DR.DADDYS JUNIOR TOOTHPASTE(MINT FLAVOR)- tetrasodium pyrophosphate, sodium monofluorophosphate, dental type silica 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.

Water

Sodium Monofluorophosphate, Dental type silica, Tetrasodium Pyrophosphate

D-Sorbitol Solution Concentrated Glycerin Xanthangum Sodium Cocoyl Glutamate Ascorbic Acid Hydroxyapatite Xylitol **Enzymatically Modified Stevia** Sodium Chloride Flavor(Apple mint) Spearmint Oil Grapefruit Seed Extract Green Tea Extract Matricaria Extract Sage Extract **Eucalyptus Extract**

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



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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76884-0007
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	15 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	0.758 g in 100 g

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

	Packaging			
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date
:	NDC:76884- 0007-1	60 g in 1 TUBE; Type 0: Not a Combination Product	02/26/2022	

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
	02/26/2022		
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date	

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(76884-0007)	

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