OBAGI NU-DERM SYSTEM NORMAL-DRY SKIN TRANSFORMATION-
hydroquinone, titanium dioxide, and zinc oxide
Obagi Cosmeceuticals LLC

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 Nu-Derm® System Normal-Dry Skin Transformation Kit

Drug Facts

Active ingredients	***Purpose***
Titanium Dioxide 6.2%	Sunscreen
Zinc Oxide 15.6%	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
- **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
- children under 6 months: Ask a doctor

Other information

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

Inactive ingredients

aqua/water/eau, caprylic/capric triglyceride, dicaprylyl carbonate, pentylene glycol, steareth-2, glycerin, silica, polyhydroxystearic acid, steareth-21, cetearyl alcohol, propanediol, alumina, stearic acid, sodium stearoyl glutamate, caprylyl glycol, triethoxycaprylylsilane, cetearyl glucoslde, xanthan gum, dipotassium glycyrrhizate,trisodium ethylenediamine disuccinate, caprylhydroxamic acid, tropaeolum majus flower/leaf/stem extract.

Questions or comments?

1.800.636.7546 Monday–Friday 9 a.m.–4 p.m. PST

Distributed by Obagi Cosmeceuticals LLC,
Long Beach, CA 90806.

PRINCIPAL DISPLAY PANEL - Kit Carton

OBAGI®
MEDICAL

OBAGI NU-DERM® SYSTEM

Addresses signs of skin aging, gently exfoliates to promote cell turnover and suppresses melanocyte activity to reduce hyperpigmentaion.

NORMAL DRY

Skin Transformation Kit

OBAGI MEDICAL

OBAGI NU-DERM® SYSTEM

Addresses signs of skin aging, gently exfoliates to promote cell turnover and suppresses melanocyte activity to reduce hyperpigmentaion.

NORMAL **DRY**

Skin Transformation Kit

Sun Shield Mineral Broad Spectrum SPF 50 Sunscreen Lotion | Net wt. 3 oz. (85 g)

Drug Facts

Active ingredients	Purpose
Titanium Dioxide 6.2%	Sunscreen
Zinc Oxide 15.6%	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
- **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
- children under 6 months: Ask a doctor

Drug Facts *(continued)*

Other information

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

Inactive ingredients

aqua/water/eau, caprylic/capric triglyceride, dicaprylyl carbonate, pentylene glycol, steareth-2, glycerin, silica, polyhydroxystearic acid, steareth-21, cetearyl alcohol, propanediol, alumina, stearic acid, sodium stearoyl glutamate, caprylyl glycol, triethoxycaprylylsilane, cetearyl glucoside, xanthan gum, dipotassium glycyrrhizate, trisodium ethylenediamine disuccinate, caprylhydroxamic acid, tropaeolum majus flower/leaf/stem extract.

Questions or comments?

1.800.636.7546 Monday–Friday 9 a.m.–4 p.m. PST

Gentle Cleanser | AM+PM

A mild daily facial cleanser that gently removes impurities, oil, and buildup for cleaner, fresher-looking skin. Helps

to soothe delicate and dry skin.

Toner | AM+PM
A toner formulated with Witch Hazel and Aloe Vera, effectively removes impurities and hydrates skin, to reveal a fresher-looking complexion. Primes skin for the next step in your skincare routine.

Exfoderm® (Skin Smoothing Lotion) | AM
A lightweight lotion that exfoliates the top layer of skin, removing dull, old skin while revealing new skin 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.

Clear (Skin Bleaching and Corrector Cream) | AM+PM
Hydroquinone USP, 4% Rx Only, NDC 62032-101-36
Dark spots may appear on the surface of your skin, but they actually start deep within the skin's layers. This effective formula absorbs into the layers of your skin to deliver prescription-strength hydroquinone, helping to correct dark spots for a healthier-looking, more even complexion.

Blender® (Skin Lightener and Blending Cream) | PM
Hydroquinone USP, 4% Rx Only, NDC 62032-100-36
A unique formula containing prescription-strength hydroquinone for the gradual lightening of sun spots or age spots and other types of hyperpigmentation (discoloration). Specially formulated to optimize the delivery of product ingredients in the Obagi Nu-Derm® System, this skin lightener helps reduce signs of aging by correcting uneven skin tone.

Obagi Hydrate® (Facial Moisturizer) | AM+PM
A lightweight facial moisturizer formulated with Tara Seed Extract, clinically proven to improve skin's moisture, and naturally derived ingredients, such as Shea Butter and Mango Seed Butter, to further combat dryness. For long-lasting hydration and smoother-looking skin. Suitable for all skin types.

Sun Shield® Mineral Broad Spectrum SPF 50 Sunscreen Lotion | AM
An elegant daily sunscreen that protects from UVA and UVB rays with a 100% mineral blend of zinc oxide and titanium dioxide, and provides a lightweight satin finish. This high protection physical sunscreen is also formulated with Nasturtium flower extract to help defend against blue and visible light. Formulated for all skin types, including sensitive skin.

