CONDITION AND ENHANCE SYSTEM TRAVEL-SIZE SURGICAL- hydroquinone, octinoxate and zinc oxide
OMP, INC.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Obagi® Condition & Enhance Clear
(Hydroquinone USP, 4%)
Skin Bleaching Cream
Obagi® Condition & Enhance Blender®
(Hydroquinone USP, 4%)
Skin Bleaching Cream

Rx Only

FOR EXTERNAL USE ONLY

DESCRIPTION

Hydroquinone is 1,4-benzenediol. Hydroquinone occurs as fine, white needles. The drug is freely soluble in water and in alcohol. Chemically, hydroquinone is designated as p-dihydroxybenzene; the empirical formula is $C_6 H_6 O_2$; molecular weight is 110.0.

Obagi® Condition & Enhance Blender contains Hydroquinone USP 40 mg/gm in a base of purified water, glycerin, cetyl alcohol, PPG-2 myristyl ether propionate, sodium lauryl sulfate, TEA-salicylate, lactic acid, phenyl trimethicone, tocopheryl acetate, sodium metabisulfite, ascorbic acid, methylparaben, saponins, disodium EDTA, BHT, and propylparaben.

$C_6H_6O_2$

Obagi® Condition & Enhance Clear contains Hydroquinone USP 40 mg/gm in a base of purified water, cetyl alcohol, glycerin, sodium lauryl sulfate, stearyl alcohol, tocopheryl acetate, ascorbic acid, sodium metabisulfite, lactic acid, saponins, disodium EDTA, methylparaben, BHT, propylparaben, and butylparaben.

CLINICAL PHARMACOLOGY

Topical application of hydroquinone produces a reversible depigmentation of the skin by inhibition of the enzymatic oxidation of tyrosine to 3, 4-dihydroxyphenylalanine (dopa) and suppression of other melanocyte metabolic processes.

Exposure to sunlight or ultraviolet light will cause repigmentation of the bleached areas, which may be prevented by the use of sunblocking agents or sunscreen agents contained in Obagi Condition & Enhance.

INDICATIONS AND USAGE

The gradual bleaching of hyperpigmented skin conditions such as chloasma, melasma, freckles, senile lentigines, and other unwanted areas of melanin hyperpigmentation.

CONTRAINDICATIONS

Prior history of sensitivity or allergic reaction to this product or any of its ingredients. The safety of topical hydroquinone use during pregnancy or in children (12 years and under) has not been established.

WARNINGS

Caution

Hydroquinone is a skin bleaching agent which may produce unwanted cosmetic effects if not used as directed. The physician should be familiar with the contents of this insert before prescribing or dispensing this medication.

Test for skin sensitivity before using by applying a small amount to an unbroken patch of skin and check in 24 hours. Minor redness is not a contraindication, but where there is itching or vesicle formation or excessive inflammatory response, further treatment is not advised. Close patient supervision is recommended.

Avoid contact with eyes. In case of accidental contact, patient should rinse eyes thoroughly with water and contact physician. A bitter taste and anesthetic effect may occur if applied to lips.

Sunscreen use is an essential aspect of hydroquinone therapy because even minimal sunlight exposure sustains melanocytic activity.

Warning

Contains sodium metabisulfite, a sulfite that may cause serious allergic type reactions (e.g., hives, itching, wheezing, anaphylaxis, severe asthma attacks) in certain susceptible persons.

PRECAUTIONS

(SEE WARNINGS)

General

Treatment should be limited to relatively small areas of the body at one time since some patients experience a transient skin reddening and a mild burning sensation which does not preclude treatment.

Pregnancy Category C

Animal reproduction studies have not been conducted with topical hydroquinone. It is also not known whether hydroquinone can cause fetal harm when used topically on a pregnant woman or affect reproductive capacity. It is not known to what degree, if any, topical hydroquinone is absorbed systemically. Topical hydroquinone should be used on pregnant women only when clearly indicated.

Nursing mothers

It is not known whether topical hydroquinone is absorbed or excreted in human milk. Caution is advised when topical hydroquinone is used by a nursing mother.

Pediatric usage

Safety and effectiveness in children below the age of 12 years have not been established.

