OPTIMAX PURPLE ANTICAVITY FLUORIDE- sodium monofluorophosphate
WHITE GLO USA INC

Optimax Purple Anticavity Fluoride Toothpaste

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 Control Center right away.

Directions

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 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.

Other information

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

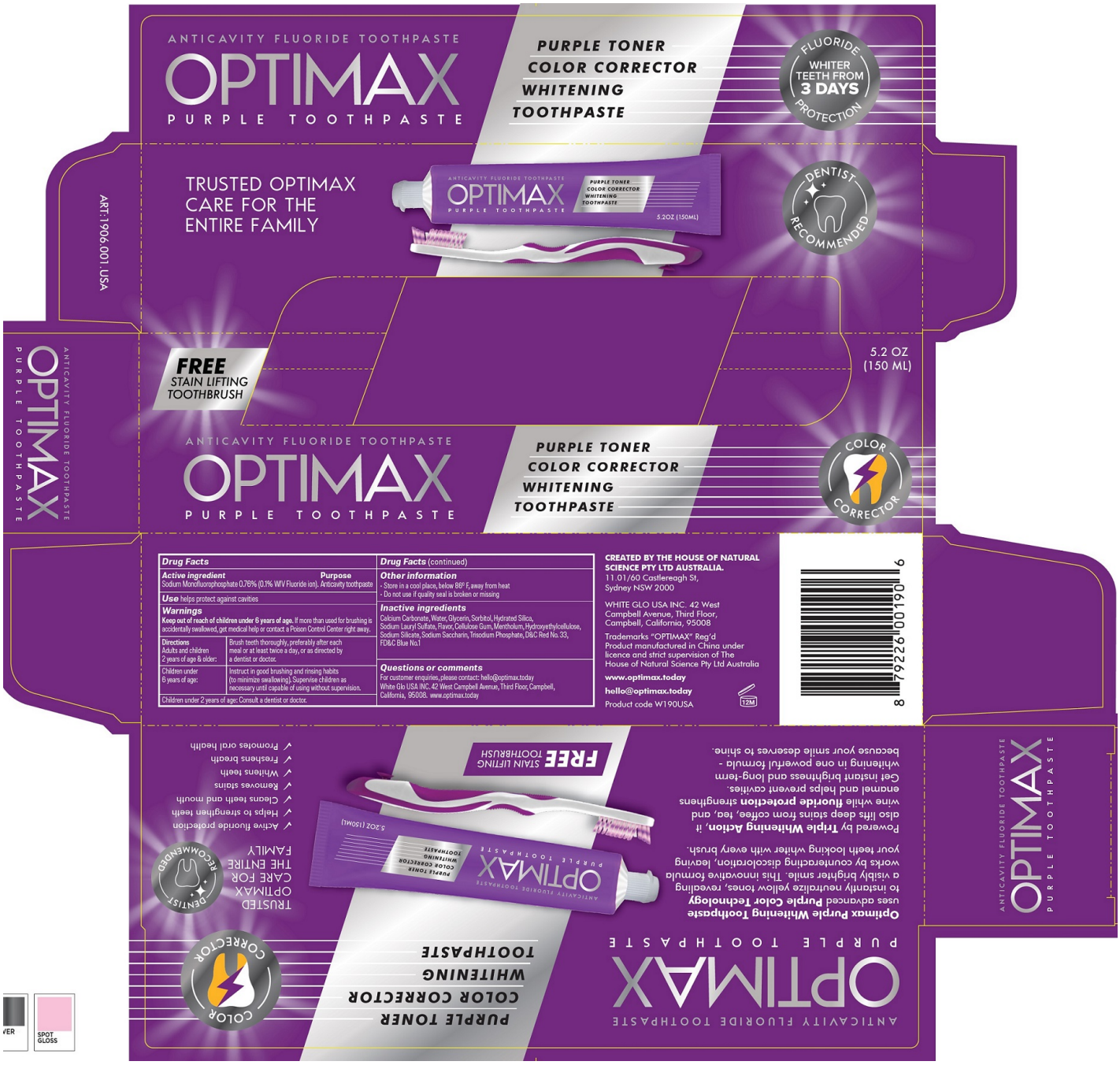
Inactive ingredients

Calcium Carbonate, Water, Glycerin, Sorbitol, Hydrated Silica, Sodium Lauryl Sulfate, Flavor, Cellulose Gum, Mentholum, Hydroxyethylcellulose, Sodium Silicate, Sodium Saccharin, Trisodium Phosphate, D&C Red No.33, FD&C Blue No.1

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:



ANTICAVITY FLUORIDE TOOTHPASTE

OPTIMAX

PURPLE TOOTHPASTE

PURPLE TONER
COLOR CORRECTOR
WHITENING
TOOTHPASTE

5.2 OZ (150 ML)

ANTICAVITY FLUORIDE TOOTHPASTE

OPTIMAX

PURPLE TOOTHPASTE

DIRECTIONS:

Adults and children 2 years of age and older: 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 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, Aroma, Cellulose Gum, Mentholum, Hydroxyethylcellulose, Sodium Silicate, Sodium Saccharin, Trisodium Phosphate, D&C Red No.33, FD&C Blue No.1.

