DR. DADDYS KIDS TOOTHPASTE(RASPBERRY FLAVOR)- dental type silica, tetrasodium 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 raspberry flavor, Water

ANTI-CAVITY

Keep out of reach of children

Put an appropriate amount on a toothbrush and brush teeth.

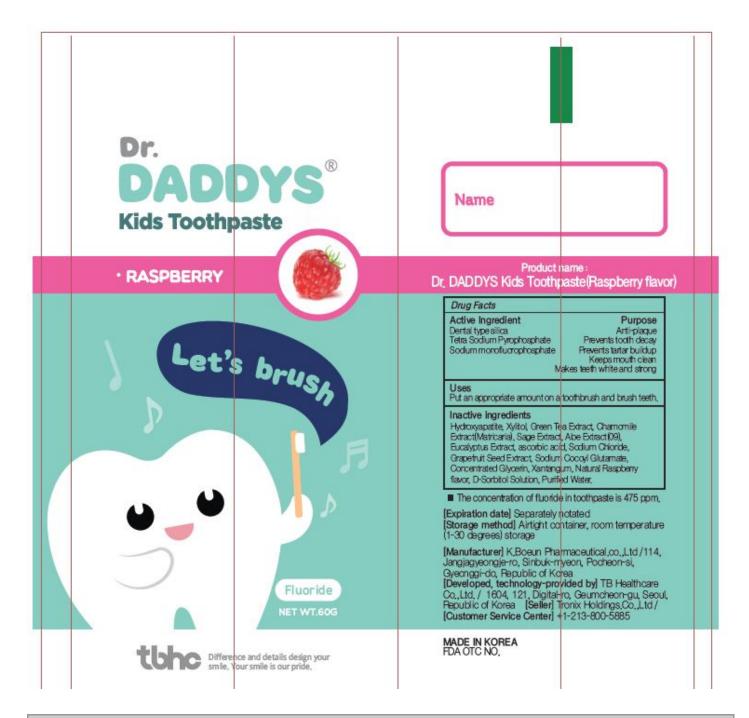
Storage method

- 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



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dental type silica, tetrasodium pyrophosphate, sodium monofluorophosphate paste, dentifrice

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76884-0011	
Route of Administration	DENTAL			

A	Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength	
SI	ILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	14 g in 100 g	

SODIUM PYROPHOSPHATE (UNII: 0352864B8Z) (PYROPHOSPHORIC ACID - UNII:4E862E7GRQ)	SODIUM PYROPHOSPHATE	0.5 g in 100 g
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ 56Q) (FLUORIDE ION - UNII: Q80VPU408O)	FLUORIDE ION	0.36 g in 100 g

Inactive Ingredients			
Ingredient Name	Strength		
XYLITOL (UNII: VCQ006KQ1E)			
WATER (UNII: 059QF0KO0R)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			

ı	Packaging				
	#	# Item Code Package Description		Marketing Start Date	Marketing End Date
		NDC:76884- 0011-1	60 g in 1 TUBE; Type 0: Not a Combination Product	09/20/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		09/20/2021		
		09/20/2021		

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(76884-0011)	

Establishment				
Name	Address	ID/FEI	Business Operations	
K.Boeun Pharmaceutical Co.,Ltd.		695674074	manufacture(76884-0011)	

Revised: 10/2021 TB Healthcare Co., Ltd.