NU-DERM SYSTEM NORMAL-DRY SKIN TRANSFORMATION- 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.

NU-DERM® SYSTEM NORMAL-DRY SKIN TRANSFORMATION KIT

Gentle Cleanser 6.7 fl. oz. (198 mL.) AM+PM

A mild facial cleanser that provides gentle yet effective cleansing for normal to dry skin. For skin that's beautifully clean and fresh, the Nu-Derm Gentle Cleanser instantly dissolves excess oil, makeup, and other everyday impurities, without stripping your skin of its natural protective oils, and prepares your skin for the next step in your skin care regimen.

Directions

Use twice daily, morning and evening. Massage a small amount of cleanser and water onto skin. Rinse with lukewarm water and gently pat dry.

Warnings

Avoid getting into eyes. **For external use only.**

Keep out of reach of children.

Ingredients

water (aqua), cocamidopropyl betaine, glycerin, sodium lauroyl oat amino acids, aloe barbadensis leaf juice (aloe barbadensis), sodium laureth sulfate, glycereth-7, prunus armeniaca (apricot) kernel oil, panthenol, acrylates/C10-30 alkyl acrylate crosspolymer, oleyl lactate, ethoxydiglycol, triethanolamine, salvia officinalis (sage) leaf extract (salvia officinalis), borago officinalis extract, saponins, phenoxyethanol, methylparaben, ethylparaben, butylparaben, propylparaben, isobutylparaben, fragrance (parfum), yellow 5 (CI 19140)

Toner 6.7 fl. oz. (198 mL.) AM+PM

An essential step in your daily skin care routine, this alcohol-free, non-drying toner helps adjust your skin's pH for increased penetration of product ingredients. Use after cleansing to remove impurities and dead skin cells and to prepare the skin for hydration or appropriate products.

Directions

Use daily, in the morning and evening after cleansing. Saturate a cotton pad and gently wipe over entire face. Do not rinse.

Warnings

Avoid getting into eyes. **For external use only.**

Keep out of reach of children.

Ingredients

water (aqua), hamamelis virginiana (witch hazel) water, aloe barbadensis leaf juice (aloe barbadensis), potassium alum, sodium PCA, panthenol, DMDM hydantoin, polysorbate 80, allantoin, glycerin, salvia officinalis (sage) leaf extract (salvia officinalis), borago officinalis extract, calendula officinalis flower extract (calendula officinalis), saponins, iodopropynyl butylcarbamate, fragrance (parfum), blue 1 (CI 42090)

Clear (Skin Bleaching and Corrector Cream) NDC 62032-101-36 Net wt. 2 oz. (57 g.)
Hydroquinone USP, 4% Rx Only AM+PM

Dark spots may appear on the surface of your skin, but they actually start deep within the skin's layers. This gentle yet effective formula absorbs into the layers of your skin to deliver prescription-strength hydroquinone, helping to correct the appearance of age and sun spots for a healthier, more even complexion.

Indications and usage

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

Dosage and administration

Use daily, in the morning and evening. Squeeze a small amount (approximately 1-2 pea-sized amounts) onto your hand. Apply evenly to the entire face, extending to the hairline, over the ears, and ending with a feathering motion, or as directed by your 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 darkening from reoccurring.

Warnings

Avoid contact with eyes, nose, mouth, or lips. In case of accidental contact, patient should rinse eyes thoroughly with water and contact physician. Sunscreen use is an essential aspect of hydroquinone therapy because even minimal sunlight exposure sustains melanocytic activity.

Contains sodium metabisulfite, a sulfite that may cause allergic-type reactions including anaphylactic symptoms and life-threatening or less severe asthmatic episodes in certain susceptible people. The overall prevalence of sulfite sensitivity in the general population is unknown and probably low. Sulfite sensitivity is seen more frequently in asthmatic than in nonasthmatic people.

Each gram of Obagi Nu-Derm Clear contains:

Active ingredient

Hydroquinone USP, 4% (40 mg/g)

Inactive ingredients

water, cetyl alcohol, glycerin, sodium lauryl sulfate, stearyl alcohol, lactic acid, tocopheryl acetate, ascorbic acid, sodium metabisulfite, disodium EDTA, methylparaben, BHT, propylparaben, saponins, butylparaben

See enclosed Package Insert for full prescribing information.