ADVERSE REACTIONS

No systemic adverse reactions have been reported. Occasional hypersensitivity (localized contact dermatitis) may occur, in which case the medication should be discontinued and the physician notified immediately.

DOSAGE AND ADMINISTRATION

A thin application should be applied to the affected area twice daily or as directed by a physician. If no improvement is seen after three (3) months of treatment, use of this product should be discontinued. Sun exposure should be limited by using a sunscreen agent, a sun blocking agent, or protective clothing to cover bleached skin when using and after using this product in order to prevent repigmentation.

HOW SUPPLIED

Obagi Condition and Enhance Blender is available as follows:

2 oz. (57 gm) bottle	NDC 62032-115-36
1 oz. (28.5 gm) bottle	NDC 62032-115-10

Obagi Condition and Enhance Clear is available as follows:

2 oz. (57 gm) bottle	NDC 62032-117-36

Store at 25°C (77°F); excursion permitted to 15°C-30°C (59°F-86°F).

OBAGI®
MEDICAL

OMP, Inc.
Long Beach, CA 90802
USA
1-800-636-7546

80707910U Rev. 6/07

PRINCIPAL DISPLAY PANEL - Kit Carton

OBAGI®
CONDITION & ENHANCE SYSTEM

For Use with Surgical Procedures

Travel Size

OBAGI®

CONDITION & ENHANCE SYSTEM

For Use with Surgical Procedures

Travel Size

OBAGI®
CONDITION & ENHANCE SYSTEM
For Use with Surgical Procedures

OBAGI MEDICAL

What to Expect

The desire for healthy, youthful and attractive skin is why you are undergoing facial aesthetic procedures. The Obagi® Condition & Enhance System uses proven ingredients and penetrating technologies to help ensure a successful outcome. This system has been shown to improve the overall results of facial aesthetic procedures.

The length of time that it takes to condition your skin varies. You should begin to notice visible improvements to your skin within a six week period.

It is important to understand that you will experience some reactions during your skin conditioning process. These reactions are normal and part of the transformation process. You may experience one or more of these symptoms:

- Dryness
- Redness
- Skin texture and appearance of wrinkles may temporarily worsen
- Exfoliation
- Sensitive skin
- Acne may temporarily worsen

In order to maximize the results of your procedure, it is very important that you follow your personal program as indicated by a skincare professional.

Your physician should discuss with you any reactions related to your facial procedure, as these vary by the type of procedure you have received.

Daily Product Routines

a.m.
Morning

	PRODUCT	QUANTITY
1	**Gentle Cleanser**	Entire face
2	**Toner**	Apply liberally with cotton pad to the entire face
3	**Clear**	☐ ½ gm ☐ 1 gm
4	**Exfoderm*** (Avoid the eye area as stinging can occur)	☐ ½ gm ☐ 1 gm
6	**Healthy Skin Protection SPF 35** or **Physical UV Block SPF 32**	Entire face

Note: Morning routine does not include step 5.

p.m.
Evening

	PRODUCT	QUANTITY
1	**Gentle Cleanser**	Entire face
2	**Toner**	Apply liberally with cotton pad to the entire face
3	**Clear**	☐ ½ gm ☐ 1 gm
5	**Blender***	☐ ½ gm
	Tretinoin Cream (apply as prescribed)	☐ ½ gm ☐ 1 gm

Note: Evening routine does not include steps 4 or 6.

Measuring

Squeeze the product onto your finger, using the bars below as reference:

½ gram

1 gram

Or use the distance from the tip of your pinky finger to the first joint as an approximate measurement of ½ gram of product.

OBAGI®
MEDICAL

OBAGI®
CONDITION & ENHANCE SYSTEM
For Use with Surgical Procedures

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For Use with Surgical Procedures

Skincare and protection should be a fundamental part of good health and beautiful, vibrant skin. Your skin needs special care to stay young and healthy. The Obagi® Condition & Enhance System is the leading prescription skincare program that works at the cellular level for skin that looks and acts younger and healthier. Whether you're planning a surgical or nonsurgical procedure, the Obagi Condition & Enhance System uses proven ingredients and penetrating technologies to help ensure a successful outcome.

Gentle Cleanser

Obagi Gentle Cleanser is a soap-free cleanser that gently removes impurities, oil and make-up. This special formula prepares skin for the transformation process. It is designed for use with surgical procedures.

