# OPTIMAX PROFESSIONAL WHITENING ANTICAVITY FLUORIDE- sodium monofluorophosphate WHITE GLO USA INC

-----

# **Optimax Professional Whitening Anticavity Fluoride Toothpaste**

## **Drug Facts**

## Active ingredient

Sodium Monoflurorophosphate 0.76% (0.1% W/V Fluoride ion).

## **Purpose**

Anticavity toothpaste

#### Use

helps protect against cavities

## Warnings

# Keep out of reach of children under 6 years of age.

If more than used for brushing is accidently swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

| Children under 6 years of age:              | Instruct in good brushing and rinsing habits (to minimize swallowing). Supervise children as necessary until capable of using without supervision. |
|---|--|
| Adults and children 2 years of age & older: | Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.                                 |

Children under 2 years of age: Consult a dentist or doctor.

#### Other information

- $\bullet~$  Store in a cool place, below 86°F, away from heat
- Do not use if quality seal is broken or missing

# Inactive ingredients

Calcium Carbonate, Water, Glycerin, Sorbitol, Hydrated Silica, Sodium Lauryl Sulfate, Flavor, Cellulose Gum, Hydroxyethylcellulose, Sodium Silicate, Sodium Saccharin, Trisodium Phosphate.

## **Questions or comments**

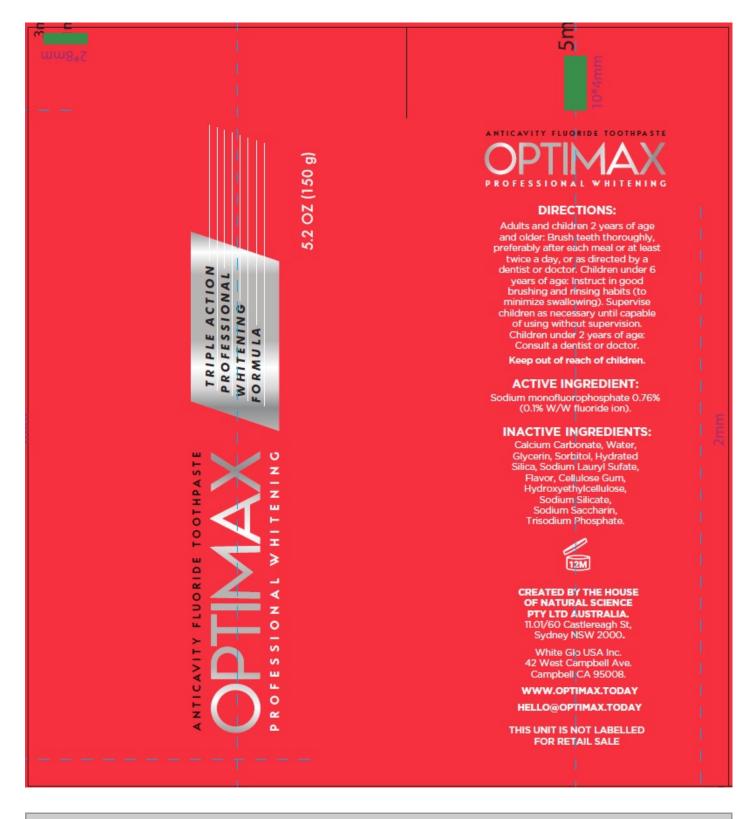
For customer enquiries, please contact: hello@optimax.today White Glo USA INC. 42 West Campbell Avenue, Third Floor, Campbell, California, 95008. www.optimax.today

# **Package Labeling:**





2 X 5.2 OZ (150 g)



## **OPTIMAX PROFESSIONAL WHITENING ANTICAVITY FLUORIDE**

sodium monofluorophosphate kit

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73656-038

#### **Packaging**

| # Item C     | ode Pa      | ackage Description | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
|--------------|-------------|--------------------|-----------------------------|---------------------------|
| 1 NDC:73656- | 038-00 1 in | 1 KIT              | 12/08/2025                  |                           |

| Quantity | of | <b>Parts</b> |
|----------|----|--------------|
|----------|----|--------------|

| ı | Part # | Package Quantity | Total Product Quantity |
|---|--------|------------------|------------------------|
|   |        |                  |                        |

**Part 1** 1 TUBE 150 g in 2

# Part 1 of 1

# **OPTIMAX PROFESSIONAL WHITENING ANTICAVITY FLUORIDE**

sodium monofluorophosphate paste

### **Product Information**

Item Code (Source) NDC:73656-037

**Route of Administration** DENTAL

|   | Active Ingredient/Active Moiety   |                      |                  |  |  |
|---|---|----------------------|------------------|--|--|
|   | Ingredient Name   | Basis of<br>Strength | Strength         |  |  |
| l | <b>SODIUM MONOFLUOROPHOSPHATE</b> (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O) | FLUORIDE ION         | 7.6 mg<br>in 1 g |  |  |

| Inactive Ingredients  |          |
|---|----------|
| Ingredient Name   | Strength |
| CALCIUM CARBONATE (UNII: H0G9379FGK)                          |          |
| WATER (UNII: 059QF0KO0R)                                      |          |
| GLYCERIN (UNII: PDC6A3C0OX)                                   |          |
| SORBITOL (UNII: 506T60A25R)                                   |          |
| HYDRATED SILICA (UNII: Y6O7T4G8P9)                            |          |
| SODIUM LAURYL SULFATE (UNII: 368GB5141J)                      |          |
| CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311) |          |
| HYDROXYETHYLCELLULOSE (UNII: T4V6TWG28D)                      |          |
| SODIUM SILICATE (UNII: IJF18F77L3)                            |          |
| SODIUM SACCHARIN (UNII: SB8ZUX40TY)                           |          |

| Pa | Packaging            |  |                         |                       |
|----|----------------------|--|-------------------------|-----------------------|
| #  | Item Code            | Package Description                                | Marketing Start<br>Date | Marketing End<br>Date |
| 1  | NDC:73656-037-<br>00 | 1 in 1 BOX   |                         |                       |
| 1  |                      | 150 g in 1 TUBE; Type 0: Not a Combination Product |                         |                       |

| Marketing Information |   |                         |                       |  |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |  |
| OTC Monograph Drug    | M021  | 12/08/2025              |                       |  |
|                       |   |                         |                       |  |

| Marketing Information |   |                         |                       |
|-----------------------|---|-------------------------|-----------------------|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |
| OTC Monograph Drug    | M021  | 12/08/2025              |                       |
|                       |   |                         |                       |

Labeler - WHITE GLO USA INC (117345666)

Registrant - WHITE GLO USA INC (117345666)

Revised: 9/2025 WHITE GLO USA INC