UV SOLUTIONS HYDRATING BROAD SPECTRUM SPF 50-avobenzone,homosalate,octisalate, and octocrylene cream
CLINIQUE LABORATORIES LLC

UV SOLUTIONS HYDRATING BROAD SPECTRUM SPF 50

Active ingredietns

Avobenzone 3%

Homosalate 7%

Octisalate 4.5%

Octocrylene 7%

Purpose

Sunscreen

Use

helps prevent sunburn

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

- For sunscreen use:
- apply liberally and evenly 15 minutes before sun exposure
- reapply at least every two hours
- use a water resistant sunscreen if swimming or sweating
- 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

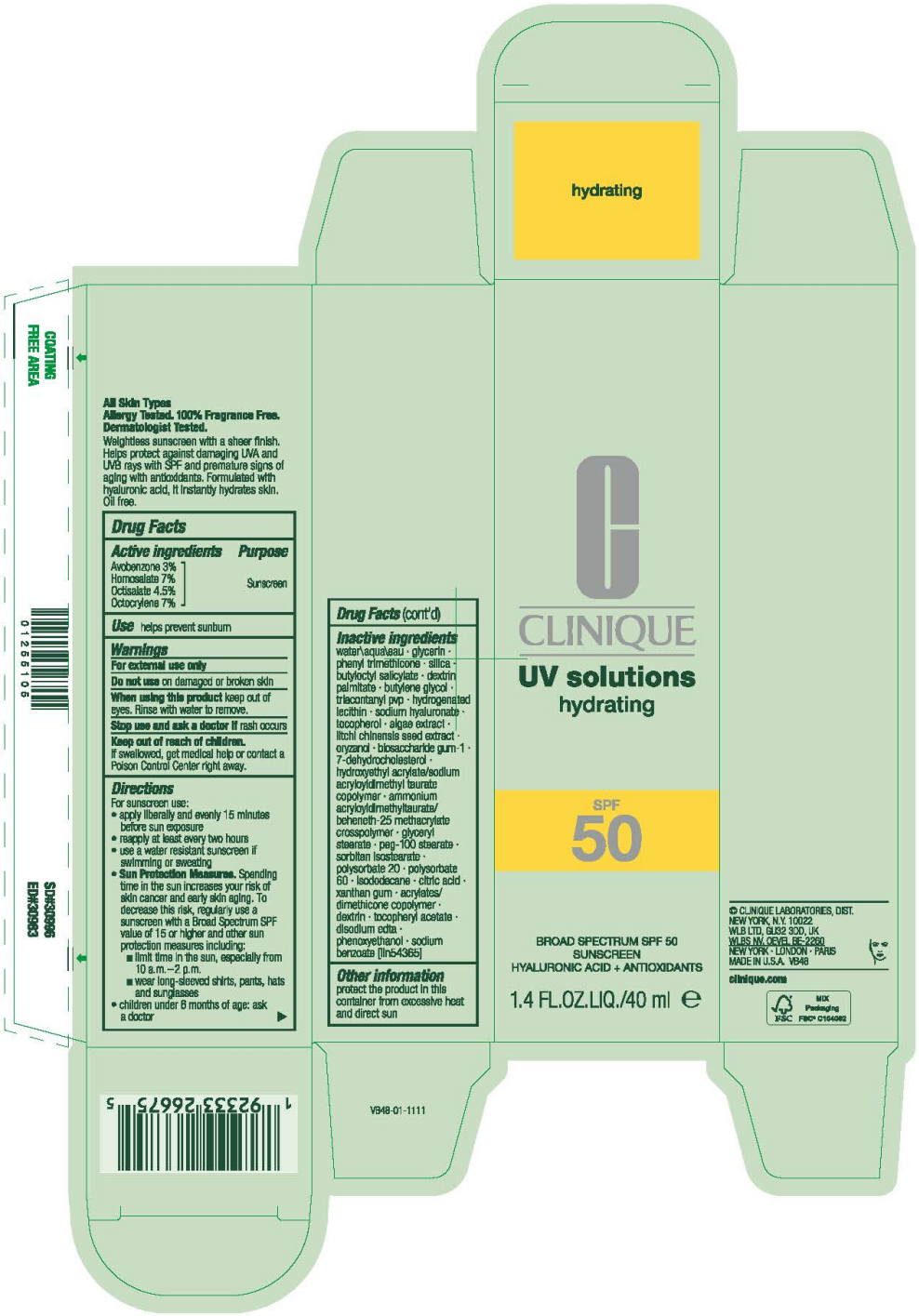
Inactive ingredients

water\aqua\eau • glycerin • phenyl trimethicone • silica • butyloctyl salicylate • dextrin palmitate • butylene glycol • triacontanyl pvp • hydrogenated lecithin • sodium hyaluronate • tocopherol • algae extract • litchi chinensis seed extract • oryzanol • biosaccharide gum-1 • 7-dehydrocholesterol • hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer • ammonium acryloyldimethyltaurate/beheneth-25 methacrylate crosspolymer • glyceryl stearate • peg-100 stearate • sorbitan isostearate • polysorbate 20 • polysorbate 60 • isododecane • citric acid • xanthan gum • acrylates/dimethicone copolymer • dextrin • tocopheryl acetate • disodium edta • phenoxyethanol • sodium benzoate [iln54365]

Other information

protect the product in this container from excessive heat and direct sun

CLINIQUE
UV solutions
hydrating
SPF 50
BROAD SPECTRUM SPF 50
SUNSCREEN
HYALURONIC ACID+ ANTIOXIDANTS
1.4 FL.OZ.LIQ./40 ml e



UV SOLUTIONS HYDRATING BROAD SPECTRUM SPF 50

avobenzone,homosalate,octisalate, and octocrylene cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49527-224
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	45 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	70 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	70 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM BENZOATE (UNII: OJ245FE5EU)	
BIOSACCHARIDE GUM-1 (UNII: BB4PU4V09H)	
7-DEHYDROCHOLESTEROL (UNII: BK1IU07GKF)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
DEXTRIN PALMITATE (CORN; 20000 MW) (UNII: 89B2BSF9I3)	
TOCOPHEROL (UNII: R0ZB2556P8)	
LITCHI CHINENSIS SEED (UNII: 9294024N9Q)	
XANTHAN GUM (UNII: TTV12P4NEE)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
WATER (UNII: 059QF0KO0R)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/BEHENETH-25 METHACRYLATE CROSSPOLYMER (52000 MPA.S) (UNII: LZ291VH90H)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
GLYCERIN (UNII: PDC6A3C0OX)	
TRIACONTANYL PVP (UNII: N0SS3Q238D)	
PORPHYRIDIUM PURPUREUM (UNII: K2P8K2558N)	
ICODEXTRIN (UNII: 2NX48Z0A9G)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ORYZANOL (UNII: SST9XCL51M)	
HYDROGENATED SOYBEAN LECITHIN (UNII: H1109Z9J4N)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SILICA (UNII: ETJ7Z6XBU4)	
ISODODECANE (UNII: A8289P68Y2)	
CITRIC ACID (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49527-224-01	1 in 1 CARTON	02/01/2026	

1		40 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:49527-224-02	1 in 1 CARTON	02/01/2026	
2		5 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	02/01/2026	

Labeler - CLINIQUE LABORATORIES LLC (044475127)

Establishment

Name	Address	ID/FEI	Business Operations
The Estee Lauder Inc		802599436	manufacture(49527-224) , label(49527-224) , pack(49527-224)

Revised: 2/2026 CLINIQUE LABORATORIES LLC