NU-DERM PHYSICAL UV BROAD SPECTRUM SPF 32- zinc oxide lotion
Obagi Cosmeceuticals LLC

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.

OBAGI
NU-DERM®
AM
PHYSICAL
UV
BROAD SPECTRUM
SPF 32
SUNSCREEN LOTION

Drug Facts

Active ingredient	***Purpose***
Zinc Oxide 18.5%	Sunscreen

Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see ***Directions***), decreases the risk of skin cancer and early skin aging caused by the sun

Warnings

- **For external use only**
- **Do not use** on damaged or broken skin
- **Stop use and ask a doctor** if rash occurs
- **When using this product** keep out of eyes. Rinse with water to remove.
- **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure
- use a water resistant sunscreen if swimming or sweating
- reapply at least every 2 hours
- children under 6 months: Ask a doctor
- **Sun Protection Measures.** Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF value of 15 or higher and other sun protection measures including:
    - limit time in the sun, especially from 10 a.m.-2 p.m.
    - wear long-sleeved shirts, pants, hats, and sunglasses

Inactive ingredients

beeswax, butylene glycol, cetyl dimethicone, cetyl PEG/PPG-10/1 dimethicone, dimethicone, disodium EDTA, epilobium angustifolium flower/leaf/stem extract, ethylhexyl stearate, glycereth-26, hydrogenated castor oil, isopropyl palmitate, methylparaben, propylparaben, sodium chloride, tocopherol acetate, triethoxycaprylylsilane, water

Other information

- store at controlled room temperature: 15°C-25°C (59°F-77°F)
- protect this product from excessive heat and direct sun

Questions or comments?

1.800.636.7546
Monday-Friday 9 a.m.-4 p.m. Pacific Time

Obagi Nu-Derm is a registered trademark of OMP, Inc.

Distributed by OMP, Inc., Long Beach, CA 90806

www.obagi.com
Made in USA

10701810Z 7018

PRINCIPAL DISPLAY PANEL - 57 g Bottle Label

OBAGI
NU-DERM®

AM
PHYSICAL
UV
BROAD SPECTRUM
SPF 32

6

Zinc Oxide 18.5%

SUNSCREEN LOTION

Net wt. 2 oz. (57 g)

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PHYSICAL UV

BROAD SPECTRUM SPF 32

6

Zinc Oxide 18.5%

SUNSCREEN LOTION

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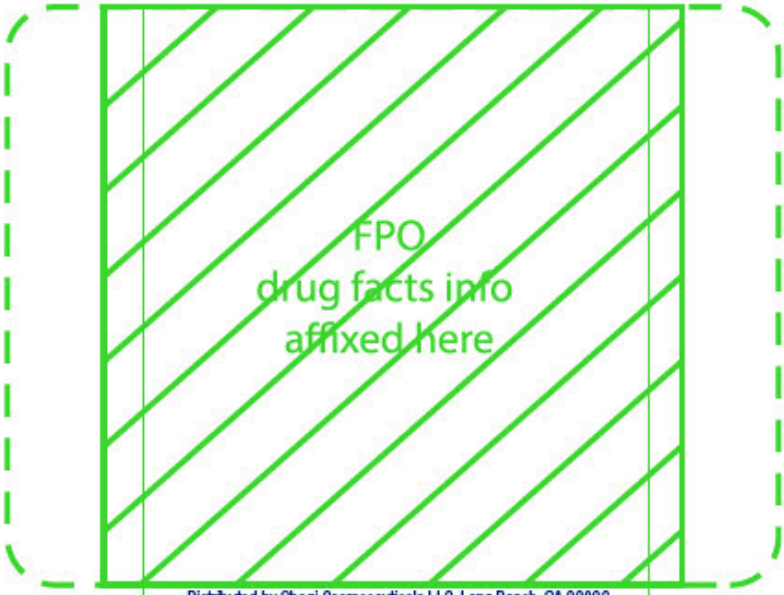

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(continued on reverse)

Drug Facts (continued)

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Questions or comments?

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Monday–Friday 9 a.m.–4 p.m. Pacific Time

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PEEL HERE for complete Drug Facts information

NU-DERM PHYSICAL UV BROAD SPECTRUM SPF 32

zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62032-070
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	185 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
WATER (UNII: 059QF0KO0R)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
GLYCERETH-26 (UNII: NNE56F2N14)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
EPILOBIUM ANGUSTIFOLIUM FLOWERING TOP (UNII: 08H094218D)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 5) (UNII: 035JKJ76MT)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	

Product Characteristics

Color	WHITE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:62032-070	57 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination		

1	NDC:62032-070-18	57 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2004	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/01/2004	

Labeler - Obagi Cosmeceuticals LLC (790553353)

Establishment

Name	Address	ID/FEI	Business Operations
Swiss- American CDMO, LLC		080170933	MANUFACTURE(62032-070)

Revised: 1/2019 Obagi Cosmeceuticals LLC