Rx ONLY. FOR EXTERNAL USE ONLY.

Exfoderm® (Skin Smoothing Lotion) Net wt. 2 oz. (57 g.) AM

A lightweight lotion that exfoliates the top layer of skin, removing dull, old skin cells while promoting new skin cells for a brighter complexion. Specifically developed for normal to dry skin, this gentle, skin-enhancing formula contains a plant acid (phytic acid) to help transform the appearance of damaged skin and reveal your skin's radiance.

Directions

Use daily, in the morning. Squeeze a small amount (approximately 1-2 pea-sized drops) onto your hands. Using your fingertips, apply evenly to the entire face. Massage until completely absorbed.

Warnings

Avoid getting into eyes. **For external use only.**

Keep out of reach of children.

Ingredients

water (aqua), ethoxydiglycol, glycerin, phytic acid, 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

Hydrate (Facial Moisturizer) Net wt. 1.7 oz. (48 g.) AM+PM

A multi-action moisturizer with Hydromanil, a natural moisturizing agent derived from tara seed known to gradually deliver moisture to the skin. Also contains shea butter, mango butter, avocado and glycerin to help combat skin dryness. Suitable for all skin types.

Directions

Apply to face in the morning and evening or as needed.

Warnings

Avoid getting into eyes. **For external use only.**

Keep out of reach of children.

Ingredients

water (aqua), glycerin, caprylic/capric triglyceride, butyrospermum parkii (shea) butter, cyclopentasiloxane, glyceryl stearate, cetyl alcohol, dimethicone, saccharide isomerate, stearic acid, polysilicone-11, glycine soja (soybean) sterols, persea gratissima (avocado) oil, mangifera indica (mango) seed butter, hydrolyzed caesalpinia spinosa gum, caesalpinia spinosa gum, hydrolyzed soybean fiber, sodium stearoyl glutamate, caprylyl glycol, bisabolol, allantoin, tocopherol, tetrahydrodiferuloylmethane, panthenol, carbomer, hexylene glycol, sodium hydroxide, laureth-12, ethylhexylglycerin, phenoxyethanol

Blender® (Skin Lightener and Blending Cream) NDC 62032-100-36 Net wt. 2 oz. (57 g.) Hydroquinone USP, 4% Rx Only PM

A unique formula containing prescription-strength hydroquinone for the gradual lightening of sun spots,

age spots, and other types of hyperpigmentation (discoloration). Specially formulated to optimize the delivery of product ingredients in the Nu-Derm System, this skin lightener helps reduce the signs of aging and correct uneven skin tone. May be used with Tretinoin Cream[1] or Refissa[2] as prescribed by a physician.

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1 Tretinoin cream is indicated for topical application in the treatment of acne vulgaris.

2 Refissa [Tretinoin Cream, USP (Emollient) 0.05%] is indicated as an adjunctive agent for use in the mitigation (palliation) of fine wrinkles, mottled hyperpigmentation, and tactile roughness of facial skin in patients who do not achieve such palliation using comprehensive skin care and sun avoidance programs. REFISSA DOES NOT ELIMINATE WRINKLES, REPAIR SUN-DAMAGED SKIN, REVERSE PHOTOAGING, or RESTORE A MORE YOUTHFUL or YOUNGER DERMAL HISTOLOGIC PATTERN.

Indications and usage

The gradual bleaching of hyperpigmented skin conditions such as chloasma, melasma, freckles, senile lentigines, and other unwanted areas of melanin hyperpigmentation. Specially formulated for blending purposes as part of the Obagi Nu-Derm System.

Dosage and administration

Use daily, in the evening. Squeeze a small amount (approximately 1-2 pea-sized drops) onto your hand. Apply evenly to the entire face, or as directed by your skin care 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 darkening from reoccurring.

