DENTI BUONGIORNO GUM CARE TOOTH- silica, sodium fluoride, tocopheryl acetate 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.

Active ingredient(s)

Silica 17.0%

Sodium Fluoride 0.22%

Tocopheryl Acetate 0.20%

Purpose

Antiplaque, Anticavity, Antigingivitis

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 protect against cavities
• Helps reduce gum inflammation and maintain healthy gums
• Aids in the prevention of periodontal disease

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, Menthol, Mentha Piperita Oil, Sodium Lauroyl Sarcosinate, Stevioside, Sodium Bicarbonate, Cooling Flavor, Aqua Mint Flavor, Xylitol, Propolis Wax, Hydrolyzed Collagen, Chamomilla Recutita Flower Extract, Salvia Officinalis Leaf Extract, Camellia Sinensis Leaf Extract, Citrus

Paradisi Fruit Extract, Centella Asiatica Extract, Aloe Barbadensis Leaf Extract, Eucalyptus Globulus Leaf Extract, Calendula Officinalis Flower Extract, Rosmarinus Officinalis Leaf Extract

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73242-1118
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	15 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
SODIUM COCOYL GLUTAMATE (UNII: BMT4RCZ3HG)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73242-1118-2	1 in 1 CARTON	04/01/2025	
1	NDC:73242-1118-1	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		04/01/2025	

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

Establishment

Name	Address	ID/FEI	Business Operations
DONG IL PHARMS CO., LTD.		557810721	manufacture(73242-1118)

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