POP GEL WHATZZUP WATERMELON- sodium fluoride paste, dentifrice
Tai Guk Pharm. Co., Ltd.

POP Gel Whatzzup Watermelon Toothpaste 160g

Drug Facts

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Active Ingredient

SODIUM FLUORIDE 0.243% (0.13% w/v fluoride ion)

Purpose

Anticavity toopaste

Use

helps protect against cavities

Warnings

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Keep out of reach of children.

Keep out of reach of children under 6 years of age. Do not swallow. If you accidentally swallow more than used for brushing, seek professional assistance or contact a poison control center right away. If an allergic reaction develops, discontinue use immediately.

Directions

- adults and children 2 years of age and and older: Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist.
- do not swallow.
- to minimize swallowing use a pea-sized amount in children under 6.
- supervise children's brushing until good habits are established.
- **children under 2 years: ask a dentist.**

Inactive Ingredients

GLYCERIN, WATER, HYDRATED SILICA, SODIUM LAURYL SULFATE, FLAVOR, XANTHAN GUM, TETRASODIUM PYROPHOSPHATE, POLOXAMER 407, SODIUM POLYACRYLATE, ZINC STEARATE, SODIUM SACCHARIN, SORBITAN OLEATE, TITANIUM DIOXIDE,

SUCRALOSE, HYDROXYAPATITE, NIACINAMIDE, TOCOPHERYL ACETATE, XYLITOL

Questions?

1-866-993-4709

Principal Display Panel - 160 g Tube Carton

POP

Playful. Original. Personal.

GEL TOOTHPASTE
Electric Toothbrush Suitable

3X faster fluoride release*
Repairs & protects enamel

NET WT. 5.64 OZ (160 G)

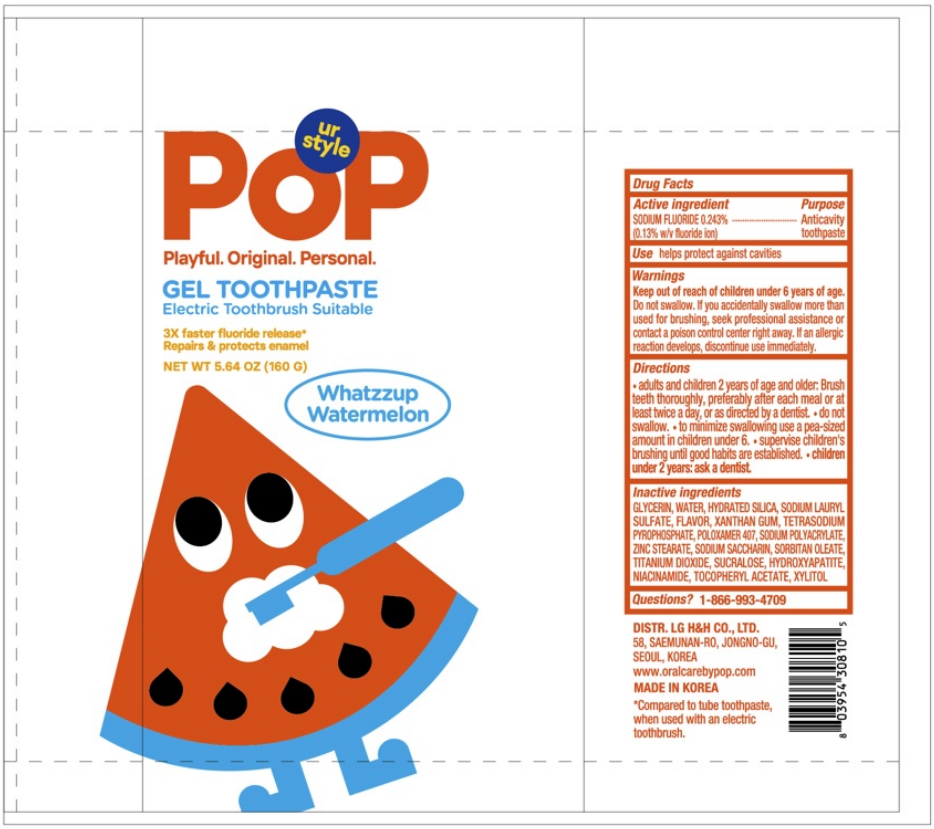
Whatzzup Watermelon

DISTR. LG H&H CO., LTD.

58, SAEMUNAN-RO, JONGNO-GU,
SEOUL, KOREA
www.oralcarebypop.com

MADE IN KOREA

*Compared to tube toothpaste,
when used with an electric
toothbrush.



POP GEL WHATZZUP WATERMELON

sodium fluoride paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43136-522
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.3 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)	
NIACINAMIDE (UNII: 25X51I8RD4)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
WATER (UNII: 059QF0KO0R)	

ZINC STEARATE (UNII: H92E6QA4FV)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
XYLITOL (UNII: VCQ006KQ1E)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	MINT (Watermelon)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43136-522-60	1 in 1 CARTON	09/18/2023	
1		160 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	09/18/2023	

Labeler - Tai Guk Pharm. Co., Ltd. (689060246)

Registrant - LG H&H CO., LTD. (688276187)

Establishment

Name	Address	ID/FEI	Business Operations
Tai Guk Pharma. Co., Ltd.		689060246	manufacture(43136-522)

Revised: 1/2025 Tai Guk Pharm. Co., Ltd.