

**OBAGI-C C-RX SYSTEM NORMAL-DRY SKIN INTERVENTION- hydroquinone, titanium dioxide, 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.*

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**Obagi-C® C-Rx System Normal-Dry Skin Intervention**

***Drug Facts***

| <b><i>Active ingredients</i></b> | <b><i>Purpose</i></b> |
|----------------------------------|-----------------------|
| 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 stearyl 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

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

### **PRINCIPAL DISPLAY PANEL - Kit Carton**

OBAGI®  
MEDICAL

OBAGI-C® Rx SYSTEM

Complete skin care system specifically formulated for normal to dry skin with hydroquinone to reduce hyperpigmentation and other essential ingredients to help address the signs of sun exposure caused by photoaging.

NORMAL DRY

Skin Intervention Kit

OBAGI  
MEDICAL

## OBAGI-C® Rx SYSTEM

Complete skin care system specifically formulated for normal to dry skin with hydroquinone to reduce hyperpigmentation and other essential ingredients to help address the signs of sun exposure caused by photoaging.



NORMAL DRY  
Skin Intervention Kit

Sun Shield Mineral Broad Spectrum SPF 50 Sunscreen Lotion | Netwt. 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

#### Other information

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

### Drug Facts (continued)

#### 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, caprylylhydroxamic acid, tropaeolum majus flower/leaf/stem extract.

#### Questions or comments?

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

#### C-Cleansing Gel | 6 fl. oz. (180 mL)

Directions: Use twice daily, morning and evening. Apply to damp face and neck with moistened fingertips, rub in circular motion and rinse completely with lukewarm water.

Ingredients: aqua/water/EAU, sodium laureth sulfate, sodium lauroyl oat amino acids, cocamidopropyl betaine, sodium chloride, aloe barbadensis leaf juice, phenoxyethanol, fragrance [parfum], ethylhexylglycerin, glycerin, saponins, xanthan gum, borago officinalis extract, medicago sativa (alfalfa) extract, ascorbic acid, chamomilla recutita (matricaria) flower extract, red 33 (CI 17200), yellow 5 (CI 19140).



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#### C-Clarifying Serum Normal to Dry (Skin Lightening Serum)

1 fl. oz. (30 mL)

NDC 62032-106-10 Hydroquinone USP, 4% Rx Only

**Dosage and Administration:** A thin application should be applied once or 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 or protective clothing to cover bleached skin when using and after using this product in

#### C-Therapy Night Cream (Skin Lightener) | Netwt. 2 oz. (57 g)

NDC 62032-105-36 Hydroquinone USP, 4% Rx Only

**Dosage and Administration:** A thin application should be applied once or 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 or protective clothing to cover bleached skin when using and after using this product in order to prevent darkening from reoccurring.

order to prevent darkening from reoccurring.

**Indications and Usage:** The gradual bleaching of hyperpigmented skin conditions such as chloasma, melasma, freckles, senile lentiginos, and other unwanted areas of melanin hyperpigmentation.

**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.

■ 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 product.

■ Test for skin sensitivity before using by applying a small amount to an unbroken patch of skin and check within 24 hours. Minor redness is not a contraindication, but where there is itching or vesicle formation or excessive inflammatory response, the product should be discontinued and a physician consulted. Close patient supervision is recommended.

Each gram of Obagi-C Rx C-Clarifying Serum contains:

Active Ingredient: Hydroquinone, USP 4% (40 mg/g)

Inactive Ingredients: propylene glycol, aqua/water/eau, ascorbic acid, propylene carbonate, sodium lauryl sulfate.

See enclosed Package Insert for full prescribing information.

Rx Only. FOR EXTERNAL USE ONLY.

**C-Exfoliating Day Lotion | 2 fl. oz. (57 mL)**

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

**Sunburn Alert:** This product contains an alpha hydroxy acid (AHA) that may increase your skin's sensitivity to the sun and particularly the possibility of sunburn. Use a sunscreen, wear protective clothing, and limit sun exposure while using this product and for a week afterwards.

