DENTI BUONGIORNO PLAQUE CARE TOOTH- silica, sodium fluoride, tocopheryl acetate, tetrasodium pyrophosphate 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 17.0%

Sodium Fluoride 0.22%

Tocopheryl Acetate 0.20%

Tetrasodium Pyrophosphate 0.50%

Purpose

Antiplaque, Anticavity, Antigingivitis, Antitartar

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 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, Citrus Mint Flavor, Menthol, Stevioside, Sodium Bicarbonate, Mentha Piperita Oil, Lemon Flavor,

Sodium Methyl Cocoyl Taurate, Titanium Dioxide, Xylitol, Propolis Wax, Chamomilla Recutita Flower Extract, Salvia Officinalis Leaf Extract, Camellia Sinensis Leaf Extract, Eucalyptus Globulus Leaf Extract, Citrus Paradisi Fruit Extract, Centella Asiatica Extract

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85601-060
Route of Administration	ORAL		

Active Ingredient/Active Moiety		

Ingredient Name	Basis of Strength	Strength
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) (.ALPHA.-TOCOPHEROL - UNII:H4N855PNZ1)	.ALPHA.-TOCOPHEROL	0.2 g in 100 g
SODIUM PYROPHOSPHATE (UNII: O352864B8Z) (PYROPHOSPHORIC ACID - UNII:4E862E7GRQ)	SODIUM PYROPHOSPHATE	0.5 g in 100 g
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	17 g in 100 g
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.22 g in 100 g

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85601-060-02	1 in 1 CARTON	04/01/2025	
1	NDC:85601-060-01	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		04/01/2025	

Labeler - ATOSAFE CO., Ltd (688995680)

Establishment

Name	Address	ID/FEI	Business Operations
DONG IL PHARMS CO., LTD.		557810721	manufacture(85601-060)

Revised: 10/2025 ATOSAFE CO., Ltd