

PERIOE POP COMPLETE PROTECTION PREBIOTIC MINT BLAST- sodium fluoride paste, dentifrice
Tai Guk Pharm. Co., Ltd.

PERIOE POP Complete Protection Prebiotic Toothpaste Mint Blast 120 g

Drug Facts

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Active Ingredient

SODIUM FLUORIDE 0.243% (0.14% w/v fluoride ion)

Purpose

Anticavity toothpaste

Use

- helps protect against cavities

Warnings

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 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

WATER, SORBITOL, HYDRATED SILICA, SODIUM LAURYL SULFATE, GLYCERIN, FLAVOR, CELLULOSE GUM, TETRASODIUM PYROPHOSPHATE, XANTHAN GUM, TITANIUM DIOXIDE, SORBITAN OLEATE, SODIUM SACCHARIN, CHARCOAL POWDER, BIFIDA FERMENT EXTRACT FILTRATE, SODIUM CHLORIDE, TOCOPHERYL ACETATE, XYLITOL

Questions?

1-800-367-2866

Distributed by

DISTR. LG H&H CO., LTD.

58, SAEMUNAN-RO, JONGNO-GU, SEOUL, KOREA

www.lghnh.com

LG H&H

MADE IN KOREA

Principal Display Panel - 120 g Tube Carton

PERIOE

POP

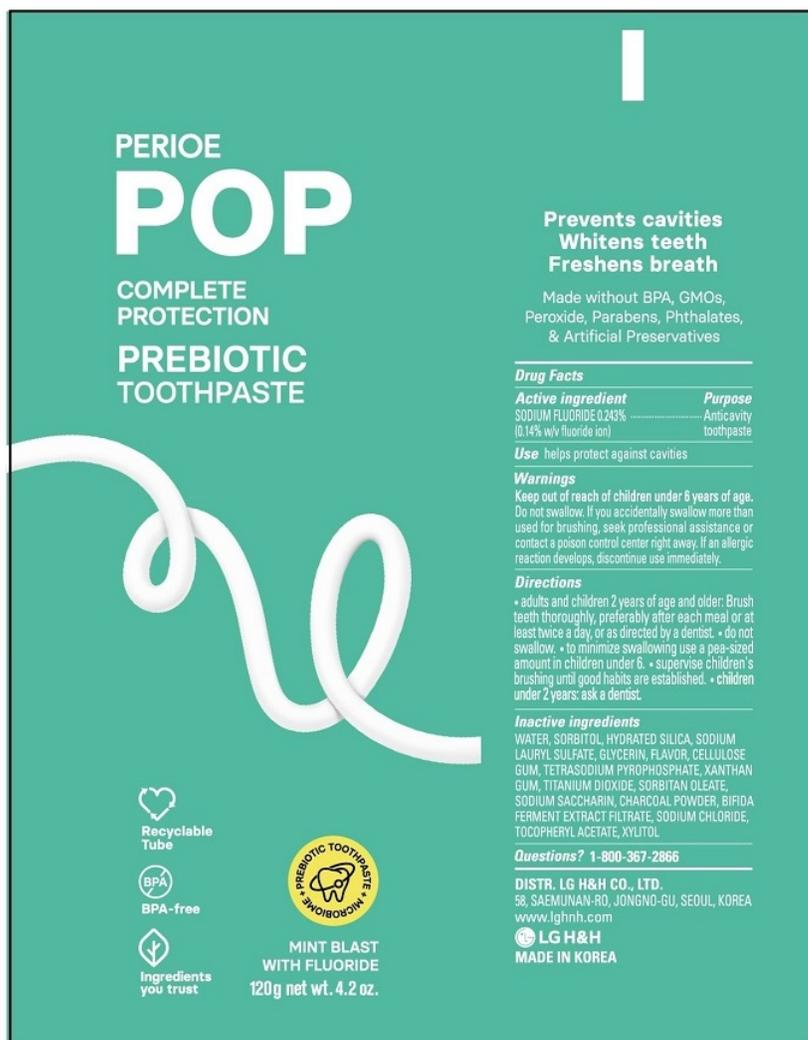
**COMPLETE
PROTECTION**

**PREBIOTIC
TOOTHPASTE**

MINT BLAST
WITH FLUORIDE

120g net wt. 4.2 oz.





PERIOE POP COMPLETE PROTECTION PREBIOTIC MINT BLAST
sodium fluoride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43136-603
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080)	FLUORIDE ION	1.4 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
MICA (UNII: V8A1AW0880)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KOOR)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
XYLITOL (UNII: VCQ006KQ1E)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ACTIVATED CHARCOAL (UNII: 2P3VWUJ3H10)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor	MINT (Blast)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43136-603-01	1 in 1 CARTON	02/17/2023	
1		120 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	02/17/2023	

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(43136-603)

Revised: 1/2025

Tai Guk Pharm. Co., Ltd.