**Ingredients:** aqua/water/eau, ethylhexyl palmitate, ethylhexyl stearate, glycolic acid, caprylic/capric triglyceride, isopropyl palmitate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, squalane, PEG-8 dimethicone, glycerin, glyceryl stearate, PEG-100 stearate, sodium hydroxide, ascorbyl glucoside, tocopheryl acetate, sodium hyaluronate, PEG-8 ricinoleate, arginine, cetearyl alcohol, dimethicone, polysorbate 60, steareth-2, ceteareth-20, bisabolol, tetrasodium EDTA, phenoxyethanol, ethylhexyl glycerin, fragrance (parfum).



**Indications and Usage:** The gradual bleaching of hyperpigmented skin conditions such as chloasma, melasma, freckles, senile lentiginos, and other unwanted areas of melanin hyperpigmentation.

**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.

■ 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 product.

■ Test for skin sensitivity before using by applying a small amount to an unbroken patch of skin and check within 24 hours. Minor redness is not a contraindication, but where there is itching or vesicle formation or excessive inflammatory response, the product should be discontinued and a physician consulted. Close patient supervision is recommended.

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-C Rx C-Therapy Night Cream contains:

Active Ingredient: Hydroquinone, USP 4% (40 mg/g)

Inactive Ingredients: aqua/water/eau, glycerin, cetyl alcohol, PPG-2 myristylether 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.

**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-C® Rx SYSTEM

Complete skin care system specifically formulated for normal to dry skin with hydroquinone to reduce hyperpigmentation and other essential ingredients to help address the signs of sun exposure caused by photoaging.

### C-Cleansing Gel | AM+PM

Helps clarify and prepare your skin for the system's product ingredients, while also removing makeup and oil.

### C-Clarifying Serum Normal to Dry (Skin Lightening Serum) | AM

NDC 62032-106-10, Hydroquinone USP, 4% Rx Only

Antioxidant Serum containing Vitamin C and prescription-strength hydroquinone. This formulation for normal to dry skin corrects dark spots for a lighter, brighter complexion.

### C-Exfoliating Day Lotion | AM

A smooth lightweight moisturizer that hydrates and gently exfoliates the skin to reveal a brighter, healthier-looking complexion.

### C-Therapy Night Cream (Skin Lightener) | PM

NDC 62032-105-36 Hydroquinone USP, 4% Rx Only

A rich moisturizer that works while you sleep to renew and rejuvenate your skin. The C-Therapy Night Cream is uniquely formulated with prescription-strength hydroquinone to gradually diminish dark spots and provide Vitamins C and E during the skin's nightly renewal process.

### 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.

## OBAGI-C C-RX SYSTEM NORMAL-DRY SKIN INTERVENTION

hydroquinone, titanium dioxide, zinc oxide kit

### Product Information

#### Product Type

HUMAN OTC DRUG

#### Item Code (Source)

NDC:62032-922

### Packaging

| # | Item Code        | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------|----------------------|--------------------|
| 1 | NDC:62032-922-01 | 1 in 1 BAG          | 06/25/2025           |                    |

### Quantity of Parts

| Part # | Package Quantity  | Total Product Quantity |
|--------|-------------------|------------------------|
| Part 1 | 1 BOTTLE, GLASS   | 30 mL                  |
| Part 2 | 1 BOTTLE, PLASTIC | 57 g                   |
| Part 3 | 1 TUBE            | 85 g                   |
| Part 4 | 1 BOTTLE, PLASTIC | 177 mL                 |
| Part 5 | 1 BOTTLE, PLASTIC | 57 g                   |

### Part 1 of 5

### OBAGI C RX SYSTEM C CLARIFYING SERUM

hydroquinone solution

### Product Information

|                         |               |
|-------------------------|---------------|
| Item Code (Source)      | NDC:62032-106 |
| Route of Administration | TOPICAL       |

### Active Ingredient/Active Moiety

| Ingredient Name                                                         | Basis of Strength | Strength      |
|-------------------------------------------------------------------------|-------------------|---------------|
| <b>HYDROQUINONE</b> (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE) | HYDROQUINONE      | 40 mg in 1 mL |

