ULOOK SUNSCREEN SPF 50- sunscreen cream
HAMEX GLOBAL COSMETİCS LLC

Active ingredients

Active ingredients ..... Purpose

Homosalate 6.44% ..... Sunscreen

Octinoxate 17.28% ..... Sunscreen

Titanium Dioxide 2% ..... Sunscreen

Octisalate 3.08% ..... Sunscreen

Avobenzone 2.8% ..... Sunscreen

Octocrylene 8.4% ..... 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 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

apply liberally 15 minutes before sun exposure

reapply at least every 2 hours

use a water-resistant sunscreen if swimming or sweating

Inactive ingredients

Aqua, Glycerin, 1,2-Hexanediol, Caprylic/Capric Triglyceride, Butylene Glycol, Aloe Barbadensis Leaf Extract, Allantoin, Centella Asiatica Extract, Ethylhexylglycerin, Panthenol, Cetyl Palmitate, Ceteareth-12, Citric Acid, Saccharide Isomerate, Simmondsia

Chinensis Seed Oil, Dehydroxanthan Gum, Magnesium Disodium EDTA, Glyceryl Stearate, Lactic Acid, Ceteareth-20, Sodium Hyaluronate, Pentaerythrityl Distearate, Tocopheryl Acetate, Micrococcus Luteus, Phenoxyethanol, Sodium PCA, Chamomilla Recutita Flower Extract, Cetearyl Alcohol, Melaleuca Alternifolia Leaf Oil, Sodium Citrate.

Keep out of reach of children

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

Purpose

Sunscreen

PRINCIPAL DISPLAY PANEL

Clean Beauty Collection uLOOK Glowy SPF 50+ Sunscreen Cream Broad Spectrum Protection (UVA+UVB) Net Wt: 1.69 fl.oz. (50ml)

ULOOK SUNSCREEN SPF 50

sunscreen cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87552-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	64.4 mg in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	84 mg in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	28 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	30.8 mg in 1 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	20 mg in 1 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	72.8 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	2 mg in 1 g
AQUA (UNII: 059QF0KO0R)	437.5 mg in 1 g
GLYCERIN (UNII: PDC6A3C0OX)	0.1 mg in 1 g
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	50 mg in 1 g
1,3-BUTANEDIOL CYCLIC SULFITE, CIS- (UNII: 1EC8ULI6OQ)	0.1 mg in 1 g
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.1 mg in 1 g
ALLANTOIN (UNII: 344S277G0Z)	1 mg in 1 g
CENTELLA ASIATICA (UNII: 7M867G6T1U)	0.1 mg in 1 g
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	2 mg in 1 g
PANTHENOL (UNII: WV9CM0O67Z)	5 mg in 1 g
CETYL PALMITATE (UNII: 5ZA2S6B08X)	20 mg in 1 g
CETEARETH-12 (UNII: 7V4MR24V5P)	20 mg in 1 g
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	2 mg in 1 g
SACCHARIDE ISOMERATE (UNII: W8K377W98I)	5 mg in 1 g
JOJOBA OIL (UNII: 724GKU717M)	10 mg in 1 g
DEHYDROXANTHAN GUM (UNII: 63ZP7I1BQO)	2 mg in 1 g
MAGNESIUM DISODIUM EDTA (UNII: NDT563S5VZ)	0.5 mg in 1 g
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	40 mg in 1 g
LACTIC ACID (UNII: 33X04XA5AT)	2 mg in 1 g
CETEARETH-20 (UNII: YRC528SWJY)	0.1 mg in 1 g
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	0.1 mg in 1 g
PENTAERYTHRITYL DISTEARATE (UNII: 697WOT8HNB)	0.1 mg in 1 g
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	0.1 mg in 1 g
MICROCOCCUS LUTEUS (UNII: LV6L29Z6AX)	0.1 mg in 1 g
PHENOXYETHANOL (UNII: HIE492ZZ3T)	8 mg in 1 g
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	5 mg in 1 g
MATRICARIA CHAMOMILLA WHOLE (UNII: G0R4UBI2ZZ)	0.1 mg in 1 g
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	50 mg in 1 g
TEA TREE OIL (UNII: VIF565UC2G)	2 mg in 1 g
SODIUM CITRATE (UNII: 1Q73Q2JULR)	2 mg in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87552-001-01	50 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/29/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	03/27/2026	

Labeler - HAMEX GLOBAL COSMETİCS LLC (142915436)

Registrant - HAMEX GLOBAL COSMETİCS LLC (142915436)

Establishment

Name	Address	ID/FEI	Business Operations
HAMAK GLOBAL KOZMETIK DIS TICARET LIMITED SIRKETI		520131567	manufacture(87552-001)

Revised: 3/2026 HAMEX GLOBAL COSMETİCS LLC