

NU-DERM SUN SHIELD SPF 50 SUNSCREEN- zinc oxide and octinoxate lotion
OMP, 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.

NU-DERM[®]
SUN SHIELD
SPF 50

Drug Facts

<i>Active Ingredients</i>	<i>Purpose</i>
Octinoxate 7.5%	Sunscreen
Zinc Oxide 10.5%	Sunscreen

Uses

- helps prevent sunburn
- higher SPF provides more sunburn protection
- provides high protection against sunburn

Warnings

For external use only.

When using this product

- keep out of eyes. rinse with water to remove.

Stop use and ask a doctor if

- rash or irritation develops and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply liberally 30 minutes before sun exposure and as needed.
- Children under 6 months of age: ask a doctor.
- Reapply as needed or after towel drying, swimming, or perspiring.

Other Information

- Store at controlled room temperature: 15°C-30° (59°F-86°F).
- Sun alert: Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

Inactive Ingredients

Water, Cyclopentasiloxane, PEG-10 Dimethicone, Pentylene Glycol, Stearyl Alcohol, Phenyl Trimethicone, PEG-40 Stearate, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Sodium Dihydroxycetyl Phosphate, Citric Acid, Squalane, Cetareth-20, Polysilicone-11, Dimethicone

Crosspolymer-3, Chlorphenesin, Cetearyl Alcohol, Ethoxylated Sorbitan Ester, Xanthan Gum, 1,2-Hexanediol, Caprylyl Glycol, Potassium Sorbate, Sodium Benzoate, Polysorbate 60, Tetrahexyldecyl Ascorbate, Disodium EDTA, Methylisothiazolinone, Hydrogenated Palm Glycerides, Ubiquinone, Tropolone.

Questions?

1.800.636.7546

Monday–Friday 9:00 AM–4:00 PM Pacific Time

Distributed by OMP, Inc.

PRINCIPAL DISPLAY PANEL - 89 mL Tube Carton

**OBAGI®
MEDICAL**

Nu-Derm®

Sun Shield

SPF 50

Sunscreen Lotion

High UVA/PA+++

UVB

6

AM

NON-COMEDOGENIC

DERMATOLOGIST TESTED

3 fl. oz. (89 mL)

OBAGI
MEDICAL

Nu-Derm®

Sun Shield
SPF 50

Sunscreen Lotion
High UVA/PA+++
UVB

6
AM



3 fl. oz. (89 mL)

Nu-Derm Sun Shield SPF 50 combines UVB absorption and UVA blockage for high sunscreen protection in an elegant, matte finish. Sheer, PABA free, and fragrance free for all skin types.

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Questions? 1.800.836.7548
Monday-Friday 9:00 AM-4:00 PM Pacific Time

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((lot #)) ((Expiration))

NU-DERM SUN SHIELD SPF 50 SUNSCREEN

zinc oxide and octinoxate lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62032-124
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	105 mg in 1 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
WATER (UNII: 059QF0KO0R)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
STEARYL ALCOHOL (UNII: 2KR8914H1Y)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
HYDROGENATED PALM GLYCERIDES (UNII: YCZ8EM144Q)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
TROPOLONE (UNII: 7L6DL16P1T)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	
UBIDECARENONE (UNII: EJ27X76M46)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SQUALANE (UNII: GW89575KF9)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62032-124-10	1 in 1 CARTON		
1		89 mL in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part352	01/05/2011	

Labeler - OMP, Inc. (790553353)**Establishment**

Name	Address	ID/FEI	Business Operations
Milbar Laboratories		195556790	MANUFACTURE(62032-124)

Revised: 1/2012

OMP, Inc.