### Inactive Ingredients

| Ingredient Name                                 | Strength |
|-------------------------------------------------|----------|
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)      |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                 |          |
| <b>ASCORBIC ACID</b> (UNII: PQ6CK8PD0R)         |          |
| <b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J) |          |
| <b>PROPYLENE CARBONATE</b> (UNII: 8D08K3S51E)   |          |

### Packaging

| # | Item Code        | Package Description                                         | Marketing Start Date | Marketing End Date |
|---|------------------|-------------------------------------------------------------|----------------------|--------------------|
| 1 | NDC:62032-106-10 | 1 in 1 CARTON                                               |                      |                    |
| 1 |                  | 30 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product |                      |                    |

## Marketing Information

| Marketing Category       | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------------|------------------------------------------|----------------------|--------------------|
| UNAPPROVED DRUG<br>OTHER |                                          | 01/01/2004           |                    |

## Part 2 of 5

### OBAGI-C RX SYSTEM C-THERAPY NIGHT SKIN LIGHTENING WITH VITAMINS C AND E

hydroquinone cream

## Product Information

|                         |               |
|-------------------------|---------------|
| Item Code (Source)      | NDC:62032-105 |
| Route of Administration | TOPICAL       |

## Active Ingredient/Active Moiety

| Ingredient Name                                                         | Basis of Strength | Strength     |
|-------------------------------------------------------------------------|-------------------|--------------|
| <b>HYDROQUINONE</b> (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE) | HYDROQUINONE      | 40 mg in 1 g |

## Inactive Ingredients

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

## Product Characteristics

|       |       |       |  |
|-------|-------|-------|--|
| Color | WHITE | Score |  |
|-------|-------|-------|--|

|          |  |              |  |
|----------|--|--------------|--|
| Shape    |  | Size         |  |
| Flavor   |  | Imprint Code |  |
| Contains |  |              |  |

| Packaging |                  |                                                              |                      |                    |
|-----------|------------------|--------------------------------------------------------------|----------------------|--------------------|
| #         | Item Code        | Package Description                                          | Marketing Start Date | Marketing End Date |
| 1         | NDC:62032-105-36 | 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<br>OTHER |                                          | 01/01/2004           |                    |

| Part 3 of 5                                                                                         |  |
|-----------------------------------------------------------------------------------------------------|--|
| <b>SUN SHIELD MINERAL BROAD SPECTRUM SPF 50 SUNSCREEN</b><br>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        |
| <b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP) | TITANIUM DIOXIDE  | 6.2 g in 100 g  |
| <b>ZINC OXIDE</b> (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)             | ZINC OXIDE        | 15.6 g in 100 g |

| Inactive Ingredients                                        |          |
|-------------------------------------------------------------|----------|
| Ingredient Name                                             | Strength |
| <b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)                    |          |
| <b>CAPRYLHYDROXAMIC ACID</b> (UNII: UPY805K99W)             |          |
| <b>MEDIUM-CHAIN TRIGLYCERIDES</b> (UNII: C9H2L21V7U)        |          |
| <b>CAPRYLYL GLYCOL</b> (UNII: 00YIU5438U)                   |          |
| <b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)               |          |
| <b>DICAPRYLYL CARBONATE</b> (UNII: 609A3V1SUA)              |          |
| <b>GLCYRRHIZINATE DIPOTASSIUM</b> (UNII: CA2Y0FE3FX)        |          |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                          |          |
| <b>PENTYLENE GLYCOL</b> (UNII: 50C1307PZG)                  |          |
| <b>POLYHYDROXYSTEARIC ACID (2300 MW)</b> (UNII: YXH47AOU0F) |          |