Warnings

Avoid contact with eyes, nose, mouth, or lips. In case of accidental contact, patient should rinse eyes thoroughly with water and contact physician. Sunscreen use is an essential aspect of hydroquinone therapy because even minimal sunlight exposure sustains melanocytic activity.

Contains sodium metabisulfite, a sulfite that may cause allergic-type reactions including anaphylactic symptoms and life-threatening or less severe asthmatic episodes in certain susceptible people. The overall prevalence of sulfite sensitivity in the general population is unknown and probably low. Sulfite sensitivity is seen more frequently in asthmatic than in nonasthmatic people.

Each gram of Obagi Nu-Derm Blender contains

Active ingredient

Hydroquinone USP, 4% (40 mg/g)

Inactive ingredients

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, disodium EDTA, propylparaben, saponins, BHT

See enclosed Package Insert for full prescribing information.

Rx ONLY. FOR EXTERNAL USE ONLY.

Sun Shield Matte Broad Spectrum SPF 50 Net wt. 3 oz. (85 g.)

This sunscreen combines UVB absorption and UVA protection in an elegant, matte finish that is non-comedogenic, allergy tested, and dermatologist tested. Sheer, PABA free, and fragrance free for all skin types.

Drug Facts

Active ingredients	***Purpose***
Octinoxate 7.5%	Sunscreen
Zinc Oxide 10.5%	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 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

1,2-hexanediol, caprylyl glycol, ceteareth-20, cetearyl alcohol, chlorphenesin, citric acid, cyclopentasiloxane, dimethicone crosspolymer-3, disodium EDTA, hydrogenated palm glycerides, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, methylisothiazolinone, PEG-10 dimethicone, PEG-40 stearate, pentylene glycol, phenyl trimethicone, polysilicone-11, polysorbate 60, potassium sorbate, sodium benzoate, sodium dihydroxycetyl phosphate, squalane, stearyl alcohol, tetrahexyldecyl ascorbate, tocopheryl acetate, tropolone, ubiquinone, water, xanthan gum

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

Travel Bag and Patient Instruction Guide

Store at controlled room temperature 15°C–25°C (59°F–77°F).

