

**CETAPHIL DAILY FACIAL MOISTURIZER WITH SUNSCREEN SPF 50- octinoxate, octisalate, octocrylene, oxybenzone, titanium dioxide lotion  
Galderma Laboratories, L.P.**

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**Cetaphil Daily Facial Moisturizer with sunscreen SPF 50+**

**Active Ingredients**

Octinoxate 7.5%  
Octisalate 5%  
Octocrylene 7%  
Oxybenzone 6%  
Titanium Dioxide 5.7%

**Purpose**

Sunscreen

**Uses**

Helps prevent sunburn.

**Warnings**

**Skin Cancer/Skin Aging Alert:**

Spending time in the sun increases your risk of skin cancer and early aging. This product has been shown to help prevent sunburn, **not** skin cancer or early aging.

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.

**Directions**

Apply liberally 15 minutes before sun exposure. • Use a water resistant sunscreen if swimming or sweating. • Reapply at least every 2 hours. • Children under 6 months: ask a doctor.

## **Inactive Ingredients**

Water, Propylene Glycol, Glycerin, Dimethicone, VP/Eicosene Copolymer, Cyclomethicone, Stearic Acid, Dimethiconol, Potassium Cetyl Phosphate, Glyceryl Stearate, PEG-100 Stearate, Aluminum Hydroxide, Disodium EDTA, Tocopherol, Triethanolamine, Phenoxyethanol, Ethylparaben, Chlorphenesin, Cetyl Alcohol, Carbomer, Methylparaben, Xanthan Gum

