

**DERMA FADE- octinoxate, octisalate, oxybenzone cream**  
**PHARMAGEL INTERNATIONAL INC**

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**PHARMAGEL - DERMA FADE (67879-301)**

**ACTIVE INGREDIENTS**

Octinoxate 7.5%

Octisalate 5.0%

Oxybenzone 3.0%

**PURPOSE**

Sunscreen

**USES**

Lightens the appearance of age spots & skin discoloration. Provides moderate protections against sunburn.

**WARNINGS**

Warnings using this product – For external use only. Avoid contact with eyes. If product gets into the eyes, rinse thoroughly with water. Do not use on children under 12 years of age unless directed by a doctor. Stop use & consult a doctor if rash or irritation develops and persists.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

**DIRECTIONS**

Apply to age spots and skin discoloration. Can be used as a full face moisturizing treatment and sunscreen before sun exposure.

**INACTIVE INGREDIENTS:**

ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER, ALLANTOIN, BIOTIN, BUTYLENE GLYCOL, CAMELLIA SINENSIS (GREEN TEA) LEAF EXTRACT, CAPRYLYL GLYCOL, CETEARYL ALCOHOL, CITRIC ACID, CITRUS UNSHIU PEEL EXTRACT, CYPERUS ROTUNDUS ROOT EXTRACT, DIMETHICONE, DISODIUM EDTA, ETHYLHEXYL PALMITATE, ETHYLHEXYLGLYCERIN, FRAGRANCE (PARFUM), GLYCERIN, GLYCERYL STEARATE, GLYCOLIC ACID, GLYCYRRHIZA GLABRA (LICORICE) ROOT EXTRACT, HEXYLENE GLYCOL, HYDROXYETHYLCELLULOSE, KOJIC ACID, LACTIC ACID, MAGNESIUM ASCORBYL PHOSPHATE, MORUS ALBA LEAF EXTRACT, MYRISTYL LAURATE, MYRISTYL



Derma Fade® contains powerful antioxidants known for their skin rejuvenating and fading properties. Derma Fade is a superior moisturizing crème with powerful skin lighteners and SPF 35 that fades age spots and skin discoloration for a more even, flawless skin tone. Paraben Free.

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www.pharmagel.net Made in USA DER-1 Rev 2

**DERMA FADE®**

**LIGHTENING AND FADING CRÈME**

Evens Skin Tone, Rejuvenates and Moisturizes  
Crème éclaircissante et estompante Crema aclaradora

**ALL SKIN TYPES | WITH SPF 35**

56 g e 2 oz.

# DERMA FADE

octinoxate, octisalate, oxybenzone cream

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67879-301
<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	7.5 g in 100 g
<b>OCTISALATE</b> (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
<b>OXYBENZONE</b> (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	3 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)</b> (UNII: 59TL3WG5CO)	
<b>ALLANTOIN</b> (UNII: 344S277G0Z)	
<b>BIOTIN</b> (UNII: 6SO6U10H04)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>GREEN TEA LEAF</b> (UNII: W2ZU1RY8B0)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>TANGERINE PEEL</b> (UNII: JU3D414057)	
<b>CYPERUS ROTUNDUS TUBER</b> (UNII: 4B51SRR959)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>ETHYLHEXYL PALMITATE</b> (UNII: 2865993309)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>GLYCOLIC ACID</b> (UNII: 0WT12SX38S)	
<b>GLYCYRRHIZA GLABRA</b> (UNII: 2788Z9758H)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>HYDROXYETHYL CELLULOSE (3000 MPA.S AT 1%)</b> (UNII: 7Q6P4JN1QT)	
<b>KOJIC ACID</b> (UNII: 6K23F1TT52)	
<b>LACTIC ACID</b> (UNII: 33X04XA5AT)	
<b>MAGNESIUM ASCORBYL PHOSPHATE</b> (UNII: 0R822556M5)	
<b>MORUS ALBA LEAF</b> (UNII: M8YIA49Q2P)	
<b>MYRISTYL LAURATE</b> (UNII: 58U0NZN2BT)	
<b>MYRISTYL MYRISTATE</b> (UNII: 4042ZC00DY)	
<b>NIACIN</b> (UNII: 2679MF687A)	
<b>PANTHENOL</b> (UNII: WW9CM0O67Z)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>PHYLLANTHUS EMBLICA FRUIT</b> (UNII: YLX4CW2576)	

<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>UNDECYLENIC ACID</b> (UNII: K3D86KJ24N)	
<b>UNDECYLENOYL GLYCINE</b> (UNII: 4D20464K2J)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67879-301-51	1 in 1 BOX	06/18/2015	
1	NDC:67879-301-11	56 g in 1 JAR; 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	06/18/2015	

**Labeler** - PHARMAGEL INTERNATIONAL INC (603215182)

Revised: 1/2024

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