurs.

- **For external use only**
- **Do not use** on damaged or broken skin
- **When using this product** keep out of eyes.If eye contact occurs, rinse thoroughly with clean running water.

Directions

- Shake well before use
- Apply liberally and evenly 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 value of 15 or higher and other sub 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

Water/Aqua/Eau, Beeswax, Aloe Barbadensis Leaf Juice, Cetearyl Alcohol, Cyclopentasiloxane, Isopropyl Palmitate, Ceteareth-20,

Hibiscus Sabdariffa Fruit Extract, Cyclohexasiloxane, Benzyl Alcohol, Hydroxyacetopenone, Phenoxyethanol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer,

Sodium Stearoyl Glutamate, Aminomethyl Propanediol, Tocopheryl Acetate, Sodium Carrageenan, Sodium Chloride.

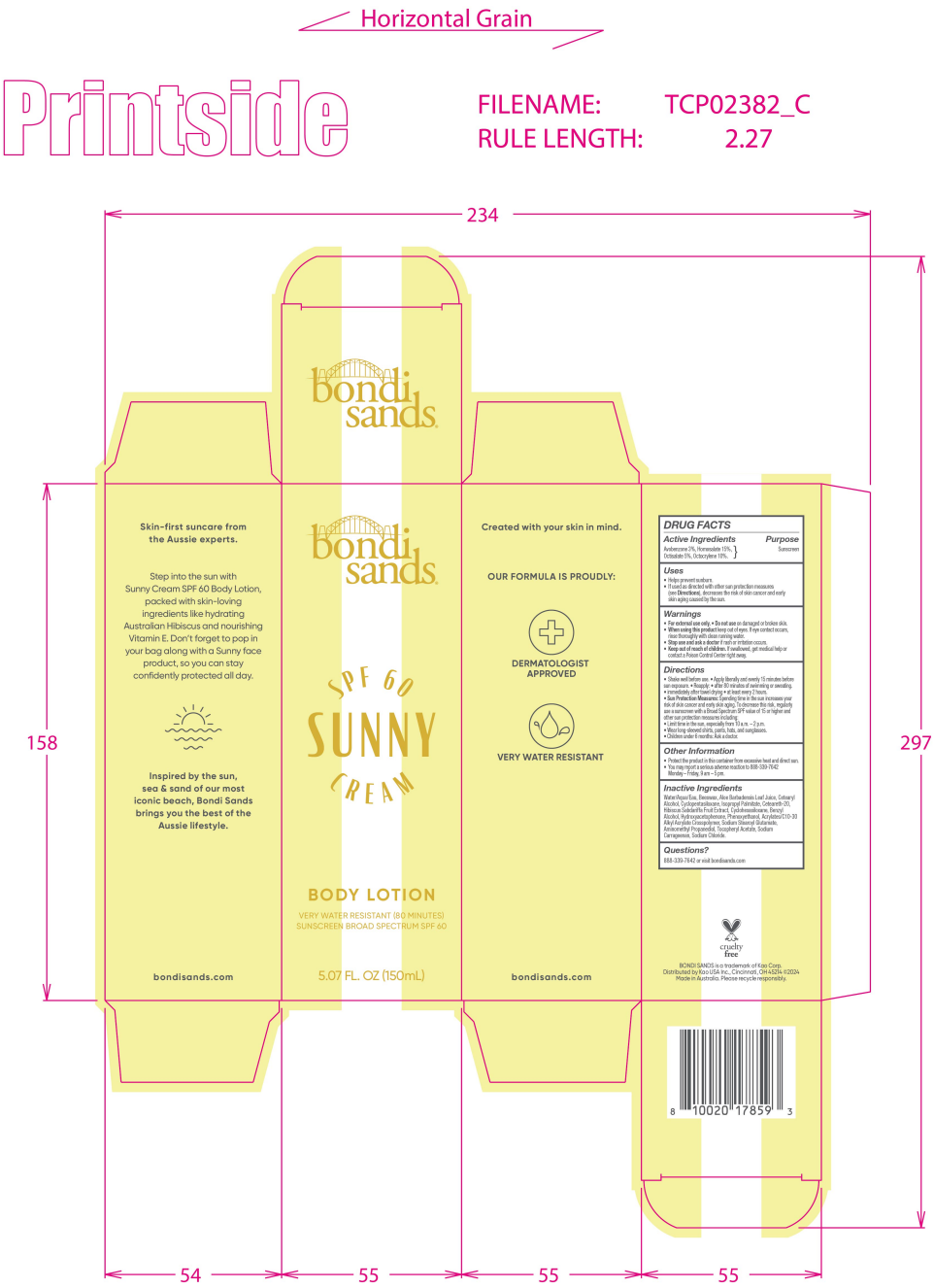
Bondi Sands

SPF 60

Sunny Cream

Body Lotion

5.07 FL. OZ (150mL)



BONDI SANDS BROAD SPECTRUM SPF 60 SUNNY CREAM BODY

avobenzone, homosalate, octisalate, octocrylene lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70157-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	15 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BEESWAX (UNII: 2ZA36H0S2V)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
CYCLOHEXASILOXANE (UNII: XHK3U310BA)	
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	
CETEARETH-20 (UNII: YRC528SWJY)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CARRAGEENAN (UNII: 7CY8BVL34N)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
HIBISCUS SABDARIFFA WHOLE (UNII: UH3Z91Y49Y)	
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)	
ALOE BARBADENSIS LEAF JUICE (UNII: ZY81Z83H0X)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70157-021-02	1 in 1 CARTON	12/03/2024	
1	NDC:70157-021-01	150 mL in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/03/2024	

Labeler - Baxter Laboratories Pty. Ltd. (740537709)

Revised: 12/2024 Baxter Laboratories Pty. Ltd.