DENTI BOUNGIORNO ORIGINAL TOOTH- silica, sodium monofluorophosphate paste, dentifrice ATOSAFE 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.

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

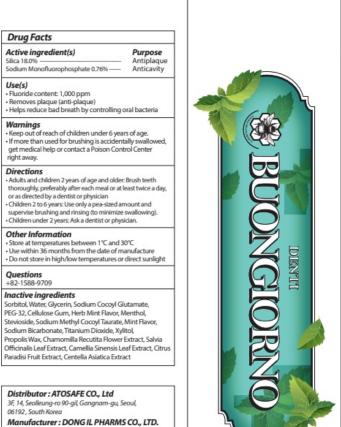
- Store at temperatures between 1°C and 30°C
- Use within 36 months from the date of manufacture
- Do not store in high/low temperatures or direct sunlight

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



Drug Facts Active ingredient(s)

Other Information

26 Rinosandan-gil, Maengdong-myeon, Eumseong-gun, Chungcheongbuk-do, South Korea

Made in Korea

Questions +82-1588-9709 Inactive ingredients

Use(s)



ORIGINAL TOOTHPASTE

3.53 oz/100 g

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DENTI BOUNGIORNO ORIGINAL TOOTH

silica, sodium monofluorophosphate paste, dentifrice

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:85601-010	
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:Q80VPU408O)	I LOOKIDE ION	in 100 g

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

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:85601-010- 02	1 in 1 CARTON	04/01/2025		
1	NDC:85601-010- 01	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 - ATOSAFE CO., Ltd (688995680)

Establishment					
Name	Address	ID/FEI	Business Operations		
DONG IL PHARMS CO., LTD.		557810721	manufacture(85601-010)		

Revised: 5/2025 ATOSAFE CO., Ltd