

SUNSOVLE MD VOLUMIZE AND RESTORE- zinc oxide lotion
Sunsolve MD Inc

Sunsofle MD Volumize and Restore

Sunsofle MD Volumize and Restore



Sunsove MD Volumize and Restore



Sunsole MD Volumize and Restore



Sunsole MD Volumize and Restore



Sunsovl MD Volumize and Restore



Sunsolve MD Volumize and Restore



Sunsovlé MD Volumize and Restore



Sunsovl MD Volumize and Restore



SUNSOVLE MD VOLUMIZE AND RESTORE

zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84878-823
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	110 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ETHYL STEARATE (UNII: C64RTC734W)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
.ALPHA.-BISABOLOL, (+)- (UNII: 10556I733Z)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
SQUALANE (UNII: GW89575KF9)	
ISODODECANE (UNII: A8289P68Y2)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
ETHYL OLEATE (UNII: Z2Z439864Y)	
RUTIN (UNII: 5G06TVY3R7)	
PENTAERYTHRITYL TETRA-DI-T-BUTYL HYDROXYHYDROCINNAMATE (UNII: 255PIF62MS)	
POLYMETHYLSILSESQUIOXANE (4.5 MICRONS) (UNII: 59Z907ZB69)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
LIMNANTHES ALBA (MEADOWFOAM) SEED OIL (UNII: 412ZHA4T4Y)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)	
ETHYL PALMITATE (UNII: IRD3M534ZM)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
C13-15 ALKANE (UNII: 114P5I43UJ)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
ETHYL LINOLEATE (UNII: MJ2YTT4J8M)	

Product Characteristics

Color	yellow	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84878-823-02	1 in 1 CARTON	11/05/2024	
1	NDC:84878-823-01	50 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 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(84878-823)

Establishment

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		118812921	manufacture(84878-823) , pack(84878-823)

Establishment

Name	Address	ID/FEI	Business Operations
Nanophase Technologies Corporation		623502044	manufacture(84878-823) , api manufacture(84878-823)

Revised: 11/2024

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