

DR. DADDYS KIDS TOOTH- dental type silica, sodium pyrophosphate, sodium monofluorophosphate paste, dentifrice
TB Healthcare 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.

Silicon Dioxide (Dental Type Silica), Sodium Pyrophosphate (Tetrasodium Pyrophosphate), Sodium Monofluorophosphate

D-Sorbitol Solution, Concentrated Glycerin, Xanthangum, Sodium Cocoyl Glutamate, Ascorbic Acid, Hydroxyapatite, Xylitol, Sodium Chloride, Green Tea Extract, Eucalyptus Extract, Matricaria Extract, Aloe Extract, Sage Extract, Grapefruit Seed Extract, Natural Green grape flavor, Water

ANTI-CAVITY

Keep out of reach of children

Put an appropriate amount on a toothbrush and brush teeth.

Storage method

1. Keep it at room temperature in a classified container.
2. Cover and store at room temperature.
3. Store in a not moisture and cool place.
4. Air may come out during use of this product, but there is no problem with its weight.

Usage Precautions

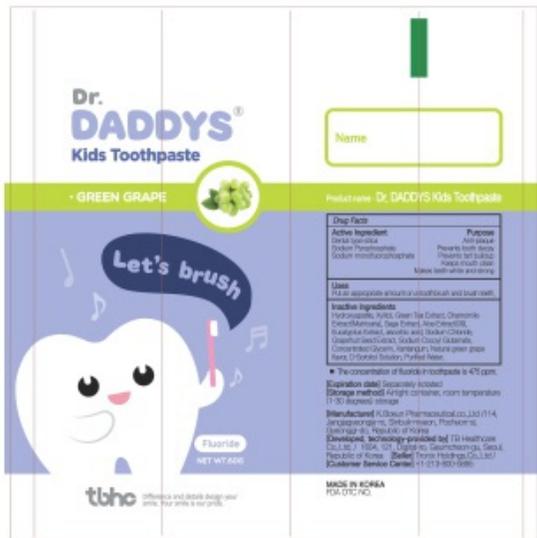
1. Be careful not to swallow. Rinse mouth thoroughly after use
2. If the use of toothpaste causes abnormalities such as gums or mouth injury, discontinue use and consult a doctor or dentist.
3. For children under 6 years of age, use a small amount of toothpaste as small as pea per use, and use under the guidance of a guardian to avoid sucking or swallowing.
4. If a child under 6 years old swallows large amount, consult with a doctor or dentist immediately.
5. Keep out of the reach of children under 6 years of age.

For dental use only

Dr. DADDYS Kids Toothpaste

Fluoride(475ppm) / NET WT.60G

For children who can spit out toothpaste



DR. DADDYS KIDS TOOTH

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76884-0010
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	13 g in 100 g
SODIUM PYROPHOSPHATE (UNII: O352864B8Z) (PYROPHOSPHORIC ACID - UNII:4E862E7GRQ)	SODIUM PYROPHOSPHATE	0.5 g in 100 g
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU4080)	FLUORIDE ION	0.36 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
XYLITOL (UNII: VCQ006KQ1E)	

WATER (UNII: 059QF0KO0R)

SODIUM CHLORIDE (UNII: 451W47IQ8X)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76884-0010-1	60 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2021	

Labeler - TB Healthcare Co., Ltd. (695035143)

Registrant - TB Healthcare Co., Ltd. (695035143)

Establishment

Name	Address	ID/FEI	Business Operations
TB Healthcare Co., Ltd.		695035143	label(76884-0010)

Establishment

Name	Address	ID/FEI	Business Operations
K.Boeun Pharmaceutical Co.,Ltd.		695674074	manufacture(76884-0010)

Revised: 10/2021

TB Healthcare Co., Ltd.