# DENTI BUONGIORNO ORIGINAL 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.

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# 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)

- Fluoride content: 1,000 ppm
- Removes plaque (anti-plaque)
- Helps reduce bad breath by controlling oral bacteria

#### **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 brushing and rinsing (to minimize swallowing).
- Children under 2 years: Ask a dentist or physician.

#### Other Information

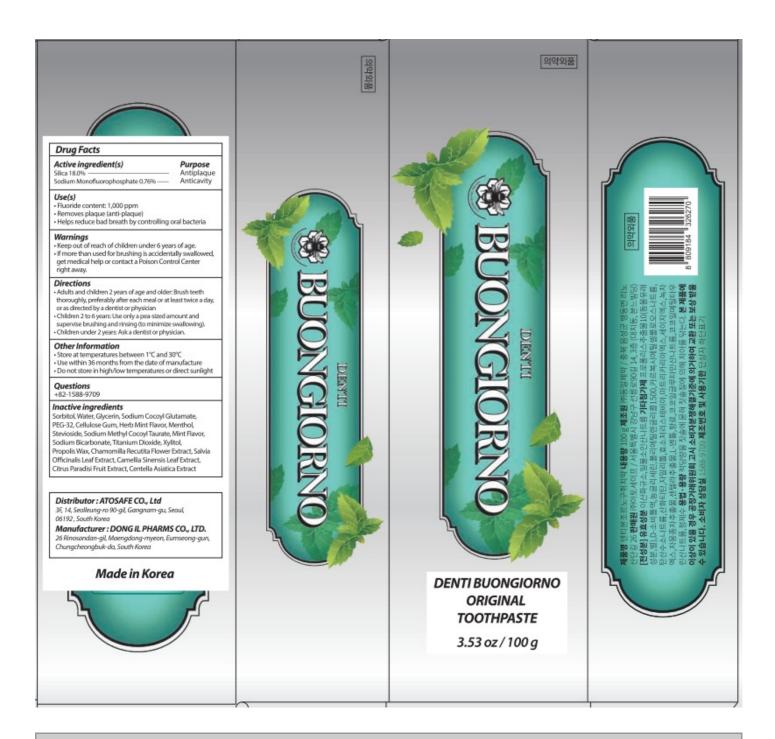
- Store at temperatures between 1°C and 30°C
- Use within 36 months from the date of manufacture
- Do not store in high/low temperatures or direct sunlight

#### **Questions**

+82-1588-9709

# Inactive ingredients

Sorbitol, Water, Glycerin, Sodium Cocoyl Glutamate, PEG-32, Cellulose Gum, Herb Mint Flavor, Menthol, Stevioside, Sodium Methyl Cocoyl Taurate, Mint Flavor, Sodium Bicarbonate, Titanium Dioxide, Xylitol, Propolis Wax, Chamomilla Recutita Flower Extract, Salvia Officinalis Leaf Extract, Camellia Sinensis Leaf Extract, Citrus Paradisi Fruit Extract, Centella Asiatica Extract



# **DENTI BUONGIORNO ORIGINAL TOOTH**

silica, sodium monofluorophosphate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73242-1120
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 -	FLUORIDE ION	0.76 g
UNII:080VPU4080)	PLOORIDE ION	in 100 a

Inactive Ingredients			
Ingredient Name	Strength		
SORBITOL (UNII: 506T60A25R)			
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			
SODIUM COCOYL GLUTAMATE (UNII: BMT4RCZ3HG)			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:73242- 1120-2	1 in 1 CARTON	04/01/2025		
1	NDC:73242- 1120-1	100 g in 1 TUBE; Type 0: Not a Combination Product			

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
	04/01/2025		
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date	

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

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
DONG IL PHARMS CO., LTD.		557810721	manufacture(73242-1120)		

Revised: 10/2025 DONG IL PHARMS CO.,LTD