DIRECTIONS: Apply to damp face and neck with moistened fingertips in the morning and evening. Rinse completely with warm water.

INGREDIENTS: Purified water, cocamidopropyl betaine, sodium lauroyl oat amino acids, sodium laureth sulfate, glycerin, aloe barbadensis leaf juice, glycereth-7 prunus armeniaca (apricot) kernel oil, panthenol, acrylates/C10-30 alkyl acrylate crosspolymer, oleyl lactate, ethoxydiglycol, triethanolamine, salvia officinalis (sage) leaf extract, borago officinalis extract, phenoxyethanol, methylparaben, ethylparaben, butylparaben, propylparaben, isobutylparaben, saponins, fragrance, FD&C Yellow No. 5.

Toner (Skin Reparation)

Obagi Toner, formulated with alum (a natural astringent), adjusts the pH of the skin for optimal penetration of the treatment ingredients in the System.

DIRECTIONS: Apply after cleansing in the morning and evening. Saturate cotton pad and wipe gently over face and neck. Avoid eyelid area. Do not rinse off.

INGREDIENTS: Purified water, aloe barbadensis leaf juice, hamamelis virginiana (witch hazel) distillate, potassium alum, sodium PCA, panthenol, DMDM hydantoin, polysorbate 80, allantoin, salvia officinalis (sage) leaf extract, borago officinalis extract, calendula officinalis flower extract, saponins, iodopropynyl butylcarbamate, fragrance, FD&C Blue No. 1.

Clear (Skin Bleaching and Corrector Cream) NDC 62032-117-96

Obagi Clear corrects uneven skin color and brown spots, continuing the transformation process.

INDICATIONS: For the gradual bleaching of hyperpigmented skin conditions such as chloasma, melasma, freckles, senile lentigines and other unwanted areas of melanin hyperpigmentation.

DOSAGE AND ADMINISTRATION: A thin layer should be applied to the affected area in the morning and evening, or as directed by a physician. If no improvement is seen after three months of treatment, use of this product should be discontinued. Sun exposure should be limited by using a sunscreen agent, a sun blocking agent, or protective clothing to cover bleached skin during and after usage of this product in order to prevent repigmentation.

WARNINGS: Avoid contact with eyes. In case of accidental contact, patient should rinse eyes thoroughly with water and contact a physician. A bitter taste and anesthetic effect may occur if applied to lips. Sunscreen use is an essential aspect of hydroquinone therapy because even minimal sunlight exposure sustains melanocytic activity.

CAUTION: Contains sodium metabisulfite, a sulfite that may cause serious allergic-type reactions (e.g., hives, itching, wheezing, anaphylaxis, severe asthma attacks) in certain susceptible persons.

INGREDIENTS: Hydroquinone USP 4% mg/gm in a base of purified water, cetyl alcohol, glycerin, sodium lauryl sulfate, stearyl alcohol, tocophenyl acetate, ascorbic acid, sodium metabisulfite, lactic acid, sapcorin, disodium EDTA, methylparaben, BHT, propylparaben and butylparaben.

See enclosed Package Insert for full prescription information.

Rx ONLY. FOR EXTERNAL USE ONLY.

Exfoderm® (Skin Smoothing Lotion)

Obagi Exfoderm exfoliates dead surface skin cells and smoothes roughness, aiding in penetration of the treatment ingredients in the System.

DIRECTIONS: Apply to face in the morning, following the application of Obagi Clear, as directed by a physician. Follow with the appropriate Obagi Condition & Enhance sun protection.

CAUTION: FOR EXTERNAL USE ONLY. Avoid contact with the eyes. A mild burning sensation of the skin is to be expected. If burning is severe, discontinue use and consult a physician.

INGREDIENTS: Purified water, ethoxydiglycol, phytic acid, glycerin, cetearyl alcohol, glyceryl stearate, PEG-100 stearate, canola oil, isohexadecane, magnesium aluminum silicate, potassium cetyl phosphate, cetyl alcohol, bis-diglyceryl polyacyladipate-2, dimethicone, polysorbate 60, PEG-150 stearate, steareth-20, xanthan gum, glycereth-7, tocopheryl acetate, saponins, phenoxyethanol, methylparaben, ethylparaben, butylparaben, propylparaben, isobutylparaben.

