DENTI BUONGIORNO FRESH BREATH HERB MINT TOOTH- silica, sodium monofluorophosphate paste, dentifrice
DONG IL PHARMS CO.,LTD

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

DENTI BUONGIORNO FRESH BREATH HERB MINT TOOTHPASTE

Active ingredient(s)

Silica 18.0%

Sodium Monofluorophosphate 0.76%

Purpose

Antiplaque, Anticavity

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.

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Use(s)

• Helps remove plaque and tartar buildup
• Helps prevent cavities
• Helps reduce gum inflammation and promote healthy teeth

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 physician

• Children 2 to 6 years : Use only a pea sized amount and supervise child's brushing and rinsing (to minimize swallowing)

• Children under 2 years : Ask a dentist or physician

Other Information

• Store at temperatures between 1°C and 30°C
• Use within 36 months from the date of manufacture
• Do not store in high/low temperatures or direct sunlight

Questions

+82-1588-9709

Inactive ingredients

D-Sorbitol Solution, Water(Aqua) ,Silicon Dioxide , Concentrated Glycerin, Sodium Cocoyl Glutamate, Polyethylene Glycol 1500, Carboxymethylcellulose Sodium, Sodium Monofluorophosphate, Sodium Bicarbonate, Titanium Oxide, Xylitol, Enzymatically

Modified Stevia, Propolis Extract, Grapefruit Seed Extract, Aloe Extract, Matricaria Extract, Eucalyptus Extract, Sage Extract, Green Tea Extract, Rosemary Extract, Calendula Extract, Leontopodium Alpinum Extract, Curcuma Longa (Turmeric), Centella Extract, Houttuynia Cordata Extract, L-Menthol, Menthol Pulegium Oil, Herb Mint Flavor, Mild Mint Flavor, Sodium Lauroyl Sarcosinate

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73242-1128
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	18 g in 100 g
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.76 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73242-1128-2	1 in 1 CARTON	01/02/2026	
1	NDC:73242-1128-1	100 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
unapproved drug other		01/02/2026	

Labeler - DONG IL PHARMS CO.,LTD (557810721)

Establishment

Name	Address	ID/FEI	Business Operations
DONG IL PHARMS CO., LTD.		557810721	manufacture(73242-1128)

Revised: 1/2026 DONG IL PHARMS CO.,LTD