|                                                                 |  |
|-----------------------------------------------------------------|--|
| <b>PROPANEDIOL</b> (UNII: 5965N8W85T)                           |  |
| <b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)                       |  |
| <b>SODIUM STEAROYL GLUTAMATE</b> (UNII: 65A9F4P024)             |  |
| <b>STEARETH-2</b> (UNII: V56DFE46J5)                            |  |
| <b>STEARETH-21</b> (UNII: 53J3F32P58)                           |  |
| <b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)                          |  |
| <b>TRIETHOXYCAPRYLYLSILANE</b> (UNII: LDC331P08E)               |  |
| <b>TRISODIUM ETHYLENEDIAMINE DISUCCINATE</b> (UNII: YA22H34H9Q) |  |
| <b>TROPAEOLUM MAJUS FLOWERING TOP</b> (UNII: RGT30824HY)        |  |
| <b>WATER</b> (UNII: 059QF0KO0R)                                 |  |
| <b>XANTHAN GUM</b> (UNII: TTV12P4NEE)                           |  |

## Product Characteristics

|                 |       |                     |  |
|-----------------|-------|---------------------|--|
| <b>Color</b>    | WHITE | <b>Score</b>        |  |
| <b>Shape</b>    |       | <b>Size</b>         |  |
| <b>Flavor</b>   |       | <b>Imprint Code</b> |  |
| <b>Contains</b> |       |                     |  |

## 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           |                    |

## Part 4 of 5

### OBAGI-C C-CLEANSING WITH VITAMIN C

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

## Product Information

|                                |         |
|--------------------------------|---------|
| <b>Route of Administration</b> | TOPICAL |
|--------------------------------|---------|

## Other Ingredients

| Ingredient Kind | Ingredient Name                                          | Quantity |
|-----------------|----------------------------------------------------------|----------|
| INGR            | <b>WATER</b> (UNII: 059QF0K00R)                          |          |
| INGR            | <b>GLYCERIN</b> (UNII: PDC6A3C00X)                       |          |
| INGR            | <b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)                 |          |
| INGR            | <b>ETHYLPARABEN</b> (UNII: 14255EXE39)                   |          |
| INGR            | <b>ISOBUTYLPARABEN</b> (UNII: 0QQJ25X58G)                |          |
| INGR            | <b>BUTYLPARABEN</b> (UNII: 3QPI1U3FV8)                   |          |
| INGR            | <b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)                  |          |
| INGR            | <b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)                  |          |
| INGR            | <b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)          |          |
| INGR            | <b>ASCORBIC ACID</b> (UNII: PQ6CK8PD0R)                  |          |
| INGR            | <b>SODIUM LAUROYL OAT AMINO ACIDS</b> (UNII: FSW2K9B9N5) |          |
| INGR            | <b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)         |          |
| INGR            | <b>SODIUM LAURETH-3 SULFATE</b> (UNII: BPV390UAP0)       |          |
| INGR            | <b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)                 |          |
| INGR            | <b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)                |          |
| INGR            | <b>MEDICAGO SATIVA WHOLE</b> (UNII: DJO934BRBD)          |          |
| INGR            | <b>CHAMOMILE</b> (UNII: FGL3685T2X)                      |          |
| INGR            | <b>XANTHAN GUM</b> (UNII: TTV12P4NEE)                    |          |
| INGR            | <b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)             |          |
| INGR            | <b>BORAGE</b> (UNII: PB618V0K2W)                         |          |

Packaging

| # | Item Code | Package Description                                            | Marketing Start Date | Marketing End Date |
|---|-----------|----------------------------------------------------------------|----------------------|--------------------|
| 1 |           | 177 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           |                                          | 01/01/2004           |                    |

