

**ILIA SUPER SERUM SKIN TINT (ST2 TULUM)- 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.*

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**ILIA SUPER SERUM SKIN TINT (ST2 TULUM)**

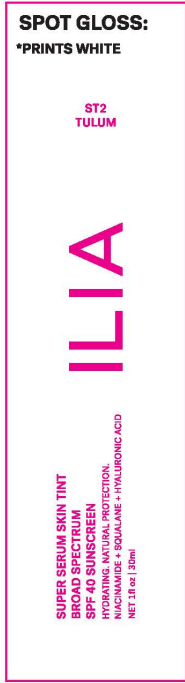
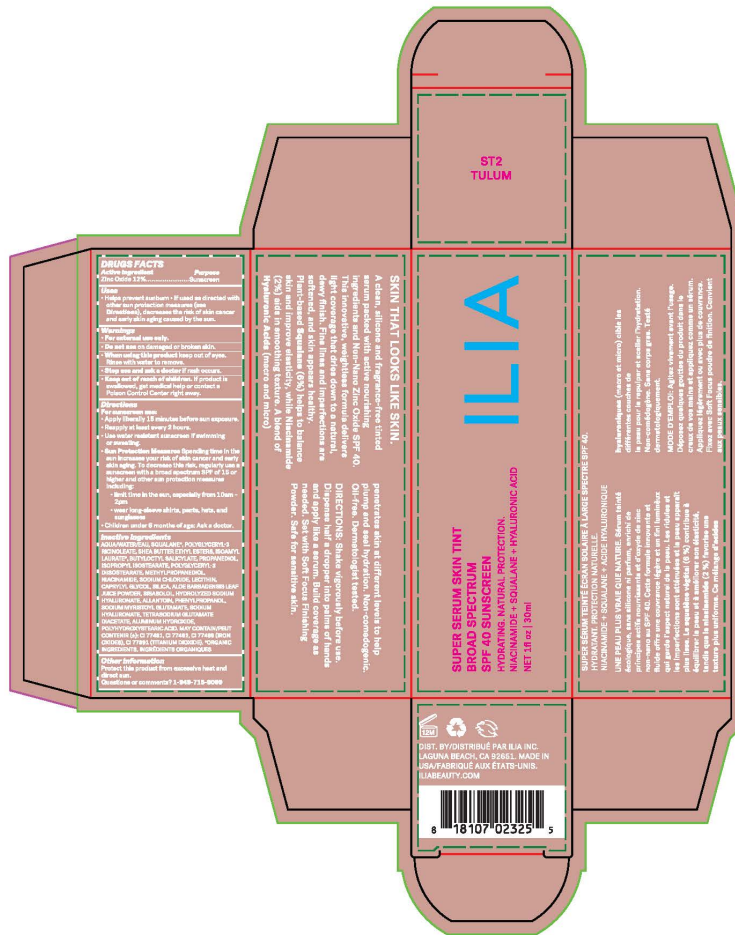
**ILIA ST2**

ILIA - SUPER SERUM SKIN TINT SPF 40 - ST2  
US

(CUSTOM 7613 U)

TEXT: WHITE

UNCOATED SIDE OF BOARD  
ON PRINT SIDE  
041A



**ILIA SUPER SERUM SKIN TINT (ST2 TULUM)**  
zinc oxide lotion

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:81110-751
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
SHEA BUTTER (UNII: K49155WL9Y)	
ISOAMYL LAURATE (UNII: M1SLX00M3M)	
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: 059QF0KO0R)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PROPANEDIOL (UNII: 5965N8W85T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENYLPROPANOL (UNII: 0F897O3O4M)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	

**Product Characteristics**

<b>Color</b>	brown	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81110-	1 in 1 CARTON	01/20/2020	

751-02	1 III I CARTON	01/20/2020	
1 NDC:81110-751-01	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
<b>Marketing Information</b>			
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)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		050383046	api manufacture(81110-751)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		623502044	manufacture(81110-751) , api manufacture(81110-751)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		118812921	manufacture(81110-751) , pack(81110-751)

Revised: 3/2023

Iliia, Inc.