

ANTIOXIDANT DAILY FACE PROTECTANT SPF 33- avobenzene, octinoxate, octisalate, oxybenzone cream
Ibr Packaging, LLC

Drug Facts

Active Ingredients	Purpose
Avobenzene 1%	Sunscreen
Octinoxate 7.5%	Sunscreen
Octisalate 5%	Sunscreen
Oxybenzone 6%	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 eye. 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

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 value 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

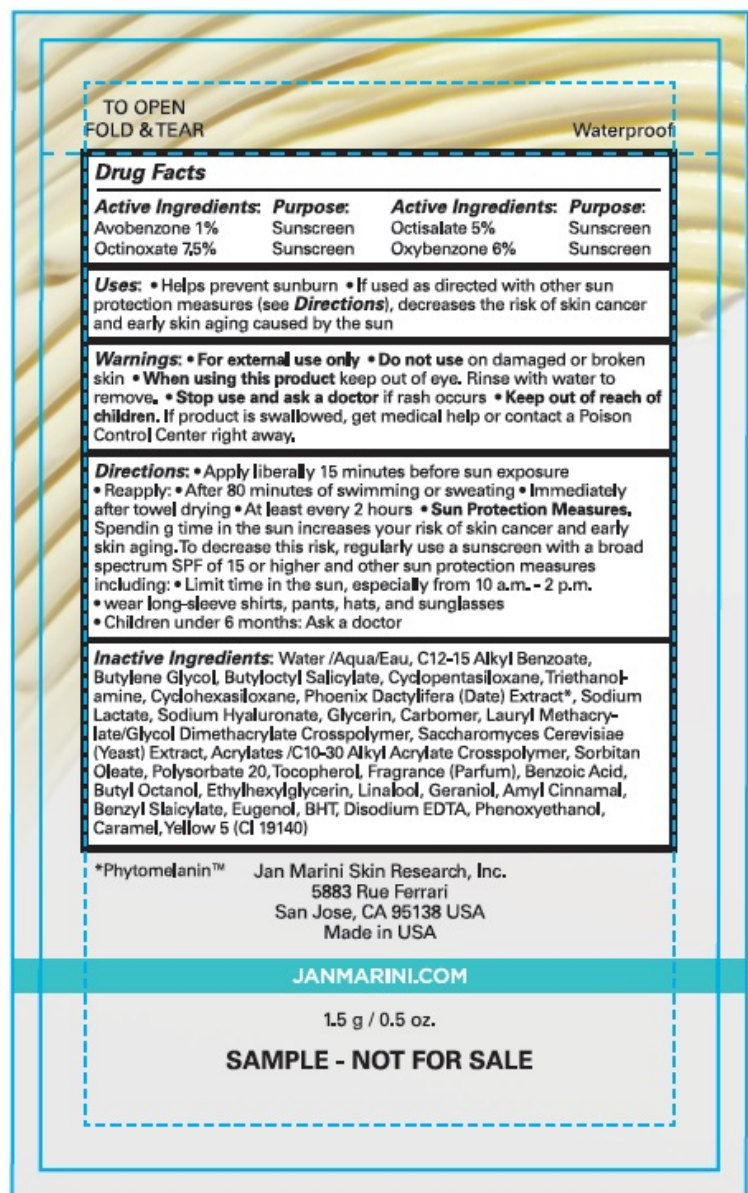
Children under 6 months of age: Ask a doctor

Inactive ingredients

Water/Aqua/Eau, C12-15 Alkyl Benzoate, Butylene Glycol, Butyloctyl Salicylate, Cyclopentasiloxane, Triethanol-amine, Cyclohexasiloxane, Phoenix Dactylifera (Date) Extract*, Sodium Lactate, Sodium Hyaluronate, Glycerin, Carbomer, Lauryl Methacrylate, /Glycol Dimethacrylate Crosspolymer, Saccharomyces Cerevisiae (Yeast) Extract, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Sorbitan Oleate, Polysorbate 20, Tocopherol, Fragrance (Parfum), Benzoic Acid, Butyl Octanol, Ethylhexylglycerin, Linalool, Geraniol, Amyl Cinnamal, Benzyl Salicylate, Eugenol, BHT, Disodium EDTA, Phenoxyethanol, Caramel, Yellow 5 (CI 19140)

*Phytomelanin™

Product label



ANTIOXIDANT DAILY FACE PROTECTANT SPF 33

avobenzone, octinoxate, octisalate, oxybenzone cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75587-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	1 g in 100 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	6 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
TOCOPHEROL (UNII: R0ZB2556P8)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
WATER (UNII: 059QF0KO0R)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
TROLAMINE (UNII: 9O3K93S3TK)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
LAURYL METHACRYLATE/GLYCOL DIMETHACRYLATE CROSSPOLYMER (UNII: EX0F4CZ66H)	
YEAST, UNSPECIFIED (UNII: 3NY3SM6B8U)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
GERANIOL (UNII: L837108USY)	
.ALPHA.-AMYL CINNAMALDEHYDE (UNII: WC51CA3418)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
EUGENOL (UNII: 3T8H1794QW)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CARAMEL (UNII: T9D99G2B1R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	

BUTYLOCTANOL (UNII: N442D9VO79)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75587-013-01	1.5 g in 1 PACKAGE; Type 0: Not a Combination Product	09/09/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	09/09/2024	

Labeler - lbr Packaging, LLC (081060205)

Revised: 9/2024

lbr Packaging, LLC