CALLIEL TINTED GLOW SPF 50 MEDIUM- sunscreen cream cream
CALLIEL TINTED GLOW SPF 50 LIGHT- sunscreen cream cream
CALLIEL TINTED GLOW SPF 50- sunscreen cream cream
CALLIEL PHARMA KOZMETIK SANAYI VE TICARET ANONIM SIRKETI

Active ingredients

Zinc Oxide 7%
Titanium Dioxide 3.5%
Avobenzone 3.5%
Octinoxate 7%

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

Do not use on damaged or broken skin

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

• apply liberally 15 minutes before sun exposure
• reapply:
• after 80 minutes of swimming or sweating
• immediately after towel drying
• at least every 2 hours
• children under 6 months: Ask a doctor

Sun Protection Measures: Spending time in the sun increases your risk of skin cancer and early skin aging caused by the sun. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF 15 or higher and other sun protection measures including:
• limit time in the sun, especially from 10 a.m. – 2 p.m.
• wear long-sleeve shirts, pants, hats, and sunglasses

Warnings

For external use only

Purpose

Sunscreen

Inactive ingredients

Water, Caprylic/Capric Triglyceride, Butyrospermum Parkii Butter, C12-15 Alkyl Benzoate, Dimethicone, Glycerin, Propylene Glycol, Isopropyl Myristate, Cyclopentasiloxane, Acrylates/Dimethicone Copolymer, Xylitylglucoside, Phenoxyethanol, PEG-10 Dimethicone, Iron Oxides, Disteardimonium Hectorite, Anhydroxylitol, Xylitol, Aloe Barbadensis Leaf Extract, Caprylyl Glycol, Triethoxycaprylylsilane

Front Panel


CALLIEL
TINTED
GLOW
+50 SPF UVA + UVB
Ultimate Sun Protection
Deeply Moisturizes
CLINICAL SOLUTION
Dermatologist Tested

CALLIEL

CALLIEL TINTED GLOW SPF 50 MEDIUM

sunscreen cream cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87411-0002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	3.5 g in 100 g

AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3.5 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	7 g in 100 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	1 g in 100 g
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	6 g in 100 g
CI 77492 (UNII: EX438O2MRT)	0.6 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	3 g in 100 g
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	2 g in 100 g
CI 77499 (UNII: XM0M87F357)	0.3 g in 100 g
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	2 g in 100 g
BIS-PEG-10 DIMETHICONE/DIMER DILINOLEATE COPOLYMER (UNII: CF5W1YCX11)	0.8 g in 100 g
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	0.5 g in 100 g
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	0.3 g in 100 g
BUTYROSPERMUM PARKII (SHEA) BUTTER (UNII: K49155WL9Y)	4 g in 100 g
DIMETHICONE (UNII: 92RU3N3Y1O)	3 g in 100 g
C12-15 ALKYL BENZOATE (UNII: A9EJ3J61HQ)	5 g in 100 g
XYLITOL (UNII: VCQ006KQ1E)	0.3 g in 100 g
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	0.4 g in 100 g
CI 77491 (UNII: 1K09F3G675)	0.4 g in 100 g
WATER (UNII: 059QF0KO0R)	45.18 g in 100 g
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.8 g in 100 g
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	0.1 g in 100 g
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	0.2 g in 100 g
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	0.12 g in 100 g
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	3 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:87411-0002-1	32 g in 1 TUBE; Type 0: Not a Combination Product	02/07/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/07/2026	

CALLIEL TINTED GLOW SPF 50 LIGHT

sunscreen cream cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87411-0003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	3.5 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3.5 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	7 g in 100 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	1 g in 100 g
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	6 g in 100 g
CI 77492 (UNII: EX438O2MRT)	0.6 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	3 g in 100 g
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	2 g in 100 g
CI 77499 (UNII: XM0M87F357)	0.3 g in 100 g
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	2 g in 100 g
BIS-PEG-10 DIMETHICONE/DIMER DILINOLEATE COPOLYMER (UNII: CF5W1YCX11)	0.8 g in 100 g
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	0.5 g in 100 g
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	0.3 g in 100 g
BUTYROSPERMUM PARKII (SHEA) BUTTER (UNII: K49155WL9Y)	4 g in 100 g

DIMETHICONE (UNII: 92RU3N3Y1O)	3 g in 100 g
C12-15 ALKYL BENZOATE (UNII: A9EJ3J61HQ)	5 g in 100 g
XYLITOL (UNII: VCQ006KQ1E)	0.3 g in 100 g
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	0.4 g in 100 g
CI 77491 (UNII: 1K09F3G675)	0.4 g in 100 g
WATER (UNII: 059QF0KO0R)	45.18 g in 100 g
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.8 g in 100 g
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	0.1 g in 100 g
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	0.2 g in 100 g
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	0.12 g in 100 g
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	3 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87411-0003-1	32 g in 1 TUBE; Type 0: Not a Combination Product	02/07/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/07/2026	

CALLIEL TINTED GLOW SPF 50

sunscreen cream cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87411-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	3.5 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3.5 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	7 g in 100 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
2-ETHYLHEXYL ACRYLATE, METHACRYLATE, METHYL METHACRYLATE, OR BUTYL METHACRYLATE/HYDROXYPROPYL DIMETHICONE COPOLYMER (30000-300000 MW) (UNII: S7ZA3CCJ4M)	1 g in 100 g
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	6 g in 100 g
CI 77492 (UNII: EX438O2MRT)	0.6 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	3 g in 100 g
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	2 g in 100 g
CI 77499 (UNII: XM0M87F357)	0.3 g in 100 g
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	2 g in 100 g
BIS-PEG-10 DIMETHICONE/DIMER DILINOLEATE COPOLYMER (UNII: CF5W1YCX11)	0.8 g in 100 g
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	0.5 g in 100 g
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	0.3 g in 100 g
BUTYROSPERMUM PARKII (SHEA) BUTTER (UNII: K49155WL9Y)	4 g in 100 g
DIMETHICONE (UNII: 92RU3N3Y1O)	3 g in 100 g
C12-15 ALKYL BENZOATE (UNII: A9EJ3J61HQ)	5 g in 100 g
XYLITOL (UNII: VCQ006KQ1E)	0.3 g in 100 g
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	0.4 g in 100 g
CI 77491 (UNII: 1K09F3G675)	0.4 g in 100 g
WATER (UNII: 059QF0KO0R)	44.8 g in 100 g
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.8 g in 100 g
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	0.1 g in 100 g
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	0.2 g in 100 g
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	0.5 g in 100 g
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	3 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87411-0001-1	50 g in 1 TUBE; Type 0: Not a Combination Product	02/15/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/15/2026	

Labeler - CALLIEL PHARMA KOZMETIK SANAYI VE TICARET ANONIM SIRKETI (751146029)

Registrant - MADDY GLOBAL LLC (144837954)

Establishment

Name	Address	ID/FEI	Business Operations
CALLIEL PHARMA KOZMETIK SANAYI VE TICARET ANONIM SIRKETI		751146029	manufacture(87411-0001, 87411-0002, 87411-0003)

Revised: 2/2026 CALLIEL PHARMA KOZMETIK SANAYI VE TICARET ANONIM SIRKETI