Blender® (Skin Lightener and Blending Cream) NDC 62032-115-10

Obagi Blender maximizes skin transformation when prescribed with tretinoin cream. Blender evens skin color and tone, making skin look and act younger and healthier.

INDICATIONS: For the gradual bleaching of hyperpigmented skin conditions such as chloasma, melasma, freckles, senile lentigines and other unwanted areas of melanin hyperpigmentation.

DOSAGE AND ADMINISTRATION: A thin layer should be applied to the affected area in the morning and evening, or as directed by a physician. If no improvement is seen after three months of treatment, use of this product should be discontinued. Sun exposure should be limited by using a sunscreen agent, a sun blocking agent or protective clothing to cover bleached skin during and after usage of this product in order to prevent repigmentation.

WARNINGS: Avoid contact with eyes. In case of accidental contact, patient should rinse eyes thoroughly with water and contact a physician. A bitter taste and anesthetic effect may occur if applied to lips. Sunscreen use is an essential aspect of hydroquinone therapy because even minimal sunlight exposure sustains melanocytic activity.

CAUTION: Contains sodium metabisulfite, a sulfite that may cause serious allergic-type reactions (e.g., hives, itching, wheezing, anaphylaxis, severe asthma attacks) in certain susceptible persons.

INGREDIENTS: Hydroquinone USP 4% mg/gm in a base of purified water, glycerin, cetyl alcohol, PPG-2 myristyl ether propionate, sodium lauryl sulfate, TEA-salicylate, lactic acid, phenyl trimethicone, tocopheryl acetate, sodium metabisulfite, ascorbic acid, methylparaben, saponins, disodium EDTA, BHT and propylparaben.

See enclosed Package Insert for full prescription information.

Rx ONLY. FOR EXTERNAL USE ONLY.

Healthy Skin Protection SPF 35

Obagi Healthy Skin Protection contains 9% micronized zinc oxide to protect the newer, healthier skin created by skin transformation. This high concentration of micronized zinc oxide provides protection against long UVA rays linked to deep premature aging.

DIRECTIONS: Apply liberally to all exposed areas. Apply at least 15 minutes before sun exposure and reapply frequently after prolonged swimming, excessive perspiration, vigorous activity or toweling.

DRUG FACTS

Active Ingredients:	**Purpose:**
Octinoxate, 25%, Zinc Oxide, 9%	Sunscreens

Uses: ■ Helps prevent sunburn. ■ Higher SPF gives more sunburn protection.

Warnings: For external use only.

When using this product: ■ Keep out of eyes. Rinse with water to remove.
Stop use and ask a physician if ■ Rash or irritation develops and persists.
Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions: ■ Apply generously and evenly 15 minutes before sun exposure. ■ For children under six months of age: Ask a physician. ■ Reapply as needed or after towel drying, swimming or perspiring.

Other Information: ■ Store at controlled room temperature: 15°-30°C (59°-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: Butylparaben, cetearyl alcohol, citric acid, C13-14 isoparaffin, diethanolamine cetyl phosphate, disodium edetate, ethylparaben, isobutylparaben, isopropyl palmitate, laureth-7, methylparaben, octyl stearate, phenoxyethanol, polyacrylamide, polysorbate 61, propylparaben, purified water, sodium hydroxide, trihydroxystearin.

Physical UV Block SPF 32

Containing 18.5% micronized zinc oxide, this chemical-free formula provides protection without irritation.

DIRECTIONS: Apply generously and evenly 15 minutes before sun exposure. For children under six months of age, ask a physician. Reapply as needed.

DRUG FACTS

Active Ingredient:	**Purpose:**
Zinc Oxide, 18.5%	Sunscreen

Warnings: For external use only.

When using this product: ■ Keep out of eyes. Rinse with water to remove.
Stop use and ask a physician if ■ Rash or irritation develops and persists.
Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions: ■ Apply liberally to all exposed areas 15 minutes prior to sun exposure and as needed. ■ For children under six months of age: Ask a physician.

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

Inactive Ingredients: Beeswax, butylene glycol, cetyl dimethicone, cetyl PEG/PPG-10/1 dimethicone, dimethicone, disodium EDTA, glycereth-26, hydrogenated castor oil, isopropyl palmitate, methylparaben, octyl stearate, propylparaben, purified water, sodium chloride, trihydroxystearin, tocopherol acetate, wheat starch extract.

