OPTIMAX ANTICAVITY FLUORIDE WATERMELON- sodium monofluorophosphate paste
WHITE GLO USA INC

Optimax Anticavity Fluoride Toothpaste, Watermelon

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 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 86ºF, away from heat
- Do not use if quality seal is broken or missing

Inactive ingredients

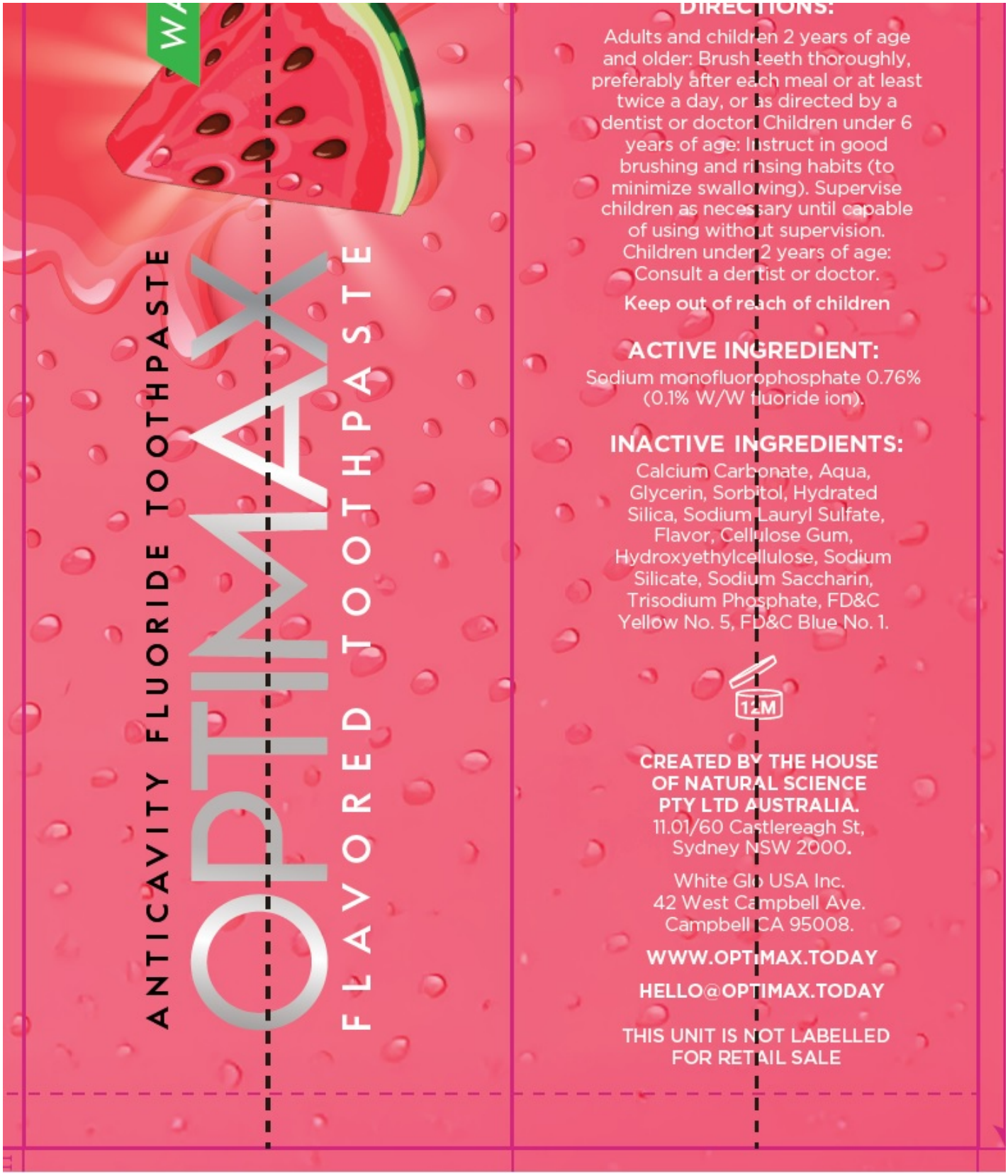
Calcium Carbonate, Aqua, Glycerin, Sorbitol, Hydrated Silica, Sodium Lauryl Sulfate, Flavor, Cellulose Gum, Hydroxyethylcellulose, Sodium Silicate, Sodium Saccharin, Trisodium Phosphate, FD&C Yellow No. 5, FD&C Blue No. 1.

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:



OPTIMAX ANTICAVITY FLUORIDE WATERMELON

sodium monofluorophosphate paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73656-032

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)	
HYDROXYETHYLCELLULOSE (UNII: T4V6TWG28D)	
SODIUM SILICATE (UNII: IJF18F77L3)	
SODIUM SACCHARIN (UNII: SB8ZUX40TY)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73656-032-00	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)

Revised: 5/2025 WHITE GLO USA INC