Part 5 of 5

OBAGI-C C-EXFOLIATING DAY WITH VITAMIN C

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            | <b>WATER</b> (UNII: 059QF0KO0R)                                                                                  |          |
| INGR            | <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                                                                               |          |
| INGR            | <b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)                                                                         |          |
| INGR            | <b>ETHYLPARABEN</b> (UNII: 14255EXE39)                                                                           |          |
| INGR            | <b>ISOBUTYLPARABEN</b> (UNII: 0QQJ25X58G)                                                                        |          |
| INGR            | <b>BUTYLPARABEN</b> (UNII: 3QPI1U3FV8)                                                                           |          |
| INGR            | <b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)                                                                          |          |
| INGR            | <b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)                                                                          |          |
| INGR            | <b>POLYSORBATE 60</b> (UNII: CAL22UVI4M)                                                                         |          |
| INGR            | <b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)                                                                  |          |
| INGR            | <b>PEG-100 MONOSTEARATE</b> (UNII: YD01N1999R)                                                                   |          |
| INGR            | <b>ETHYLHEXYL PALMITATE</b> (UNII: 2865993309)                                                                   |          |
| INGR            | <b>SQUALANE</b> (UNII: GW89575KF9)                                                                               |          |
| INGR            | <b>HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%)</b> (UNII: 86FQE96TZ4) |          |
| INGR            | <b>PEG-8 DIMETHICONE (600 CST)</b> (UNII: GIA7T764OD)                                                            |          |
| INGR            | <b>PEG-8 RICINOLEATE</b> (UNII: DM36F4D2OU)                                                                      |          |
| INGR            | <b>GLYCOLIC ACID</b> (UNII: 0WT12SX38S)                                                                          |          |
| INGR            | <b>ARGININE</b> (UNII: 94ZLA3W45F)                                                                               |          |
| INGR            | <b>ETHYLHEXYL STEARATE</b> (UNII: EG3PA2K3K5)                                                                    |          |
| INGR            | <b>MEDIUM-CHAIN TRIGLYCERIDES</b> (UNII: C9H2L21V7U)                                                             |          |
| INGR            | <b>ISOPROPYL PALMITATE</b> (UNII: 8CRQ2TH63M)                                                                    |          |
| INGR            | <b>HYALURONATE SODIUM</b> (UNII: YSE9PPT4TH)                                                                     |          |
| INGR            | <b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)                                                                       |          |
| INGR            | <b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)                                                                    |          |
| INGR            | <b>POLYOXYL 20 CETOSTEARYL ETHER</b> (UNII: YRC528SWJY)                                                          |          |
| INGR            | <b>DIMETHICONE, UNSPECIFIED</b> (UNII: 92RU3N3Y1O)                                                               |          |
| INGR            | <b>STEARETH-2</b> (UNII: V56DFE46J5)                                                                             |          |
| INGR            | <b>ASCORBYL GLUCOSIDE</b> (UNII: 2V52R0NHXW)                                                                     |          |
| INGR            | <b>LEVOMENOL</b> (UNII: 24WE03BX2T)                                                                              |          |
| INGR            | <b>EDETATE SODIUM</b> (UNII: MP1J8420LU)                                                                         |          |
| INGR            | <b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)                                                             |          |

### 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           |                                          | 01/02/2004           |                    |

## Marketing Information

| Marketing Category       | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------------|------------------------------------------|----------------------|--------------------|
| UNAPPROVED DRUG<br>OTHER |                                          | 12/02/2019           |                    |

**Labeler** - Obagi Cosmeceuticals LLC (790553353)

## Establishment

| Name                             | Address | ID/FEI    | Business Operations                |
|----------------------------------|---------|-----------|------------------------------------|
| Bay Cities Container Corporation |         | 118417470 | PACK(62032-922) , LABEL(62032-922) |

## Establishment

| Name                          | Address | ID/FEI    | Business Operations    |
|-------------------------------|---------|-----------|------------------------|
| G. S. Cosmeceutical USA, Inc. |         | 017014734 | MANUFACTURE(62032-922) |

## Establishment

| Name                     | Address | ID/FEI    | Business Operations    |
|--------------------------|---------|-----------|------------------------|
| Swiss American CDMO, LLC |         | 080170933 | MANUFACTURE(62032-922) |

## Establishment

| Name                                           | Address | ID/FEI    | Business Operations    |
|------------------------------------------------|---------|-----------|------------------------|
| Universal Packaging Systems, Inc. (DBA Paklab) |         | 177711082 | MANUFACTURE(62032-922) |

## Establishment

| Name                | Address | ID/FEI    | Business Operations    |
|---------------------|---------|-----------|------------------------|
| PURETEK CORPORATION |         | 785961046 | MANUFACTURE(62032-922) |

Revised: 11/2025

Obagi Cosmeceuticals LLC