

**DR. C. TUNA SUNSCREEN SPF30 BROAD SPECTRUM- avobenzone, octinoxate, octocrylene, homosalate lotion**  
**Farmasi US LLC**

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**Dr. C. Tuna Sunscreen SPF30 Broad Spectrum**

***Drug Facts***

***Active Ingredients***

Butyl Methoxydibenzoylmethane 1.5%

Ethylhexyl Methoxycinnamate 5%

Octocrylene 5%

Homosalate 7%

***Purpose***

Sunscreen

***Uses***

Helps prevent sunburn - if used as directed with other sun protection measures (see ), decreases the risk of skin cancer and early skin aging caused by the sun. **Directions**

***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 doctor if**

rash occurs.

**- Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away

***Directions***

**For Sunscreen Use:**

-Apply generously 15 minutes before sun exposure.-Reapply; - after 80 minutes of swimming or sweating - immediately after towel drying - at least every 2 hours

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 - children under 6 months of age: Ask a doctor - **Sun Protection Measures:**

### ***Other Information***

- protect this product from excessive heat and direct sun. - may stain some fabrics

### ***Inactive Ingredients***

Water/Aqua, C12-15 Alkyl Benzoate, Cetearyl Alcohol, Glyceryl Stearate, Peg-100 Stearate, Glycerin, Ethylhexyl Palmitate, Cetearith-20, Cyclopentasiloxane, Phenoxyethanol, Dimethicone, Acrylates, C10-30 Alkyl Acrylate Crosspolymer, Glycine Soja Oil, Daucus Carota Sativa Root Extract, Beta-Carotene, Tocopherol, Triethanolamine, Tocopheryl Acetate, Ethylhexylglycerin.

### ***Questions or Comments?***

info@farmasius.com +1 (833) 432 76 27

Monday - Friday (9 a.m - 6p.m. EST)

### ***Package Labeling***

F



## SUNSCREEN

### Drug Facts

Active Ingredients	PURPOSE
Butyl Methoxydibenzoylmethane 1.5%.....	Sunscreen
Ethylhexyl Methoxycinnamate 5%.....	Sunscreen
Octocrylene 5%.....	Sunscreen
Homosalate 7%.....	Sunscreen

### Uses

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### Warnings

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### Directions

#### For Sunscreen Use:

- Apply generously 15 minutes before sun exposure. - Reapply; - after 80 minutes of swimming or sweating - immediately after towel drying - at least every 2 hours - **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 - children under 6 months of age: Ask a doctor

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### Distributed By:

FARMASI US LLC 2315 NW 107 TH AVE, STE 1B 12,  
DORAL , FL 33172.

4801178

8690131109899



# SUN SCREEN

# 30<sup>SPF</sup>

BROAD SPECTRUM

Face & Body

**UVA + UVB**  
Protection

**CLEAN FEEL**  
**LIGHT WEIGHT**

75 ml e 2,5 fl.oz.

## DR. C. TUNA SUNSCREEN SPF30 BROAD SPECTRUM

avobenzone, octinoxate, octocrylene, homosalate lotion

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:74690-001
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>AVOBENZONE</b> (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	15 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>HOMOSALATE</b> (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	70 mg in 1 mL

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALKYL (C12-15) BENZOATE</b> (UNII: A9EJ3J61HQ)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>ETHYLHEXYL PALMITATE</b> (UNII: 2865993309)	
<b>POLYOXYL 20 CETOSTEARYL ETHER</b> (UNII: YRC528SWUY)	
<b>CYCLOMETHICONE 5</b> (UNII: 0THT5PCI0R)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)</b> (UNII: 59TL3WG5CO)	
<b>ACRYLIC ACID</b> (UNII: J94PBK7X8S)	
<b>SOYBEAN OIL</b> (UNII: 241ATL177A)	
<b>CARROT</b> (UNII: L56Z1JK48B)	
<b>BETA CAROTENE</b> (UNII: 01YAE03M7J)	
<b>TOCOPHEROL</b> (UNII: R0ZB2556P8)	
<b>TROLAMINE</b> (UNII: 9O3K93S3TK)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:74690-001-01	1 in 1 BOX	06/01/2020	
1		75 mL in 1 TUBE; Type 0: Not a Combination Product		

## Marketing Information

<b>Marketing</b>	<b>Application Number or Monograph</b>	<b>Marketing Start</b>	<b>Marketing End</b>
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Category	Citation	Date	Date
OTC Monograph Drug	M020	06/01/2020	

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

Revised: 12/2023

Farmasi US LLC