DAILY FACIAL MOISTURIZER SPF 35- octinoxate and avobenzone lotion Pharmco Laboratories Inc.

Daily Facial Moisturizer SPF 35

Drug Facts

Active Ingredient:	Purpose
Avobenzone 2.5%	Sunscreen
Octinoxate 4%	Sunscreen

Uses

• Helps prevent sunburn.

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 or irritation develops and lasts
- Keep out of reach of children. If product is swallowed, get medical help or contact Poison Control Center right away.

Directions

- After cleansing apply evenly to face and neck every morning, at least 15 minutes before sun exposure.
- Use a water resistant sunscreen if swimming or sweating.
- Reapply at least every 2 hours or after sweating.
- Children under 6 months of age: Ask a doctor. (Not intended for use on children)

Other Ingredients

Water, Carthamus Tinctorius (Safflower) Oleosomes, C12-15 Alkyl Benzoate, C13-14 Isoparrafin, Laureth 7, Polyacrylamide, Benzyl Alcohol, Fragrance, Sodium Hydroxide, Methylisothiazolinone, Tocopheryl Acetate, Aloe Barbadensis Leaf Extract, Citric Acid

Skin Cancer/Skin Aging Alert & 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

This product has been shown to help prevent sun burn, not **skin cancer or early skin aging.**

Other Information

Store at 15-30°C (59-86°F). Protect the product in this container from excessive heat and or freezing conditions.

Manufactured by:

Pharmco Laboratories Inc. • Titusville, FL 32780 www.pharmcolabs.com • 1.800.635.0712 • Reorder CPL16-3

PRINCIPAL DISPLAY PANEL - 87 g Tube Label

PHARMCO

SKINCARE LABS

Daily Facial Moisturizer

Broad Spectrum SPF 30+

Net wt. 3 oz. (87 g)



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 When using this product, keep out of eyes. Rinse with water to remove

Stop use and ask a doctor if

- rash or irritation develops and lasts
- Keep out of reach of children. If product is swallowed. get medical help or contact Poison Control Center right away.

Directions

- After cleansing apply liberally to face and neck. at least 15 minutes before sun exposure.
- Use a water resistant sunscreen if swimming or sweating.
- Reapply at least every 2 hours or after sweating.
- Children under 6 months of age: Ask a doctor. (Not intended for use on children)

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PLOTC-REV0001

DAILY FACIAL MOISTURIZER SPF 35

octinoxate and avobenzone lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58400-007

Route	- E A			
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TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Octinoxate (UNII: 4Y5P7MUD51) (Octinoxate - UNII:4Y5P7MUD51)	Octinoxate	40 mg in 1 g	
Avobenzone (UNII: G63QQF2NOX) (Avobenzone - UNII:G63QQF2NOX)	Avobenzone	25 mg in 1 g	

Inactive Ingredients			
Ingredient Name	Strength		
Aloe (UNII: V5VD430YW9)			
Benzyl Alcohol (UNII: LKG8494WBH)			
Carthamus Tinctorius Seed Oleosomes (UNII: 9S60Q72309)	Carthamus Tinctorius Seed Oleosomes (UNII: 9S60Q72309)		
Alkyl (C12-15) Benzoate (UNII: A9EJ3J61HQ)			
C13-14 Isoparaffin (UNII: E4F12ROE70)			
Laureth-7 (UNII: Z95S6G8201)			
Methylisothiazolinone (UNII: 229D0E1QFA)			
Sodium Hydroxide (UNII: 55X04QC32I)			
.AlphaTocopherol Acetate (UNII: 9E8X80D2L0)			
Water (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58400- 007-01	87 g in 1 TUBE; Type 0: Not a Combination Product	06/01/2012	
2	NDC:58400- 007-02	3900 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/01/2012	

Labeler - Pharmco Laboratories Inc. (096270814)

Establishment			
Name	Address	ID/FEI	Business Operations
Pharmco Laboratories Inc.		096270814	MANUFACTURE(58400-007), LABEL(58400-007), PACK(58400-007), ANALYSIS (58400-007)

Revised: 1/2024 Pharmco Laboratories Inc.