Store at 25°C (77°F); excursion permitted to 15°-30°C (59°-86°F).

Dist. by OMP, Inc., Long Beach, CA 90806
Made in U.S.A. 4070801 0U
Obagi is a registered trademark of OMP, Inc.

Obagi products are physician dispensed and should be used under the guidance of your skincare specialist.

RECOMMENDED
SKIN CANCER FOUNDATION

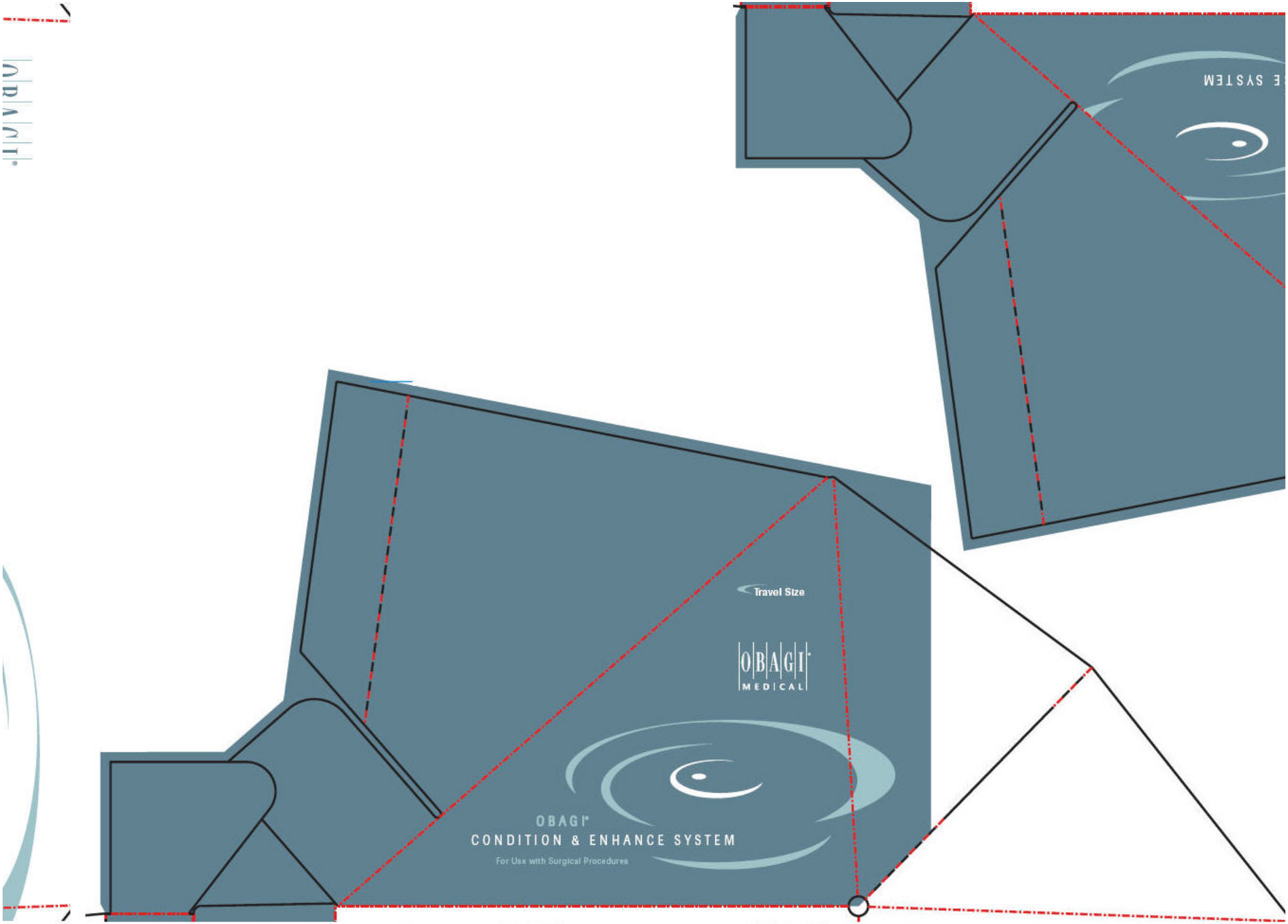
OBAGI®
MEDICAL

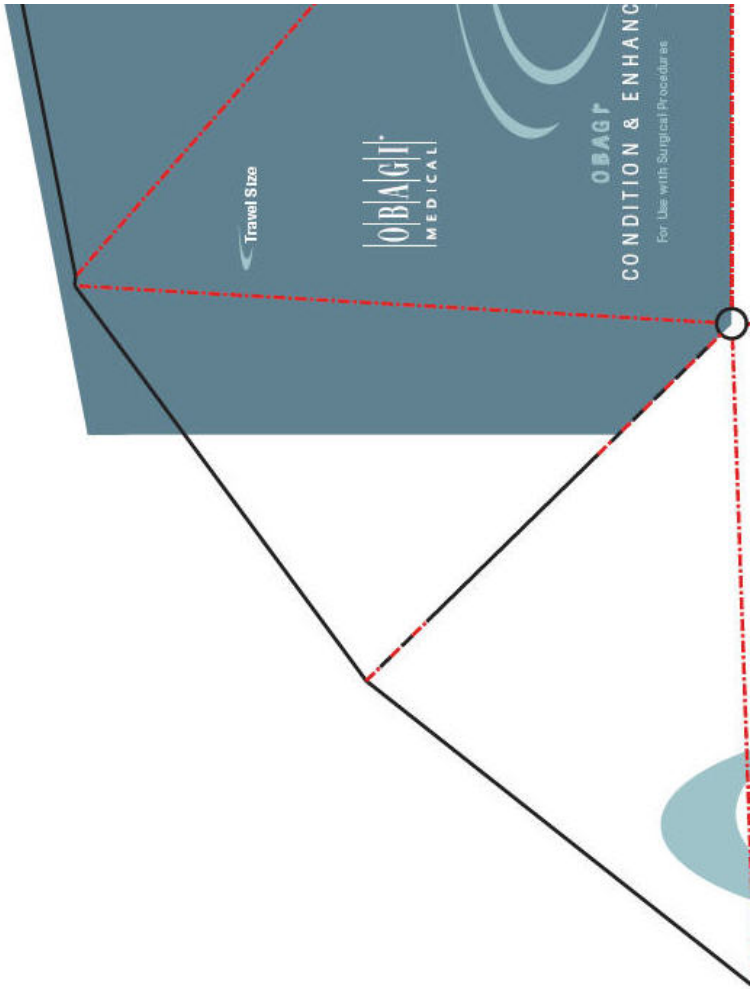
This Obagi® Condition & Enhance System includes:

Gentle Cleanser	Toner	Clear	Exfoderm®	Blender®	Healthy Skin Protection	Physical UV Block	Patient Instructions
2 FL. OZ. (60 mL)	2 FL. OZ. (60 mL)	NET WT. 2 OZ. (57 g) Hydroquinone USP, 4% **Rx Only**	NET WT. 1 OZ. (28.5 g)	NET WT. 1 OZ. (28.5 g) Hydroquinone USP, 4% **Rx Only**	1 FL. OZ. (30 mL) **SPF 35**	NET WT. 2 OZ. (57 g) **SPF 32**	

OBAGI®
CONDITION & ENHANCE SYSTEM
For Use with Surgical Procedures

OBAGI®
MEDICAL


Travel Size
OBAGI MEDICAL
OBAGI
CONDITION & ENHANCE SYSTEM
For Use with Surgical Procedures



CONDITION AND ENHANCE SYSTEM TRAVEL-SIZE SURGICAL

hydroquinone, octinoxate and zinc oxide kit

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:62032-508

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62032-508-04	1 in 1 CARTON		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 BOTTLE, PLASTIC	28.5 g
Part 2	1 BOTTLE, PLASTIC	57 g
Part 3	1 BOTTLE, PLASTIC	30 mL
Part 4	1 BOTTLE, PLASTIC	60 mL
Part 5	1 BOTTLE, PLASTIC	60 mL
Part 6	1 BOTTLE, PLASTIC	57 g
Part 7	1 BOTTLE, PLASTIC	28.5 g

