

**OPTIMAX ANTICAVITY FLUORIDE COTTON CANDY- sodium monofluorophosphate paste**  
**WHITE GLO USA INC**

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**Optimax Anticavity Fluoride Toothpaste, Cotton Candy**

**Drug Facts**

**Active ingredient**

Sodium Monofluorophosphate 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 accidentally swallowed, get medical help or contact a Poison

**Directions**

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Adults and children 2 years &	Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.
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

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

**Other information**

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

**Inactive ingredients**



ANTICAVITY FLUORIDE TOOTHPASTE

OPTIMAX

FLAVORED TOOTHPASTE

preferably after each meal or at least twice a day, or as directed by a dentist or doctor. 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. Children under 2 years of age: Consult a dentist or doctor.

Keep out of reach of children

**ACTIVE INGREDIENT:**

Sodium monofluorophosphate 0.76% (0.1% W/W fluoride ion).

**INACTIVE INGREDIENTS:**

Calcium Carbonate, Aqua, Glycerin, Sorbitol, Hydrated Silica, Sodium Lauryl Sulfate, Flavor, Cellulose Gum, Hydroxyethylcellulose, Sodium Silicate, Sodium Saccharin, Trisodium Phosphate, D&C Red No. 33



CREATED BY THE HOUSE  
OF NATURAL SCIENCE  
PTY LTD AUSTRALIA.

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THIS UNIT IS NOT LABELLED  
FOR RETAIL SALE

## OPTIMAX ANTICAVITY FLUORIDE COTTON CANDY

sodium monofluorophosphate paste

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73656-033
Route of Administration	DENTAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
<b>SODIUM MONOFLUOROPHOSPHATE</b> (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)			FLUORIDE ION	7.6 mg in 1 g
Inactive Ingredients				
Ingredient Name				Strength
<b>CALCIUM CARBONATE</b> (UNII: H0G9379FGK)				
<b>WATER</b> (UNII: 059QF0KO0R)				
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)				
<b>SORBITOL</b> (UNII: 506T60A25R)				
<b>HYDRATED SILICA</b> (UNII: Y6O7T4G8P9)				
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)				
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED</b> (UNII: K679OBS311)				
<b>HYDROXYETHYLCELLULOSE</b> (UNII: T4V6TWG28D)				
<b>SODIUM SILICATE</b> (UNII: IJF18F77L3)				
<b>SODIUM SACCHARIN</b> (UNII: SB8ZUX40TY)				
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73656-033-00	1 in 1 BOX	04/04/2025	
1		80 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		M021	04/04/2025	

**Labeler** - WHITE GLO USA INC (117345666)

**Registrant** - WHITE GLO USA INC (117345666)