

**CONDITION AND ENHANCE PHYSICAL UV BLOCK SPF 32- zinc oxide cream
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.

**OBAGI®
MEDICAL
CONDITION & ENHANCE
physical uv block
SPF 32**

DRUG FACTS

Active Ingredient

Zinc Oxide 18.5%.

Warning

For external use only.

See bottom of outer package.

Directions

- Apply liberally to all exposed areas 15 minutes prior to sun exposure and as needed.
- For children less than six months of age, consult a physician.

Other Information

Store at controlled room temperature: 15°-30°C (59°-86°F).

Inactive Ingredients

See bottom of outer package.

Dist. by OMP Inc. Long Beach, CA 90802

PRINCIPAL DISPLAY PANEL - 57 g Bottle Label

**OBAGI®
MEDICAL
CONDITION & ENHANCE
AM 6**

physical uv block

SPF 32

Broad-Spectrum UVA/UVB
Physical Sunblock
18.5% Zinc Oxide

NET WT. 2 OZ. (57 g)

OBAGI[®]
MEDICAL

CONDITION & ENHANCE
6
AM

physical uv block
SPF 32

Broad-Spectrum UVA/UVB
Physical Sunblock
18.5% Zinc Oxide

NET WT. 2.0Z. (57 g)

Physical UV Block SPF 32 is specially formulated for use with the Obagi[®] Condition & Enhance System for skin transformation. It provides exceptional protection against sunburn caused by UVB rays and premature aging caused by UVA-1 rays.

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Dist. by OMP Inc. Long Beach, CA 90802
Made in U.S.A. 7843 10784311U

CONDITION AND ENHANCE PHYSICAL UV BLOCK SPF 32

zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62032-118
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: A2I8C7H9T)	
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
1	NDC:62032-118-36	57 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part352	01/01/2004	

Labeler - OMP, Inc. (790553353)

Establishment

Name	Address	ID/FEI	Business Operations
Swiss-American Products		611921669	MANUFACTURE(62032-118)

Revised: 12/2011

OMP, Inc.