

**DR. C. TUNA RESURFACE ESSENTIAL DAY CREAM SPF 30- homosalate,  
octinoxate, octocrylene, avobenzone  
Farmasi US LLC**

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**Dr. C. Tuna Resurface Essential Day Cream SPF 30**

**Drug Facts**

**Active Ingredient:**

Homosalate 7%

Ethylhexyl Methoxycinnamate 5%

Octocrylene 5%

Butyl Methoxydibenzoylmethane 2%

**Purpose**

Sunscreen

**Use:**

Helps to moisturize and protect the skin from sun damage.

**Warnings:**

For external use only. Do not use on damaged or broken skin.

**Keep out of reach of children.**

**When using this product**

keep out of eyes. Rinse with water to remove

**Stop use and ask a doctor**

if skin irritation develops when using this product. If swallowed, get medical help or contact a Poison Control Center right away.

**Directions:**

Use daily, every morning 15 minutes before sun exposure. Take enough amounts with the enclosed spatula on your hands; gently apply on clean face and neck.

**INACTIVE INGREDIENTS:**

Water/Aqua, Cetyl Alcohol, Dicaprylyl Carbonate, Caprylic/Capric Triglyceride, Cichorium Intybus Leaf Extract, Hexylene Glycol, Caprylyl Glycol, Glyceryl Stearate SE, Phenoxyethanol, Cetyl Palmitate, Hexyldecanol, Hexyldecyl Laurate, Sodium Stearoyl Glutamate, Glycerin, Sodium Polyacrylate, Helianthus Annuus Seed Oil, Chlorella Vulgaris Extract, Rosmarinus Officinalis Leaf Extract, Niacinamide, Polymethylsilsesquioxane, Disodium EDTA, Xanthan Gum, Ethylhexylglycerin.

## Questions or Comments?

info@farmasius.com (833) 432-7627 Monday - Friday (9 a.m - 9 p.m. EST)

## Package Labeling:



## DR. C. TUNA RESURFACE ESSENTIAL DAY CREAM SPF 30

homosalate, octinoxate, octocrylene, avobenzone kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:74690-025
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74690-025-00	1 in 1 KIT	08/15/2021	

### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 PACKET	2.5 mL

### Part 1 of 1

## DR. C. TUNA LUMI RADIANCE BRIGHTENING CREAM 22 SPF

homosalate, octinoxate, octocrylene, avobenzone cream

### Product Information

<b>Item Code (Source)</b>	NDC:74690-026
<b>Route of Administration</b>	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	70 mg in 1 mL
<b>OCTINOXATE</b> (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	50 mg in 1 mL
<b>OCTOCRYLENE</b> (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	50 mg in 1 mL
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	20 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0K00R)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>DICAPRYLYL CARBONATE</b> (UNII: 609A3V1SUA)	
<b>MEDIUM-CHAIN TRIGLYCERIDES</b> (UNII: C9H2L21V7U)	
<b>CHICORY LEAF</b> (UNII: WBQ249COFR)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	
<b>GLYCERYL STEARATE SE</b> (UNII: FCZ5MH785I)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>CETYL PALMITATE</b> (UNII: 5ZA2S6B08X)	
<b>HEXYLDECANOL</b> (UNII: 151Z7P1317)	
<b>HEXYLDECYL LAURATE</b> (UNII: 0V595C1P6M)	
<b>SODIUM STEAROYL GLUTAMATE</b> (UNII: 65A9F4P024)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>SUNFLOWER OIL</b> (UNII: 3W1JG795YI)	
<b>CHLORELLA VULGARIS</b> (UNII: RYQ4R60M02)	
<b>ROSEMARY</b> (UNII: IJ67X351P9)	
<b>NIACINAMIDE</b> (UNII: 25X51I8RD4)	
<b>POLYMETHYLSILSESQUIOXANE (4.5 MICRONS)</b> (UNII: 59Z907ZB69)	
<b>EDETATE DISODIUM ANHYDROUS</b> (UNII: 8NLQ36F6MM)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74690-026-00	2.5 mL in 1 PACKET; 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	08/15/2021	

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OTC Monograph Drug	M020	08/15/2021	

**Labeler** - Farmasi US LLC (113303351)

Revised: 12/2024

Farmasi US LLC