SUNRIGHT 50- avobenzone, homosalate, and octinoxate lotion
NSE Products, Inc.

Sunright® 50

Drug Facts

Active Ingredient

Avobenzone 3.0%, Homosalate 15.0%, Octisalate 5.0%.

Purpose

Sunscreen

Uses

- Helps prevent sunburn. Higher SPF gives more sunburn protection.
- 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.

Warning

- **For external use only.** Do not use on broken or damaged skin. Stop use and ask a doctor if rash occurs. When using this product, keep out of eyes. Rinse with water to remove.

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

Directions

- Apply liberally 15 minutes before sun or water exposure.
- Reapply: After 80 minutes of swimming or sweating
  - Immediately after towel drying
  - At least every 2 hours
- Children under 6 months: Ask a doctor

Inactive Ingredients

Water (Aqua), Aloe Barbadensis Leaf Juice, Acrylates Copolymer, Butyloctyl Salicylate, Propanediol, Caprylyl Methicone, Pentylene Glycol, Triacontanyl PVP, Potassium Cetyl Phosphate, Haematococcus Pluvialis Extract, Physalis Angulata Extract, Allantoin, Bisabolol, Panthenol, Tocopherol, Caprylic/Capric Triglyceride, VP/Acrylates/Lauryl Methacrylate Copolymer, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Acrylic Acid/VP Crosspolymer, Fragrance (Parfum), Hydroxyacetophenone, Sodium Dehydroacetate, Aminomethyl Propanol, Disodium EDTA.

Questions?

1-800-487-1000

PRINCIPAL DISPLAY PANEL - 100 mL Tube Label

NU SKIN®

SUNRIGHT® 50

BROAD SPECTRUM
SPF 50

FACE & BODY SUNSCREEN
WATER RESISTANT 80 MINUTES

100 mL ℮ 3.4 Fl Oz

SUNRIGHT 50

avobenzone, homosalate, and octinoxate lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62839-2011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Avobenzone (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	30 mg in 1 mL
Homosalate (UNII: V06SV4M95S) (Homosalate - UNII:V06SV4M95S)	Homosalate	150 mg in 1 mL
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
Butyloctyl Salicylate (UNII: 2EH13UN8D3)	
Propanediol (UNII: 5965N8W85T)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
Pentylene Glycol (UNII: 50C1307PZG)	
TRICONTANYL POVIDONE (UNII: N0SS3Q238D)	
Potassium Cetyl Phosphate (UNII: 03KCY6P7UT)	
Hydroxyacetophenone (UNII: G1L3HT4CMH)	
Sodium Dehydroacetate (UNII: 8W46YN971G)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
PHYSALIS ANGULATA WHOLE (UNII: W4TKW9D5GG)	
LEVOMENOL (UNII: 24WE03BX2T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
Allantoin (UNII: 344S277G0Z)	
Panthenol (UNII: WV9CM0O67Z)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
HAEMATOCOCCUS PLUVIALIS (UNII: 31T0FF0472)	
Tocopherol (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62839-2011-1	100 mL in 1 TUBE; Type 0: Not a Combination Product	05/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M020	05/01/2021	

Labeler - NSE Products, Inc. (803486393)

Revised: 4/2026 NSE Products, Inc.