PRINCIPAL DISPLAY PANEL - Kit Carton

NDC# 62032-514-00

OBAGI®
MEDICAL

NU-DERM® SYSTEM

NORMAL DRY

Skin Transformation Kit

OBAGI MEDICAL

NU-DERM® SYSTEM

OBAGI® MEDICAL

NU-DERM® SYSTEM

NORMAL DRY

Skin Transformation Kit

OBAGI® MEDICAL

NU-DERM® SYSTEM

OBAGI® MEDICAL

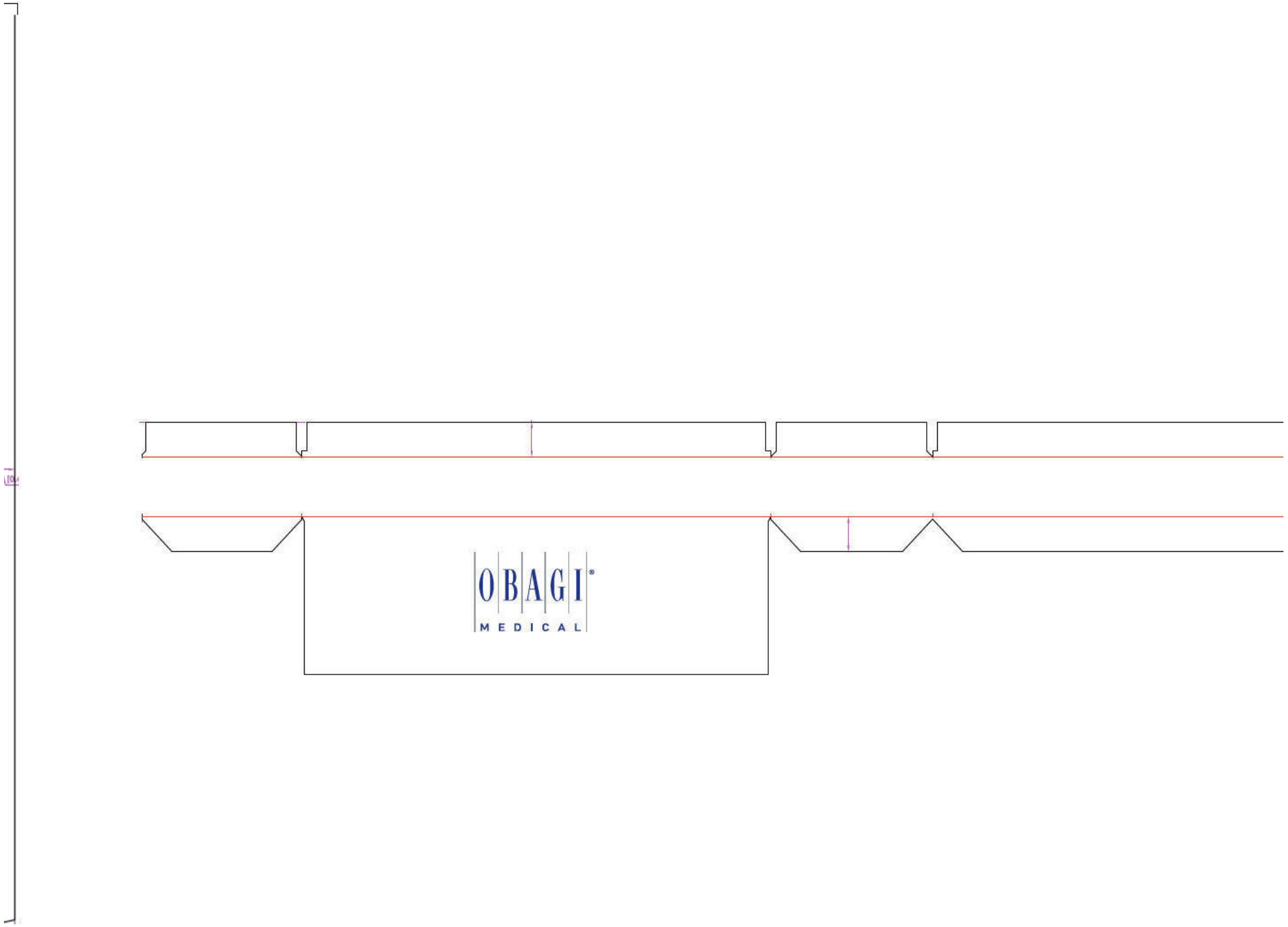
NU-DERM® SYSTEM

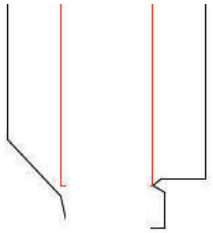
The #1 prescription-strength, physician-dispensed skincare system that transforms cellular functions at all layers of the skin, helping to make skin look and act younger and healthier.

NORMAL DRY

Skin Transformation Kit

FPO
Barcode Area
6201251400


OBAGI®
MEDICAL



NU-DERM SYSTEM NORMAL-DRY SKIN TRANSFORMATION

hydroquinone, octinoxate,and zinc oxide kit

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:62032-514

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62032-514-00	1 in 1 CARTON		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 BOTTLE, PLASTIC	57 g
Part 2	1 BOTTLE, PLASTIC	57 g
Part 3	1 TUBE	85 g
Part 4	1 BOTTLE, PLASTIC	198 mL
Part 5	1 BOTTLE, PLASTIC	198 mL
Part 6	1 BOTTLE, PLASTIC	57 g
Part 7	1 BOTTLE, PLASTIC	48 g

Part 1 of 7

NU-DERM BLENDER SKIN LIGHTENER AND BLENDING

hydroquinone cream

Product Information	
Item Code (Source)	NDC:62032-100
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:62032-100-36	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 2 of 7

NU-DERM CLEAR SKIN BLEACHING AND CORRECTOR

hydroquinone cream

Product Information	
Item Code (Source)	NDC:62032-101
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:62032-101-36	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

NU-DERM SUN SHIELD BROAD SPECTRUM SPF 50 MATTE SUNSCREEN

octinoxate and zinc oxide lotion

Product Information

Item Code (Source)	NDC:62032-104
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 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	105 mg in 1 g