Part 1 of 7

CONDITION AND ENHANCE BLENDER SKIN LIGHTENER AND BLENDING

hydroquinone cream

Product Information	
Item Code (Source)	NDC:62032-115
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	40 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PPG-2 MYRISTYL ETHER PROPIONATE (UNII: 88R97D8U8A)	
TROLAMINE SALICYLATE (UNII: H8O4040BHD)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTIC ACID (UNII: 33X04XA5AT)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
WATER (UNII: 059QF0KO0R)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62032-115-10	28.5 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		01/01/1988	

Part 2 of 7

CONDITION AND ENHANCE CLEAR SKIN BLEACHING AND CORRECTOR

hydroquinone cream

Product Information

Item Code (Source)	NDC:62032-117
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	40 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTIC ACID (UNII: 33X04XA5AT)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
WATER (UNII: 059QF0KO0R)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

Product Characteristics

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62032-117-36	57 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		01/01/1988	

Part 3 of 7

CONDITION AND ENHANCE HEALTHY SKIN PROTECTION SPF 35

octinoxate and zinc oxide cream

Product Information	
Item Code (Source)	NDC:62032-119
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	90 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
WATER (UNII: 059QF0KO0R)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
LAURETH-7 (UNII: Z95S6G8201)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
DIETHANOLAMINE CETYL PHOSPHATE (UNII: 4UG0316V9S)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLPARABEN (UNII: 14255EXE39)	
ISOBUTYLPARABEN (UNII: 0QQJ25X58G)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Product Characteristics			
Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62032-119-10	30 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/01/2002	

Part 4 of 7

CONDITION AND ENHANCE GENTLE CLEANSER

cleansing (cold creams, cleansing lotions, liquids, and pads) liquid

Product Information	
Route of Administration	TOPICAL

Other Ingredients		
Ingredient Kind	Ingredient Name	Quantity
INGR	**WATER** (UNII: 059QF0KO0R)	
INGR	**GLYCERIN** (UNII: PDC6A3C0OX)	
INGR	**PHENOXYETHANOL** (UNII: HIE492ZZ3T)	
INGR	**METHYLPARABEN** (UNII: A2I8C7HI9T)	
INGR	**PROPYLPARABEN** (UNII: Z8IX2SC1OH)	
INGR	**BUTYLPARABEN** (UNII: 3QPI1U3FV8)	
INGR	**ETHYLPARABEN** (UNII: 14255EXE39)	
INGR	**ISOBUTYLPARABEN** (UNII: 0QQJ25X58G)	
INGR	**SODIUM LAUROYL OAT AMINO ACIDS** (UNII: FSW2K9B9N5)	
INGR	**COCAMIDOPROPYL BETAINE** (UNII: 5OCF3O11KX)	
INGR	**SODIUM LAURETH SULFATE** (UNII: BPV390UAP0)	
INGR	**ALOE VERA LEAF** (UNII: ZY81Z83H0X)	
INGR	**GLYCERETH-7** (UNII: 3D2Y91QZ2H)	
INGR	**PANTHENOL** (UNII: WV9CM0O67Z)	
INGR	**DIETHYLENE GLYCOL MONOETHYL ETHER** (UNII: A1A1I8X02B)	
INGR	**TROLAMINE** (UNII: 9O3K93S3TK)	

INGR	**SAGE** (UNII: 065C5D077J)	
INGR	**FD&C YELLOW NO. 5** (UNII: I753WB2F1M)	
INGR	**CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)** (UNII: 809Y72KV36)	
INGR	**APRICOT KERNEL OIL** (UNII: 54JB35T06A)	

Product Characteristics			
Color	YELLOW	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		60 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Cosmetic		01/01/1988	

Part 5 of 7

CONDITION AND ENHANCE TONER

face and neck (excluding shaving preparations) liquid

Product Information	
Route of Administration	TOPICAL

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
INGR	**WATER** (UNII: 059QF0KO0R)	
INGR	**GLYCERIN** (UNII: PDC6A3C0OX)	
INGR	**SODIUM PYRROLIDONE CARBOXYLATE** (UNII: 469OTG57A2)	
INGR	**DMDM HYDANTOIN** (UNII: BYR0546TOW)	
INGR	**IODOPROPYNYL BUTYLCARBAMATE** (UNII: 603P14DHEB)	
INGR	**POTASSIUM ALUM** (UNII: 1L24V9R23S)	
INGR	**PANTHENOL** (UNII: WV9CM0O67Z)	
INGR	**SAGE** (UNII: 065C5D077J)	
INGR	**CALENDULA OFFICINALIS FLOWER** (UNII: P0M7O4Y7YD)	
INGR	**POLYSORBATE 80** (UNII: 6OZP39ZG8H)	
INGR	**ALLANTOIN** (UNII: 344S277G0Z)	

