

**NEXGEN SUNSCREEN BROAD SPECTRUM SPF 36 DOC MARTINS- octinoxate, oxybenzone, octocrylene, avobenzone cream
Cosmetic Enterprises Ltd.**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

NexGen Sunscreen Broad Spectrum SPF 36

☐Active Ingredients:

Purpose:

Octinoxate 7.0% Benzophenone-3 ☐**Sunscreen**
(Oxybenzone) 5.0% Octocrylene 4.0% Avobenzone 3.0%

☐Uses:

Helps prevent sunburn. If used as described with other sun protection measures (see directions), decreases the risk of skin cancer and early skin aging caused by the sun.

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

Stop use and ask a doctor if rash occurs.

☐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.

☐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.
- Children under 6 months: ask a doctor.

Beeswax, Caprylic/Capric Triglyceride, Cetyl Alcohol, Cetyl Dimethicone, Cetyl Lactate, Cpernicia Cerifera (Carnauba) Wax, C10-30 Cholesterol/Lanosterol, Euphorbia Cerifera (Candelilla) Wax, Fragrance, Neopentyl Glycol Diheptanoate, Phenoxyethanol, Polybutene, Propylene Carbonate, Propylene Glycol Dibenzoate, Sea Whip Extract, Silica, Stearalkonium Hectorite

Doc Martin's of Maui

NexGen Sunscreen

New Skin Bonding Technology

Broad Spectrum SPF 36

Water Resistant (80 minutes)

2.0 FL OZ (60 mL)



**Doc Martin's
OF MAUI™**

NexGen™ Sunscreen
NEW SKIN BONDING TECHNOLOGY

FRONT
PANEL

Broad Spectrum SPF 36
Water Resistant (80 minutes)

BEND PT.

DEVELOPED BY MAUI DERMATOLOGIST "DOC" MARTIN
2.0 FL OZ (60 mL)

Graphics area-color ok / text not recommended

Drug Facts

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Inactive Ingredients:

Beeswax, Caprylic/Capric Triglyceride, Cetyl Alcohol, Cetyl Dimethicone, Cetyl Lactate, Copernicia Cerifera (Carnauba) Wax, C10-30 Cholesterol / Lanosterol, Euphorbia Cerifera (Candelilla) Wax, Fragrance, Neopentyl Glycol Diheptanoate, Phenoxyethanol, Polybutene, Propylene Carbonate, Propylene Glycol Dibenzoate, Sea Whip Extract, Silica, Stearalkonium Hectorite.

Protect this product from excessive heat and direct sun.

Other Information:

Sunscreen may stain or damage some fabrics/materials or surfaces. PABA Free. Reef Safe.

Questions? (808) 875-8888

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DocMartinsMaui.com



Void of Ink

NEXGEN SUNSCREEN BROAD SPECTRUM SPF 36 DOC MARTINS

octinoxate, oxybenzone, octocrylene, avobenzone cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56 152-50 10
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7 g in 100 mL
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:950OS7VE0Y)	OXYBENZONE	5 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	4 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
YELLOW WAX (UNII: 2ZA36H0S2V)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CETYL LACTATE (UNII: A7EVH2RK4O)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
C10-30 CHOLESTEROL/LANOSTEROL ESTERS (UNII: 137SL7IL0Y)	
CANDELILLA WAX (UNII: WL0328HX19)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYBUTENE (1400 MW) (UNII: 1NA5AO9GH7)	
PROPYLENE CARBONATE (UNII: 8D08K3S51E)	
PROPYLENE GLYCOL DIBENZOATE (UNII: EQY32Z1AN4)	
PSEUDOPTEROGORGIA ELISABETHAE (UNII: UDY3H10UX5)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARALKONIUM HECTORITE (UNII: O LX698AH5P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56152-5010-1	60 mL in 1 TUBE; Type 0: Not a Combination Product	03/21/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	03/21/2017	

Labeler - Cosmetic Enterprises Ltd. (017701475)

Registrant - Cosmetic Enterprises Ltd. (017701475)

Establishment

Name	Address	ID/FEI	Business Operations
Cosmetic Enterprises Ltd.		017701475	manufacture(56152-5010)