

**CELL FUSION C LASER SUNSCREEN BROAD SPECTRUM SPF 50- homosalate, octisalate, titanium dioxide, zinc oxide cream
CMS LAB Inc.**

Cell Fusion C Laser Sunscreen Broad Spectrum SPF 50

Drug Facts

ACTIVE INGREDIENTS

Homosalate 7.5%
Octisalate 4.5%
Titanium Dioxide 2.2%
Zinc Oxide 12.7%

PURPOSE

Sunscreen

USES

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

WARNINGS

For external use only

- **Do not use on** damaged or broken skin
- **When using this product** keep out of the eyes. Rinse with water to remove.
- **Stop use and ask a 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 and evenly 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

OTHER INFORMATION

- protect the product in this container from excessive heat and direct sun
- may stain some fabrics or surfaces.

INACTIVE INGREDIENTS

WATER

QUESTIONS?

Call 212-804-7608

Outside U.S. +82 080-889-7760

www.cellfusionnc.com

Dist. by Dkcos

20 W 33rd St New York, NY 10001

www.dkcos.com

CMS LAB Inc.

Made in Korea

PRINCIPAL DISPLAY PANEL

Cell Fusion C

Dermatological Laboratory

LASER

SUNSCREEN

Broad Spectrum SPF 50+

Effective UV Protection

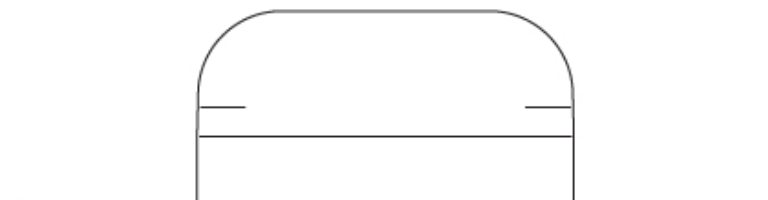
Improve Overall

Skin Condition

WATER RESISTANT

(80 minutes)

50ML / 1.69 FL.OZ.



Drug Facts

Active Ingredients	Purpose
Homosalate (7.5%)	Sunscreen
Octisalate (4.5%)	Sunscreen
Titanium Dioxide (2.2%)	Sunscreen
Zinc Oxide (12.7%)	Sunscreen

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Drug Facts (Continued)

Inactive Ingredients

Water, Cyclopentasiloxane, Dipropylene Glycol, Disiloxane, Lauryl PEG-10 Tris(trimethylsiloxy) Silylethyl Dimethicone, Butyloctyl Salicylate, Glycerin, Silica, Dimethicone, Distearidimonium Hectorite, Magnesium Sulfate, Methyl Methacrylate Crosspolymer, 1,2-Hexanediol, Triethoxycaprylylsilane, Stearic Acid, Aluminum Hydroxide, Sorbitan Caprylate, Dimethicone/PEG-10/15 Crosspolymer, Lavandula Angustifolia (Lavender) Oil, Dimethicone/Vinyl Dimethicone Crosspolymer, Glyceryl Caprylate, Ethylhexylglycerin, Sodium Citrate, Hydrolyzed Collagen, Hydroxydecyl Ubiquinone, Tocopherol, Hydrogenated Lecithin, Ceramide NP, Sucrose Stearate, Cholesterol, Palmitic Acid, Cholesteryl Macadamiate, Biosaccharide Gum-4, Hydrolyzed Lupine Protein, Saccharide Isomerate



Cell Fusion C
Dermatological Laboratory

**LASER
SUNSCREEN**
Broad Spectrum SPF 50+

*Effective UV Protection
Improves Overall
Skin Condition*

WATER RESISTANT
(80 minutes)
50ML / 1.69 FL.OZ.



10ML / 0.33 FL.OZ.

Cell Fusion C
Dermatological Laboratory

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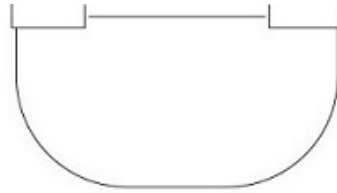
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Improves Overall
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(80 minutes)

10ML / 0.33 FL.OZ.



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CELL FUSION C LASER SUNSCREEN BROAD SPECTRUM SPF 50

homosalate, octisalate, titanium dioxide, zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52554-9051
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7.5 g in 50 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	12.7 g in 50 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	4.5 g in 50 mL
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	2.2 g in 50 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52554-9051-1	1 in 1 BOX	05/15/2024	
1		50 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:52554-9051-2	1 in 1 BOX	05/15/2024	
2		10 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	05/15/2024	

Labeler - CMS LAB Inc. (557795012)

