DENTI BUONGIORNO PEACH MINT WHITENING TOOTH- hydrogen peroxide, silica 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)

Hydrogen Peroxide 2.14%

Silica 7%

Purpose

Whitening, Anti-plaque

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)

• Teeth whitening.
• Remove dental plaque (anti-prag).
• Keep your teeth white and strong.

• Keeps the oral cavity clean.

• Refreshes the oral cavity.

• Prevents cavities and eliminates bad breath.

• Enhance the aesthetic effect.

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

Glycerin, PEG-32, Xanthan gum, Citric Acid, Sodium Citrate, Disodium Pyrophosphate, Sodium metaphosphate, Disodium EDTA, Stevioside, Aloe Barbadensis Leaf Extract, Chamomilla Recutita Flower Extract, Eucalyptus Globulus Leaf Extract, Salvia Officinalis Leaf Extract, Camellia Sinensis Leaf Extract, Rosmarinus Officinalis Leaf Extract, Chaenomeles Sinensis Fruit Extract, Leontopodium Alpinum Extract, Menthol, Mentha Pulegium Oil, Peach Mint Flavor, Honey Peach Flavor, Poloxamer 407, Sodium Lauryl Sulfate, Water

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hydrogen peroxide, silica paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73242-1119
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	7 g in 100 g
HYDROGEN PEROXIDE (UNII: BBX060AN9V) (HYDROGEN PEROXIDE - UNII:BBX060AN9V)	HYDROGEN PEROXIDE	2.14 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
XANTHAN GUM (UNII: TTV12P4NEE)	
CITRIC ACID (UNII: 2968PHW8QP)	
PEG-32 (UNII: 1212Z7S33A)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73242-1119-2	1 in 1 CARTON	07/01/2025	
1	NDC:73242-1119-1	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		07/01/2025	

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

Establishment

Name	Address	ID/FEI	Business Operations
DONG IL PHARMS CO., LTD.		557810721	manufacture(73242-1119)

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