DAILY FACIAL MOISTURIZER SPF 25- sunscreen cream
Person and Covey

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.

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

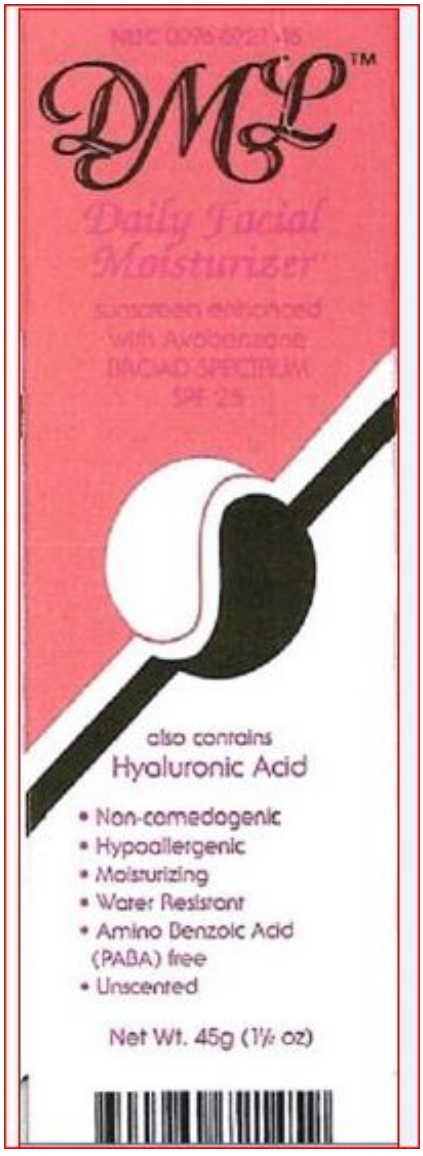
-Apply DML Daily Facial Moisturizer liberally and evenly 15-20 minutes before sun exposure and message in gently, allowing it to dry. Reapply as needed or as directed by your Dermatologist.

-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
DIMETHICONE 1000 (UNII: MCU2324216)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

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	
2	NDC:0096-0721-45	45 g in 1 BOTTLE; Type 0: Not a Combination Product	08/01/1991	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	08/01/1991	

Labeler - Person and Covey (008482473)

Establishment

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

Revised: 11/2018 Person and Covey