Gentle Cleanser | 6.7 fl. oz. (200 mL)
Directions: Apply to damp face and neck. Rinse completely.
Ingredients: aqua/water/eau, cocamidopropyl betaine, glycerin, sodium lauroyl oat amino acids, aloe barbadensis leaf juice, sodium laureth sulfate, sodium chloride, glycereth-7, tocopherol, panthenol, prunus armeniaca (apricot) kernel oil, borago officinalis extract, salvia officinalis (sage) leaf extract, saponins, oleyl lactate, acrylates/c10-30 alkyl acrylate crosspolymer, ethoxydiglycol, triethanolamine, phenoxyethanol, ethylhexylglycerin, fragrance (parfum), sodium benzoate, tetrasodium EDTA, citric acid, potassium sorbate, methylchloroisothiazolinone, methylisothiazolinone, yellow 5 (CI 19140).

12 M

Toner | 6.7 fl. oz. (200 mL)
Directions: After cleansing, apply toner onto face and neck with a cotton ball.
Ingredients: aqua/water/eau, hamamelis virginiana (witch hazel) water, aloe barbadensis leaf juice, tocopherol, borago officinalis extract, salvia officinalis (sage) leaf extract, calendula officinalis flower extract, allantoin, panthenol, glycerin, saponins, sodium PCA, phenoxyethanol, potassium alum, polysorbate 80, ethylhexylglycerin, fragrance (parfum), citric acid, benzoic acid, sodium benzoate, potassium sorbate, blue 1 (CI 42090), alpha-isomethyl-ionone, amyl cinnamal, benzyl salicylate, hydroxycitronellal, limonene, linalool, linalyl acetate.

12 M

Exfoderm® (Skin Smoothing Lotion) | Net wt. 2 oz. (57 g)
Directions: Apply topically in a thin layer to the skin once a day, in the morning.
Ingredients: aqua/water/eau, ethoxydiglycol, glycerin, phytic acid, cetearyl alcohol, glyceryl stearate, PEG-100 stearate, canola oil, isohexadecane, magnesium aluminum silicate, potassium cetyl phosphate, sodium hydroxide, bis-diglyceryl polyacyladipate-2, cetyl alcohol, dimethicone, xanthan gum, phenoxyethanol, glycereth-7, polysorbate 60, steareth-20, ethylhexylglycerin, tocopheryl acetate, PEG-150 distearate, saponins.

12 M

PRECAUTIONS For external use only. Avoid getting into eyes. If product gets into eyes, rinse with water to remove. **Keep out of reach of children.**
STORAGE Store at room temperature: 15°-25°C (59°-77°F). Keep out of direct sunlight.
QUESTIONS OR COMMENTS? 1.800.636.7546

Obagi Hydrate® (Facial Moisturizer) | Net wt. 1.7 oz. (48 g)
Directions: Apply to face in the morning and evening or as needed.
Ingredients: aqua/water/eau, glycerin, caprylic/capric triglyceride, butyrospermum parkii (shea) butter, cyclopentasiloxane, glyceryl stearate, cetyl alcohol, dimethicone, saccharide isomerate, polysilicone-11, glycine soja (soybean) sterols, persea gratissima (avocado) oil, stearic acid, tetrahydrodiferuloylmethane, mangifera indica (mango) seed butter, hydrolyzed caesalpinia spinosa gum, caesalpinia spinosa gum, hydrolyzed soybean fiber, panthenol, tocopherol, allantoin, bisabolol, sodium stearoyl glutamate, laureth-12, phenoxyethanol, ethylhexylglycerin, hexylene glycol, caprylyl glycol, carbomer, sodium hydroxide.

12 M

Clear (Skin Bleaching and Corrector Cream) | Net wt. 2 oz. (57 g)
Hydroquinone USP, 4% Rx only NDC 62032-101-36
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-size 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 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, and lips. In case of accidental contact, patient should rinse thoroughly with water and contact a 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: aqua/water/eau, 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 prescribing information.
Rx ONLY. FOR EXTERNAL USE ONLY.

Blender® (Skin Lightener and Blending Cream) | Net wt. 2 oz. (57 g)
Hydroquinone USP, 4% Rx only NDC 62032-100-36
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 evening. Squeeze a small amount (approximately 1-2 pea-size 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 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, and lips. In case of accidental contact, patient should rinse thoroughly with water and contact a 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: aqua/water/eau, 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 prescribing information.
Rx ONLY. FOR EXTERNAL USE ONLY.