12M

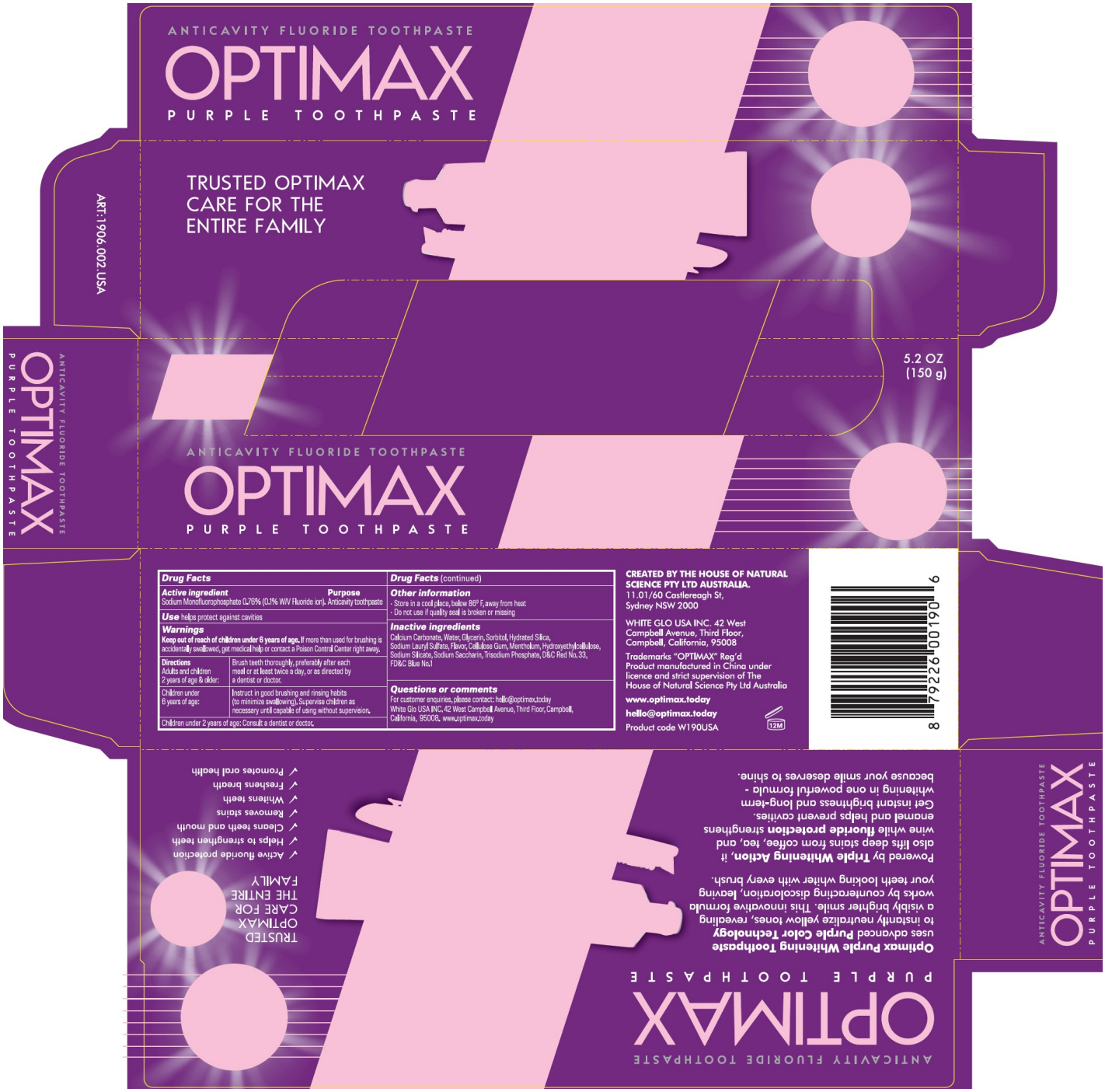
CREATED BY THE HOUSE OF NATURAL SCIENCE PTY LTD AUSTRALIA.
11.01/60 Castlereagh St,
Sydney NSW 2000.

White Glo USA Inc.
42 West Campbell Ave.
Campbell CA 95008.

WWW.OPTIMAX.TODAY

HELLO@OPTIMAX.TODAY

THIS UNIT IS NOT LABELLED FOR RETAIL SALE



OPTIMAX PURPLE ANTICAVITY FLUORIDE

sodium monofluorophosphate kit

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73656-039

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73656-039-00	1 in 1 KIT	12/08/2025	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 TUBE	150 g

Part 1 of 1

OPTIMAX PURPLE ANTICAVITY FLUORIDE

sodium monofluorophosphate paste

Product Information	
Item Code (Source)	NDC:73656-040
Route of Administration	DENTAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	7.6 mg 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: K679OBS311)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
SODIUM SILICATE (UNII: IJF18F77L3)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73656-040-00	1 in 1 BOX		
1		150 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	12/08/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	12/08/2025	

Labeler - WHITE GLO USA INC (117345666)

Registrant - WHITE GLO USA INC (117345666)

Revised: 10/2025 WHITE GLO USA INC