





**Sunsolve MD Mineral Shade Amber 002**



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# SUNSOLVE MD MINERAL SHADE AMBER 002

zinc oxide lotion

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:84878-402
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

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



## Inactive Ingredients

Ingredient Name	Strength
<b>TRILAURETH-4 PHOSPHATE</b> (UNII: M96W2OLL2V)	
<b>BUTYLOCTYL SALICYLATE</b> (UNII: 2EH13UN8D3)	
<b>OCTYLDODECYL NEOPENTANOATE</b> (UNII: X8725R883T)	
<b>ISODODECANE</b> (UNII: A8289P68Y2)	
<b>.ALPHA.-BISABOLOL, (+)-</b> (UNII: 105S6I733Z)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>NIACINAMIDE</b> (UNII: 25X5118RD4)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>DIMETHICONOL/PROPYLSILSESQUIOXANE/SILICATE CROSSPOLYMER (450000000 MW)</b> (UNII: 9KB5R958PB)	
<b>PROPANEDIOL</b> (UNII: 5965N8W85T)	
<b>TETRASODIUM GLUTAMATE DIACETATE</b> (UNII: 5EHL50I4MY)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>POLYMETHYLSILSESQUIOXANE (4.5 MICRONS)</b> (UNII: 59Z907ZB69)	
<b>ALKYL (C12-15) BENZOATE</b> (UNII: A9EJ3J61HQ)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>ALLANTOIN</b> (UNII: 344S277G0Z)	
<b>PEG-10</b> (UNII: 761NX2Q08Y)	
<b>CAPRYLYL METHICONE</b> (UNII: Q95M2P1KJL)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>LAURYL PEG-8 DIMETHICONE (300 CPS)</b> (UNII: ELL2U7K8T8)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FERROSFERRIC OXIDE</b> (UNII: XM0M87F357)	
<b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)	

## Product Characteristics

<b>Color</b>	pink	<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:84878-402-02	1 in 1 CARTON	11/05/2024	
1	NDC:84878-402-01	28 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

## Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M020	11/05/2024	

**Labeler** - Sunsolve MD Inc (119376976)

**Registrant** - Nanophase Technologies Corporation (623502044)

**Establishment**

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		050383046	api manufacture(84878-402)

**Establishment**

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		118812921	pack(84878-402) , manufacture(84878-402)

**Establishment**

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		623502044	api manufacture(84878-402) , manufacture(84878-402)

Revised: 11/2024

Sunsolve MD Inc