

DERMA SOLEIL NATURAL AGE-FREE FACIAL SUNBLOCK SPF 30- octocrylene, octyl methoxycinnamate, octyl salicylate, oxybenzone lotion
Deserving Health International Corp.

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.

DERMA SOLEIL NATURAL AGE-FREE FACIAL SUNBLOCK SPF 30

Octocrylene 4%
Octyl Methoxycinnamate 7.5%,
Octyl Salicylate 4%,
Oxybenzone 5%

Sunscreen

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.

For external use only.

Do not use on damaged or broken 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 swallowed, get medical help or contact a poison control center right away.

Aloe Vera Gel, C12-15 Alkyl Benzoate, Aqua and Glycerin and Sodium Levulinate and Sodium Anisate, Fragrance, Acrylates/C10-30 Alkyl Acrylates Crosspolymer, Aminomethyl Propanol, Carbomer, Titanium Dioxide, Hydroxypropyl Methylcellulose, Tetrasodium EDTA, Evening Primrose Oil, Avocado Oil, Jojoba Oil, Grape Seed Oil, Blue Chamomile Oil, Calendula Oil, Hydrolyzed Wheat Protein, Retinyl Palmitate, Ascorbic Acid, Tocopherol Acetate.

Natural Age-Free Facial Sunblock

Broad Spectrum SPF 30+

80% natural

UVA / UVB Protection
Water Resistant (40 minutes)
PABA Free

For All Skin Types

e120mL Net 4.0 fl. oz

Age-Free Facial Sunblock Broad Spectrum SPF 30+

Drugs Facts	
Active Ingredient	Purpose
Octocrylene 4 %	Sunscreen
Octyl Methoxycinnamate 7.5 %	Sunscreen
Octyl Salicylate 4 %	Sunscreen
Oxybenzone 5 %	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 eyes. Rinse with water to remove. • **Stop use and ask a doctor** if rash occurs • **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply liberally 15 minutes before sun exposure • Reapply: after 40 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: Aloe Vera Gel, C12-15 Alkyl Benzoate, Aqua and Glycerin and Sodium Levulinate and Sodium Anisate, Fragrance, Acrylates/C10-30 Alkyl Acrylates Crosspolymer, Aminomethyl Propanol, Carbomer, Titanium Dioxide, Hydroxypropyl Methylcellulose, Tetrasodium EDTA, Evening Primrose Oil, Avocado Oil, Jojoba Oil, Grape Seed Oil, Blue Chamomile Oil, Calendula Oil, Hydrolyzed Wheat Protein, Retinyl Palmitate, Ascorbic Acid, Tocopherol Acetate.

Questions? Call toll-free from US & CA +1-844-881-2882 or visit www.dermaed.com.

Distributed in the US by: Biological Health Group

Made in Canada

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octocrylene, octyl methoxycinnamate, octyl salicylate, oxybenzone lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69711-905
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYBENZONE (UNII: 95OOS7VE0 Y) (OXYBENZONE - UNII:95OOS7VE0 Y)	OXYBENZONE	5 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6 WM) (OCTOCRYLENE - UNII:5A68WGF6 WM)	OCTOCRYLENE	4 g in 100 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 mL
OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	4 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE SODIUM (UNII: MP1J8420LU)	
JOJOBA OIL (UNII: 724GKU717M)	
GRAPE SEED OIL (UNII: 930MLC8XGG)	
EVENING PRIMROSE OIL (UNII: 3Q9L08K71N)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
SODIUM ANISATE (UNII: F9WFJ28MV9)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
HYDROLYZED WHEAT PROTEIN (ENZYMATIC; 3000 MW) (UNII: J2S07SB0YL)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
SODIUM LEVULINATE (UNII: VK44E1MQU8)	
AVOCADO OIL (UNII: 6VNO72PFC1)	
CHAMOMILE FLOWER OIL (UNII: 60F80Z61A9)	
GLYCERYL CAPRYLATE (UNII: TM2TZD4G4A)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0K00R)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69711-905-01	120 mL in 1 TUBE; Type 0: Not a Combination Product	02/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	02/21/2020	

Labeler - Deserving Health International Corp. (079767886)

Establishment

Name	Address	ID/FEI	Business Operations

Revised: 2/2020

Deserving Health International Corp.