

DOCTRUST K TOOTH- silicon dioxide, sodium monofluorophosphate gel, 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

Silicon dioxide 16.0%
Sodium monofluorophosphate 0.740%
(Total Fluoride Content 977 ppm Approx.)

Inactive ingredients

Carboxymethyl cellulose, D-sorbitol solution, Sodium Hydrogencarbonate, Enzymatically modified stevia, Xylitol, Hydroxyapatite, Turmeric root extract, Centella asiatica extract, Houttuynia cordata Thunberg extract, Green tea extract, Salvia officinalis (sage) extract, Eucalyptus extract, natural aromatics (Lime, Green grape) Sodium cocoyl glutamate, Purified water

PURPOSE

Anticavity Toothpaste

WARNINGS

- Do not swallow. Rinse mouth thoroughly after use.
- For children under 6 years of age, use only a pea-sized amount of toothpaste under the supervision of a guardian to avoid sucking or swallowing.
- If a child under 6 years of age swallows a large amount of toothpaste, consult a physician or dentist immediately.
- Keep out of reach of children under 6 years of age.

KEEP OUT OF REACH OF CHILDREN

- Keep out of the reach of children under 6 years of age.

Uses

- Helps protect against cavities

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 doctor.
- Children under 6 years: To minimize swallowing, use a pea-sized amount and supervise brushing until good habits are established.
- Children under 2 years: Consult a dentist or doctor.

Other Information

- Do not store this product in an inappropriate place such as high or low temperatures or under direct sun light

Questions

- <https://docsmedi.kr>
- Tel: 82-031-922-2240, Fax: 82-031-365-4596

DOCSMEDI K TOOTHPASTE		CAVITY CARE	Harmful Ingredients 13無 Toothpaste SLS, Artificial Dyes, Artificial Sweetener, Benzoic Acid, Saccharin, Ethanol, Paraben, MIT /CMIT, Microplastics, Tar Dye, Synthesis Oil, Antibiotics																																														
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73242-1123
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	16 g in 100 g
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.74 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	
SORBITOL (UNII: 506T60A25R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73242-1123-2	1 in 1 CARTON	12/01/2022	
1	NDC:73242-1123-1	60 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		12/01/2022	

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

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

Establishment

Name	Address	ID/FEI	Business Operations
DONG IL PHARMS CO., LTD.		557810721	manufacture(73242-1123)