Distributed by Obagi Cosmeceuticals LLC,
Long Beach, CA 90806. www.obagi.com

Made in USA with US and imported components.
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OBAGI NU-DERM SYSTEM NORMAL-DRY SKIN TRANSFORMATION
hydroquinone, titanium dioxide, and zinc oxide kit

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:62032-918

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62032-918-01	1 in 1 BAG	06/25/2025	

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

Part 1 of 7

NU-DERM GENTLE CLEANSER

cleansing (cold creams, cleansing lotions, liquids, and pads) [skin care preparations (creams, lotions, powder, and sprays)] 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	**CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)** (UNII: 59TL3WG5CO)	
INGR	**SODIUM LAUROYL OAT AMINO ACIDS** (UNII: FSW2K9B9N5)	
INGR	**COCAMIDOPROPYL BETAINE** (UNII: 5OCF3O11KX)	
INGR	**SODIUM LAURETH-3 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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		198 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

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

Part 2 of 7

NU-DERM TONER

cleansing (cold creams, cleansing lotions, liquids, and pads) [skin care preparations (creams, lotions, powder, and sprays)] liquid

Product Information

Route of Administration	TOPICAL

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
INGR	**WATER** (UNII: 059QF0KO0R)	
INGR	**GLYCERIN** (UNII: PDC6A3C0OX)	
INGR	**HAMAMELIS VIRGINIANA TOP WATER** (UNII: NT00Y05A2V)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		198 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

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

Part 3 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, UNSPECIFIED FORM (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; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		01/01/1988	

Part 4 of 7

NU-DERM EXFODERM SKIN SMOOTHING

face and neck (excluding shaving preparations), leave-on [skin care preparations (creams, lotions, powder, and sprays)] 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 MONOSTEARATE** (UNII: YD01N1999R)	
INGR	**DIMETHICONE, UNSPECIFIED** (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)	
INGR	**BIS-DIGLYCERYL POLYACYLADIPATE-2** (UNII: 6L246LAM9T)	
INGR	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	

Packaging

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

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 HYDRATE FACIAL MOISTURIZER

moisturizing [skin care preparations (creams, lotions, powder, and sprays)]

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	**SODIUM HYDROXIDE** (UNII: 55X04QC32I)	
INGR	**TARA SPINOSA RESIN** (UNII: WL3883U2PO)	
INGR	**SHEA BUTTER** (UNII: K49155WL9Y)	
INGR	**DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE)** (UNII: 9E4CO0W6C5)	
INGR	**CYCLOMETHICONE 5** (UNII: 0THT5PCI0R)	
INGR	**CETYL ALCOHOL** (UNII: 936JST6JCN)	
INGR	**SACCHARIDE ISOMERATE** (UNII: W8K377W98I)	
INGR	**DIMETHICONE, UNSPECIFIED** (UNII: 92RU3N3Y1O)	
INGR	**TOCOPHEROL** (UNII: R0ZB2556P8)	
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	**TETRAHYDRODIFERULOYLMETHANE** (UNII: 00U0645U03)	
INGR	**PANTHENOL** (UNII: WV9CM0O67Z)	
INGR	**MANGIFERA INDICA SEED BUTTER** (UNII: 4OXD9M35X2)	
INGR	**SODIUM STEAROYL GLUTAMATE** (UNII: 65A9F4P024)	
INGR	**CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE** (UNII: 0A5MM307FC)	
INGR	**ALLANTOIN** (UNII: 344S277G0Z)	
INGR	**GLYCERYL MONOSTEARATE** (UNII: 230OU9XXE4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		48 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

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

Part 6 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, UNSPECIFIED FORM (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; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		01/01/1988	

Part 7 of 7

SUN SHIELD MINERAL BROAD SPECTRUM SPF 50 SUNSCREEN

titanium dioxide and zinc oxide lotion

Product Information

Item Code (Source)	NDC:62032-811
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	6.2 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	15.6 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALUMINUM OXIDE (UNII: LMI26O6933)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETEARYL GLUCOSIDE (UNII: 09FUA47KNA)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
GLYCYRRHIZINATE DIPOTASSIUM (UNII: CA2Y0FE3FX)	
GLYCERIN (UNII: PDC6A3C0OX)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
PROPANEDIOL (UNII: 5965N8W85T)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
TROPAEOLUM MAJUS FLOWERING TOP (UNII: RGT30824HY)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

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

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62032-811-09	1 in 1 CARTON		
1		85 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M020	12/02/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		06/25/2025	

Labeler - Obagi Cosmeceuticals LLC (790553353)

Establishment

Name	Address	ID/FEI	Business Operations
Bay Cities Container Corporation		118417470	PACK(62032-918) , LABEL(62032-918)

Establishment

Name	Address	ID/FEI	Business Operations
Swiss American CDMO, LLC		080170933	MANUFACTURE(62032-918)

Establishment

Name	Address	ID/FEI	Business Operations
G. S. Cosmeceutical USA, Inc.		017014734	MANUFACTURE(62032-918)

Establishment

Name	Address	ID/FEI	Business Operations
PURETEK CORPORATION		785961046	MANUFACTURE(62032-918)