

**CABANACLEAR ILLUMINATING SPF 50- homosalate, octisalate,
avobenzone lotion
NAKED SUNDAYS PTY LTD**

CabanaClear Illuminating

Active Ingredients (% W/W): Avobenzone 3%, Homosalate 6%, Octyl Salicylate 5%

Sunscreen

Use:

- Helps prevent sunburn.
- If used as directed with other sun protection measures (see Directions), decreases the risk of skin cancer and early skin ageing 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

Directions:

- Shake well before use.
- Apply liberally and evenly 20 minutes before sun exposure.
- Re-apply every 2 hours or immediately after swimming, sweating or towel drying.
- Use a water-resistant sunscreen if swimming or sweating.

Children under 6 months of age: Ask a doctor

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 sun protection measures including:

- Limit time in the sun, especially from 10.00AM - 2.00PM
- Wear long-sleeved shirts, pants, hats, and sunglasses

Other Information:

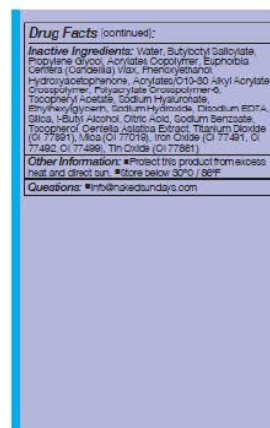
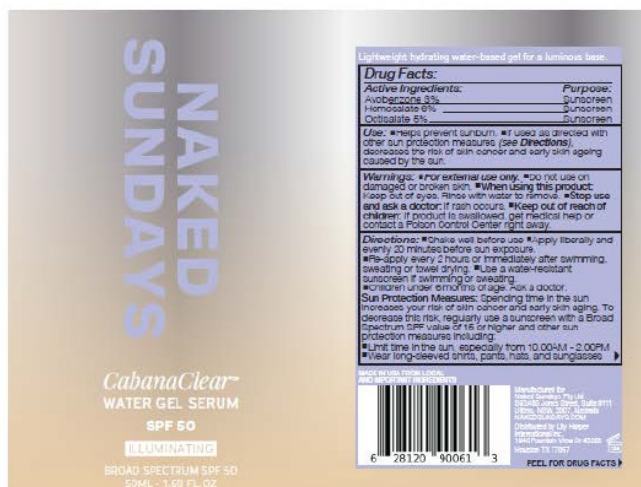
- Protect this product from excess heat and direct sun.
- Store below 30°C / 86°F

Questions:

- info@nakedSundays.com

Inactive Ingredients: Water, Butyloctyl Salicylate, Propylene Glycol, Acrylates Copolymer, Euphorbia Cerifera (Candelilla) Wax, Phenoxyethanol,

Hydroxyacetophenone, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Polyacrylate Crosspolymer-6, Tocopheryl Acetate, Sodium Hyaluronate, Ethylhexylglycerin, Sodium Hydroxide, Disodium EDTA, Silica, t-Butyl Alcohol, Citric Acid, Sodium Benzoate, Tocopherol, Centella Asiatica Extract, Titanium Dioxide (CI 77891), Mica (CI 77019), Iron Oxide (CI 77491, CI 77492, CI 77499), Tin Oxide (CI 77861)



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81104-594
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	6 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
SILICA (UNII: ETJ7Z6XBU4)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
EUPHORBIA CERIFERA (CANDELILLA) WAX (UNII: WL0328HX19)	

EDETATE DISODIUM (UNII: 7FLD91C86K)	
MICA (UNII: V8A1AW0880)	
CI 77491 (UNII: 1K09F3G675)	
CI 77492 (UNII: EX438O2MRT)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CITRIC ACID (UNII: 2968PHW8QP)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
POLYACRYLATE CROSSPOLYMER-6 (UNII: Q7UI015FF9)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BUTYL ACRYLATE/METHYL METHACRYLATE/METHACRYLIC ACID COPOLYMER (18000 MW) (UNII: JZ1374NL9E)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
CENTELLA ASIATICA TRITERPENOIDS (UNII: 4YS74Q4G4J)	
CI 77891 (UNII: 15FIX9V2JP)	
CI 77499 (UNII: XM0M87F357)	
TIN OXIDE (UNII: KM7N50LOS6)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81104-594-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/19/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	12/19/2025	

Labeler - NAKED SUNDAYS PTY LTD (746918845)