INGR	**ALOE VERA LEAF** (UNII: ZY81Z83H0X)	
INGR	**FD&C BLUE NO. 1** (UNII: H3R47K3TBD)	
INGR	**HAMAMELIS VIRGINIANA TOP WATER** (UNII: NT00Y05A2V)	

Product Characteristics			
Color	BLUE	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		60 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Cosmetic		01/01/1988	

Part 6 of 7

CONDITION AND ENHANCE PHYSICAL UV BLOCK SPF 32

zinc oxide cream

Product Information	
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: 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
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	

Part 7 of 7

CONDITION AND ENHANCE EXFODERM

face and neck (excluding shaving preparations) lotion

Product Information	
Route of Administration	TOPICAL

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
INGR	**WATER** (UNII: 059QF0KO0R)	
INGR	**GLYCERIN** (UNII: PDC6A3C0OX)	
INGR	**METHYLPARABEN** (UNII: A2I8C7HI9T)	
INGR	**PROPYLPARABEN** (UNII: Z8IX2SC1OH)	
INGR	**POLYSORBATE 60** (UNII: CAL22UVI4M)	
INGR	**CETOSTEARYL ALCOHOL** (UNII: 2DMT128M1S)	
INGR	**STEARETH-20** (UNII: L0Q8IK9E08)	

INGR	**CANOLA OIL** (UNII: 331KBJ17RK)	
INGR	**ISOHEXADECANE** (UNII: 918X1OUF1E)	
INGR	**MAGNESIUM ALUMINUM SILICATE** (UNII: 6M3P64V0NC)	
INGR	**CETYL ALCOHOL** (UNII: 936JST6JCN)	
INGR	**FYTIC ACID** (UNII: 7IGF0S7R8I)	
INGR	**GLYCERYL MONOSTEARATE** (UNII: 230OU9XXE4)	
INGR	**PEG-100 STEARATE** (UNII: YD01N1999R)	
INGR	**DIMETHICONE** (UNII: 92RU3N3Y1O)	
INGR	**PEG-150 STEARATE** (UNII: 7BSG7DF10Q)	
INGR	**PHENOXYETHANOL** (UNII: HIE492ZZ3T)	
INGR	**BUTYLPARABEN** (UNII: 3QPI1U3FV8)	
INGR	**ETHYLPARABEN** (UNII: 14255EXE39)	
INGR	**ISOBUTYLPARABEN** (UNII: 0QQJ25X58G)	
INGR	**POTASSIUM CETYL PHOSPHATE** (UNII: 03KCY6P7UT)	
INGR	**XANTHAN GUM** (UNII: TTV12P4NEE)	
INGR	**.ALPHA.-TOCOPHEROL ACETATE** (UNII: 9E8X80D2L0)	
INGR	**GLYCERETH-7** (UNII: 3D2Y91QZ2H)	
INGR	**DIETHYLENE GLYCOL MONOETHYL ETHER** (UNII: A1A1I8X02B)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		28.5 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Cosmetic		01/01/1988	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		08/20/2007	

Labeler - OMP, INC. (790553353)

Establishment

Name	Address	ID/FEI	Business Operations
PURETEK CORPORATION		785961046	MANUFACTURE(62032-508) , LABEL(62032-508) , PACK(62032-508)

Establishment

Name	Address	ID/FEI	Business Operations
Ei INC.		105803274	MANUFACTURE(62032-508) , LABEL(62032-508) , PACK(62032-508) , ANALYSIS(62032-508)

Establishment

Name	Address	ID/FEI	Business Operations
Swiss-American Products		611921669	MANUFACTURE(62032-508)

Establishment

Name	Address	ID/FEI	Business Operations
Bay Cities Container Corporation		066229618	RELABEL(62032-508) , REPACK(62032-508)

Revised: 5/2012 OMP, INC.