DR. DADDYS GUMTOOTHPASTE- silicon dioxide, tetrasodium pyrophosphate, tocopherol acetate, 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 Dioxidel,Tetrasodium Pyrophosphate,Tocopherol Acetate,Sodium Monofluorophosphate

D-Sorbitol Solution,Concentrated Glycerin,Xanthan gum,Sodium Cocoyl Glutamate,Ascorbic Acid,Hydroxyapatite,Xylitol,Sodium Chloride,Grapefruit Seed Extract,Glycyrrhiza Extract,Green Tea Extract,Matricaria Extract,Sage Extract,Aloe Extract,Eucalyptus Extract,Ginseng Extract,Propolis Extract,Ginger Powder,Myrrh Tincture,l-Menthol,Flavor,Water

Reduces bad breath, Prevents tartar buildup, Keeps mouth clean, Makes teeth white and strong, Prevent periodontal disease, gum disease

Keep out of reach of children

Put an appropriate amount on a toothbrush and brush teeth

1. Be careful not to swallow. Rinse mouth thoroughly after use

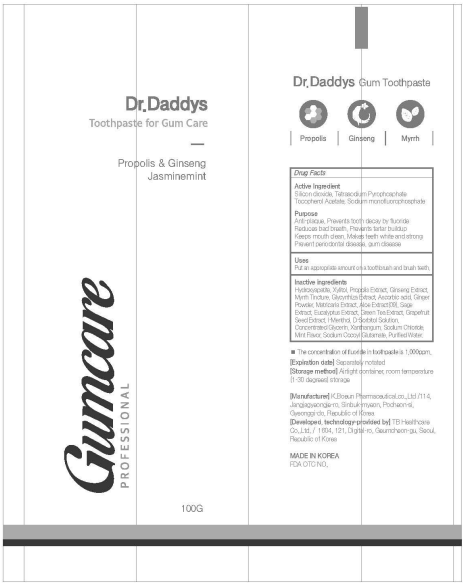
2. If the use of toothpaste causes abnormalities such as gums or mouth injury, discontinue use and consult a doctor or dentist.

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 GUMTOOTHPASTE

silicon dioxide, tetrasodium pyrophosphate, tocopherol acetate, sodium monofluorophosphate paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76884-0012
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0) (.ALPHA.-TOCOPHEROL - UNII:H4N855PNZ1)	.ALPHA.-TOCOPHEROL ACETATE	0.2 g in 100 g
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	17 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:Q80VPU408O)	FLUORIDE ION	0.76 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
XYLITOL (UNII: VCQ006KQ1E)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76884-0012-1	100 g in 1 TUBE; Type 0: Not a Combination Product	04/13/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/06/2021	

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

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

Establishment

Name	Address	ID/FEI	Business Operations
K.Boeun Pharmaceutical Co.,Ltd.		695674074	manufacture(76884-0012)

Revised: 4/2022 TB Healthcare Co., Ltd.