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.

DERMA SOLEIL NATURAL AGE-FREE FACIAL SUNBLOCK SPF 30

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: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	5 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	4 g in 100 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	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: 059QF0KO0R)	
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

Deserving Health International Corp.		202617023	manufacture(69711-905)