

DAILY FACIAL MOISTURIZER SPF 25- sunscreen cream

Person and Covey

DML Daily Facial Moisturizer

Uses

- Moisturizes and helps to relieve irritated and sensitive skin
- Higher SPF gives more sunburn protection
- Water resistant

Keep Out of the Reach of Children

If swallowed, get medical help or contact a Poison Control Center right away.

Warnings

For external use only

.Stop Use

Stop Use and ask a Doctor if a rash rash or irritation develops and lasts

Questions

Questions? Please call 1 (800) 423-2341

Inactive Ingredients

Purified Water, Propylene Glycol Dioctanoate, Petrolatum, Glycerin, Glyceryl Stearate, PEG-100 Stearate, DEA-Cetyl Phosphate, Stearic Acid, Hyaluronic Acid, Benzyl Alcohol, Dimethicone, Cyclomethicone, PVP/Eicosene Copolymer, Silica, Sodium Carbomer 941, Allantoin, Disodium EDTA, Magnesium Aluminum Silicate

Other Information

- High sun protection product
- Sun Alert: limiting sun exposure, wearing sun protective clothing and using sunscreens may reduce the risk of skin aging, skin cancer and other harmful effects of the sun.
- Avoid exposure to extreme heat or cold.

Active Ingredients

Octinoxate.....7.5% Sunscreen

Homosalate....6.0% Sunscreen

Avobenzone...3.0% Sunscreen

Octocrylene....1.5% Sunscreen

Directions

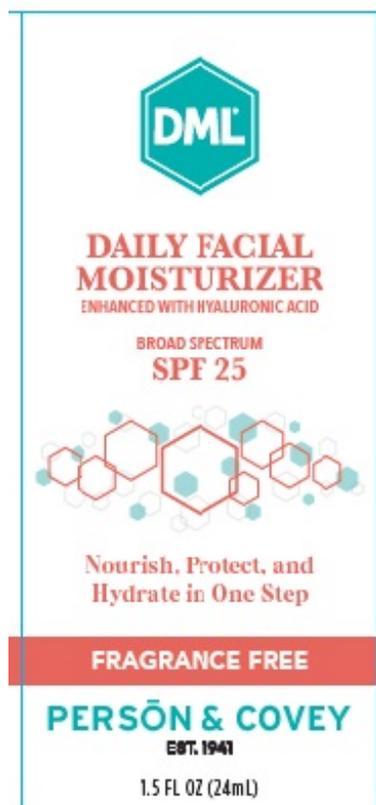
-Apply DML Daily Facial Moisturizer liberally and evenly 15-20 minutes before sun exposure and message in gently, allowing it to dry. Reapply as needed or as directed by your Dermatologist.

-DML Daily Facial is an excellant make-up base.

Purpose

Sunscreen

Package Label Principal Display Panel



DAILY FACIAL MOISTURIZER SPF 25

sunscreen cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:0096-0721

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	0.06 g in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.03 g in 1 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	0.075 g in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	0.015 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
VINYLPYRROLIDONE/EICOSENE COPOLYMER (UNII: 035MV9S1C3)	
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	
CETYL PHOSPHATE (UNII: VT07D6X67O)	
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	
ALLANTOIN (UNII: 344S277G0Z)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
DIMETHICONE 1000 (UNII: MCU2324216)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
GLYCERYL STEARATE/PEG-100 STEARATE (UNII: RD25J5V947)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
PROPYLENE GLYCOL DIETHYLHEXANOATE (UNII: 8D8I9Z0F1Z)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
MAGNESIUM ALUMINUM SILICATE TYPE IA (UNII: SUS08ZOA9S)	
DIMETHICONE 200 (UNII: RGS4T2AS00)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0096-0721-07	7 g in 1 BOTTLE; Type 0: Not a Combination Product	08/01/1991	01/31/2025
2	NDC:0096-0721-45	45 g in 1 BOTTLE; Type 0: Not a Combination Product	08/01/1991	
3	NDC:0096-0721-10	10 g in 1 TUBE; Type 0: Not a Combination Product	01/23/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	08/01/1991	

Labeler - Person and Covey (008482473)

Establishment

Name	Address	ID/FEI	Business Operations
Person and Covey		008482473	manufacture(0096-0721)

Revised: 1/2026

Person and Covey