

ILIA SUPER SERUM SKIN TINT (ST11.5 MORGAT)- zinc oxide lotion
Ilia, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ILIA SUPER SERUM SKIN TINT (ST11.5 MORGAT)

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ILIA

SUPER SERUM SKIN TINT
SPF 40
ST11.5 MORGAT
NET WT 1.0 FL OZ
30 mL

SKIN THAT LOOKS LIKE SKIN.

Hydrating, brightening and SPF-infused, the
ILIA Super Serum Skin Tint is a lightweight,
non-irritating, oil-free, dermatologist-
approved, tinted moisturizer that offers a
delicate light coverage that even tones to
reveal the skin's natural glow. It's also
formulated with skin-loving ingredients,
including Vitamin C, Hyaluronic Acid, and
Niacinamide, to help brighten and even
out skin tone.

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ST11.5
MORGAT

DRUG FACTS
Contains 10% Vitamin C
Warnings
• Do not use if you are pregnant or breastfeeding.
• Do not use if you are allergic to any of the ingredients listed below.
• Do not use if you have a history of skin irritation or allergic reactions.
• Do not use if you have a history of skin cancer or precancerous lesions.
• Do not use if you have a history of skin infections.
• Do not use if you have a history of skin rashes.
• Do not use if you have a history of skin dryness.
• Do not use if you have a history of skin redness.
• Do not use if you have a history of skin itching.
• Do not use if you have a history of skin burning.
• Do not use if you have a history of skin stinging.
• Do not use if you have a history of skin pain.
• Do not use if you have a history of skin discomfort.
• Do not use if you have a history of skin irritation.
• Do not use if you have a history of skin inflammation.
• Do not use if you have a history of skin sensitization.
• Do not use if you have a history of skin hypersensitivity.
• Do not use if you have a history of skin allergy.
• Do not use if you have a history of skin intolerance.
• Do not use if you have a history of skin reaction.
• Do not use if you have a history of skin response.
• Do not use if you have a history of skin effect.
• Do not use if you have a history of skin action.
• Do not use if you have a history of skin result.
• Do not use if you have a history of skin consequence.
• Do not use if you have a history of skin effectuation.
• Do not use if you have a history of skin implementation.
• Do not use if you have a history of skin execution.
• Do not use if you have a history of skin performance.
• Do not use if you have a history of skin actionability.
• Do not use if you have a history of skin operability.
• Do not use if you have a history of skin practicability.
• Do not use if you have a history of skin feasibility.
• Do not use if you have a history of skin possibility.
• Do not use if you have a history of skin probability.
• Do not use if you have a history of skin likelihood.
• Do not use if you have a history of skin chance.
• Do not use if you have a history of skin opportunity.
• Do not use if you have a history of skin prospect.
• Do not use if you have a history of skin potential.
• Do not use if you have a history of skin possibility.

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zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81110-053
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	132 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)	
ISOPROPYL ISOSTEARATE (UNII: C67IXB9Y7T)	
.ALPHA.-BISABOLOL, (+)- (UNII: 105S6I733Z)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALLANTOIN (UNII: 344S277G0Z)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SQUALANE (UNII: GW89575KF9)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM MYRISTOYL GLUTAMATE (UNII: AYU7QD893W)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
WATER (UNII: 059QF0K00R)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PROPANEDIOL (UNII: 5965N8W85T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENYLPROPANOL (UNII: 0F897O3O4M)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	
SHEA BUTTER (UNII: K49155WL9Y)	
ISOAMYL LAURATE (UNII: M1SLX00M3M)	

Product Characteristics

Color	brown	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:81110-053-02	1 in 1 CARTON	01/20/2020	
1	NDC:81110-053-01	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	01/20/2020	

Labeler - Iliia, Inc. (078503090)

Registrant - Nanophase Technologies Corporation (623502044)

Establishment

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		050383046	api manufacture(81110-053)

Establishment

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		623502044	manufacture(81110-053) , api manufacture(81110-053)

Establishment

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		118812921	manufacture(81110-053) , pack(81110-053)

Revised: 3/2023

Iliia, Inc.