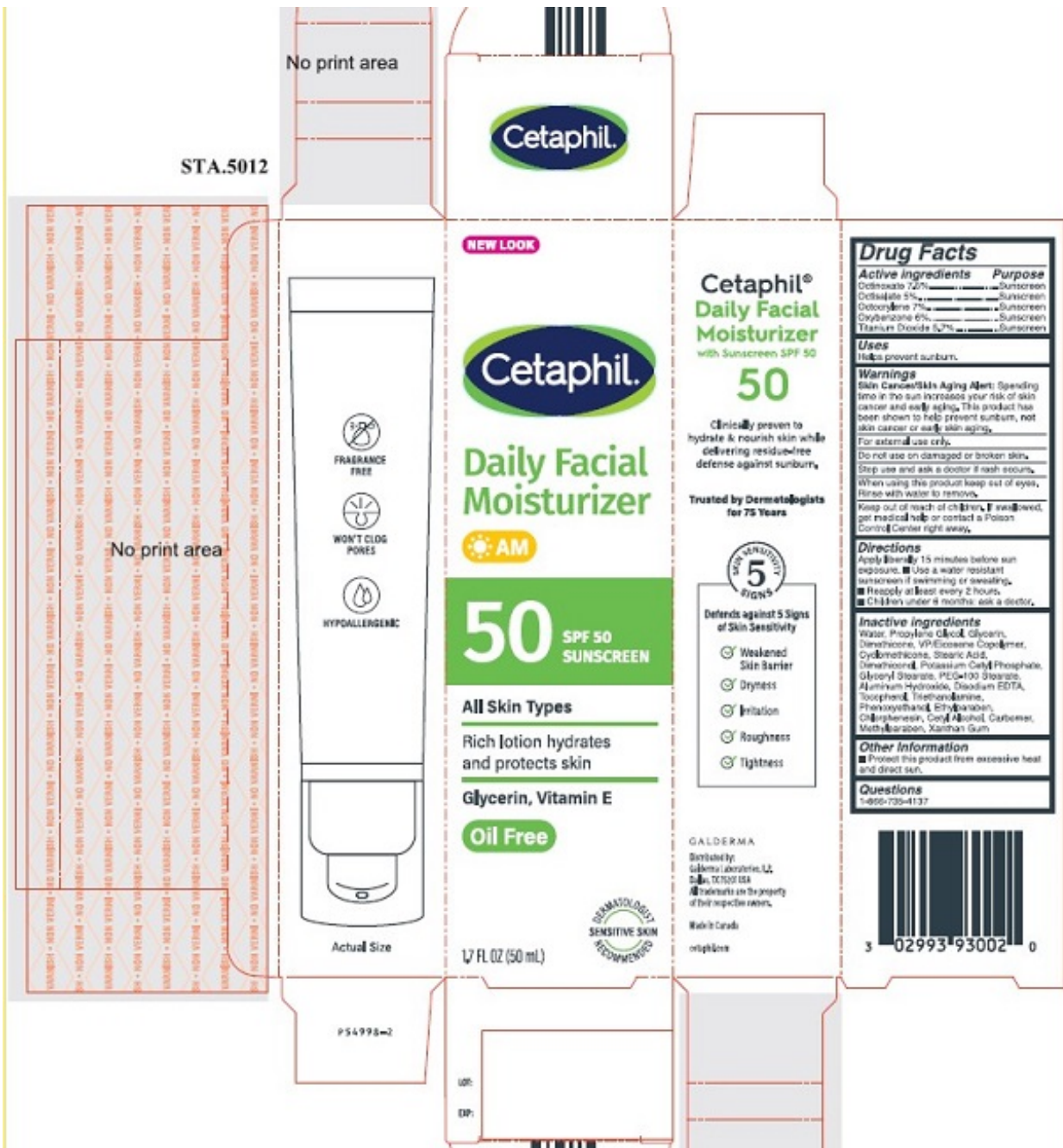
## **Other Information**

- Protect this product from excessive heat and direct sun.

## **Questions?**

1-866-735-4137

**Principal Display Panel - 1.7oz carton**



NEW LOOK

**Cetaphil®**

**Daily Facial  
Moisturizer  
AM**

50 SPF 50  
Sunscreen

All Skin Types

Rich lotion hydrates  
and protects skin

## Glycerin, Vitamin E

Oil Free

Dermatologist Recommended  
Sensitive Skin

1.7 FL OZ (50 mL)

Distributed by:

Galderma Laboratories, L.P.

Dallas, TX 75201 USA

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Made in Canada

cetaphil.com

P54998-2

### CETAPHIL DAILY FACIAL MOISTURIZER WITH SUNCREEN SPF 50

octinoxate, octisalate, octocrylene, oxybenzone, titanium dioxide lotion

#### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0299-4930
<b>Route of Administration</b>	TOPICAL		

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OCTINOXATE</b> (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	70 mg in 1 mL
<b>OXYBENZONE</b> (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:950OS7VE0Y)	OXYBENZONE	60 mg in 1 mL
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	57 mg in 1 mL

#### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>VINYLPYRROLIDONE/EICOSENE COPOLYMER</b> (UNII: 035MV9S1C3)	
<b>CYCLOMETHICONE</b> (UNII: NMQ347994Z)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>DIMETHICONOL (40 CST)</b> (UNII: 343C7U75XW)	
<b>POTASSIUM CETYL PHOSPHATE</b> (UNII: 03KCY6P7UT)	

<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)
<b>ALUMINUM HYDROXIDE</b> (UNII: 5QB0T2IUN0)
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)
<b>TOCOPHEROL</b> (UNII: R0ZB2556P8)
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)
<b>ETHYLPARABEN</b> (UNII: 14255EXE39)
<b>CHLORPHENESIN</b> (UNII: I670DAL4SZ)
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)
<b>CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)</b> (UNII: 809Y72KV36)
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0299-4930-02	1 in 1 CARTON	01/01/2012	
1		50 mL in 1 TUBE; Type 0: Not a Combination Product		

### Marketing Information

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

**Labeler** - Galderma Laboratories, L.P. (047350186)

### Establishment

Name	Address	ID/FEI	Business Operations
G Production Inc.		251676961	manufacture(0299-4930)

Revised: 11/2023

Galderma Laboratories, L.P.