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: 2KR89I4H1Y)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
PEG-40 STEARATE (UNII: ECU18C66Q7)	
SODIUM DIHYDROXYCETYL PHOSPHATE (UNII: YWI33EV595)	
HYDROGENATED PALM GLYCERIDES (UNII: YCZ8EM144Q)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
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)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
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:62032-104-90	85 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	11/07/2012	

Part 4 of 7

NU-DERM 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	**APRICOT KERNEL OIL** (UNII: 54JB35T06A)	
INGR	**OLEYL LACTATE** (UNII: B3AWW0N3GM)	
INGR	**CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)** (UNII: 809Y72KV36)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		198 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

NU-DERM 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		198 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

NU-DERM 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		57 g in 1 BOTTLE, PLASTIC		

Marketing Information

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

Part 7 of 7

NU-DERM HYDRATE FACIAL MOISTURIZER

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	**MEDIUM-CHAIN TRIGLYCERIDES** (UNII: C9H2L21V7U)	
INGR	**CAESALPINIA SPINOSA RESIN** (UNII: WL3883U2PO)	
INGR	**SHEA BUTTER** (UNII: K49155WL9Y)	
INGR	**GLYCERYL MONOSTEARATE** (UNII: 230OU9XXE4)	
INGR	**CYCLOMETHICONE 5** (UNII: 0THT5PCI0R)	
INGR	**CETYL ALCOHOL** (UNII: 936JST6JCN)	
INGR	**SACCHARIDE ISOMERATE** (UNII: W8K377W98I)	
INGR	**DIMETHICONE** (UNII: 92RU3N3Y1O)	
INGR	**LAURETH-12** (UNII: OAH19558U1)	
INGR	**PHENOXYETHANOL** (UNII: HIE492ZZ3T)	

INGR	**ETHYLHEXYLGLYCERIN** (UNII: 147D247K3P)	
INGR	**STEARIC ACID** (UNII: 4ELV7Z65AP)	
INGR	**AVOCADO OIL** (UNII: 6VNO72PFC1)	
INGR	**SOY STEROL** (UNII: PL360EPO9J)	
INGR	**CAPRYLYL GLYCOL** (UNII: 00YIU5438U)	
INGR	**LEVOMENOL** (UNII: 24WE03BX2T)	
INGR	**HEXYLENE GLYCOL** (UNII: KEH0A3F75J)	
INGR	**PANTHENOL** (UNII: WV9CM0O67Z)	
INGR	**MANGIFERA INDICA SEED BUTTER** (UNII: 4OXD9M35X2)	
INGR	**SODIUM STEAROYL GLUTAMATE** (UNII: 65A9F4P024)	
INGR	**CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)** (UNII: 59TL3WG5CO)	
INGR	**ALLANTOIN** (UNII: 344S277G0Z)	
INGR	**TETRAHYDRODIFERULOYLMETHANE** (UNII: 00U0645U03)	
INGR	**TOCOPHEROL** (UNII: R0ZB2556P8)	
INGR	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		48 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Cosmetic		11/07/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		11/07/2012	

Labeler - OMP, INC. (790553353)

Establishment

Name	Address	ID/FEI	Business Operations
MILBAR LABORATORIES		195556790	MANUFACTURE(62032-514)

Establishment

Name	Address	ID/FEI	Business Operations
PURETEK CORPORATION		785961046	MANUFACTURE(62032-514) , LABEL(62032-514) , PACK(62032-514)

Establishment

Name	Address	ID/FEI	Business Operations
Ei INC.		105803274	MANUFACTURE(62032-514) , LABEL(62032-514) , PACK(62032-514) , ANALYSIS(62032-514)

Establishment

Name	Address	ID/FEI	Business Operations
G.S. COSMECEUTICAL USA, INC.		017014734	MANUFACTURE(62032-514)

Establishment

Name	Address	ID/FEI	Business Operations
Bay Cities Container Corporation		066229618	RELABEL(62032-514) , REPACK(62032-514)

Revised: 12/2012 OMP, INC.