

SUNSOLVE MD BALANCE AND CLEAR- zinc oxide lotion

Sunsolve MD Inc

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Portia Antonia Alexis
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Sunsolve MD Balance and Clear



Designed by
Portia Antonia Alexis
 -OTCFD3AD17448-

1/30/2024 | 12:32 AM CST

SUNSOLVE MD BALANCE AND CLEAR

zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84878-997
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
CAMELLIA SINENSIS LEAF OIL (UNII: VC855RRT77)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
EUTERPE OLERACEA FRUIT OIL (UNII: Z0W6766A2W)	
CARAPA GUIANENSIS SEED OIL (UNII: Y82418EH2I)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
COPAIFERA OFFICINALIS (BALSAM COPAIBA) RESIN (UNII: 1VH544O5AT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
JOJOBA OIL (UNII: 724GKU717M)	
ORYZA SATIVA WHOLE (UNII: 84IVV0906Z)	
TRIDECYL SALICYLATE (UNII: AZQ08K38Z1)	
TOCOPHEROL (UNII: R0ZB2556P8)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
COCO-CAPRYLATE (UNII: 4828G836N6)	
NYLON-12 (UNII: 446U8J075B)	
PHENYLPROPANOL (UNII: 0F897O3O4M)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
METHYLPROPANEDIOL (UNII: N8F53B3R4R)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
ALLANTOIN (UNII: 344S277G0Z)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ROSMARINUS OFFICINALIS FLOWER (UNII: NR1A27F29O)	
.ALPHA.-BISABOLOL, (+)- (UNII: 105S6I733Z)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PROPANEDIOL (UNII: 5965N8W85T)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POLYGLYCERYL-4 DIISOSTEARATE/POLYHYDROXYSTEARATE/SEBACATE (UNII: 687U3PEB2Y)	
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)	
STEARALKONIUM HECTORITE (UNII: O LX698AH5P)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84878-997-02	1 in 1 CARTON	11/06/2024	
1	NDC:84878-997-01	48 mL in 1 BOTTLE, PUMP; 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	11/06/2024	

Labeler - Sunsolve MD Inc (119376976)

Revised: 11/2025

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