

POP HINT OF MINT- sodium fluoride paste, dentifrice
LG H&H Co., Ltd.

53208-571-20 POP Hint Of Mint Toothpaste 120g

Drug Facts

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Active Ingredient

SODIUM FLUORIDE 0.243% (0.14% 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

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

HYDROXYAPATITE, NIACINAMIDE, TOCOPHERYL ACETATE, XYLITOL

Questions?

1-866-993-4709

Principal Display Panel - 120 g Tube Carton

POP

Playful. Original. Personal

**REPAIRS ENAMEL
PREVENTS CAVITIES
FRESHENS BREATH**

HINT OF MINT

Fluoride Toothpaste
with Hydroxyapatite &
Vitamins E and B3

NET WT. 4.2 OZ. (120 G)

DIST. LG H&H CO., LTD.
58, SAEMUNAN-RO, JONGNO-GU, SEOUL, KOREA
www.oralcarebypop.com

MADE IN KOREA



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POP

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MADE IN KOREA



POP HINT OF MINT

sodium fluoride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53208-571
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
Sorbitol (UNII: 506T60A25R)	
Sorbitan monooleate (UNII: 06XEA2VD56)	
Tribasic calcium phosphate (UNII: 91D9GV0Z28)	
Niacinamide (UNII: 25X5118RD4)	
.alpha.-tocopherol acetate, DL- (UNII: WR1WPI7EW8)	
Water (UNII: 059QF0KO0R)	

ZINC STEARATE (UNII: H92E6QA4FV)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
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: Y607T4G8P9)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53208-571-20	1 in 1 CARTON	09/18/2023	
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	09/18/2023	

Labeler - LG H&H Co., Ltd. (688276187)

Establishment

Name	Address	ID/FEI	Business Operations
Tai Guk Pharma. Co., Ltd.		689060246	manufacture(53208-571)

Revised: 1/2025

LG H&H Co., Ltd.