KILL FIRETOOTHPASTE- sodium fluoride paste
Pharmacal-International. 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

Sodium Fluoride 0.02%

Purpose

- Anticavity Toothpaste

Use

- Aids in the prevention of dental cavities

Directions

- Apply appropriate amount on a toothbrush and brush teeth.

Warnings

- Be careful not to swallow. Rinse mouth thoroughly after use.
- If pain occurs when using this product, stop using and consult your dentist.
- For children under 6 years of age, use a pea size amount of toothpaste as recommended by the KFDA. Use under the guidance of a guardian.
- If a child under 6 years old swallows a large amount, consult a physician or dentist immediately.
- Keep out of the reach of children under 6 years old.

Inactive ingredients

Silica,Sorbitol, Tetrasodium Pyrophosphate, Tocopheryl Acetate,
Xylitol, Glucosyl Stevioside, Hydrated Silica, Cellulose Gum, Sodium lauryl Sulfate, Menthol,
Chamomilla Recutita (Matricaria) Extract, Artemisia Princeps Extract, Zingiber Officinale (Ginger) Root Extract,
Melaleuca Alternifolia (Tea Tree) Leaf Oil, Flavor, Gardenia Red, Water

Effect- Efficacy

- Whitens teeth while fighting germs
- Long-lasting freshness and clean mouth feel
- Protects against cavities and bad breath
- Natural brightness of your teeth shines
- Anti plaque
- Helps prevent plaque, gingivitis cavities and gum disease

Product label

Drug Facts

Active Ingredients **Purpose**

SODIUM FLUORIDE 0.02% ..... Anticavity Toothpaste

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Expiration date

- Separately marked

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Melaleuca Alternifolia (Tea Tree) Leaf Oil, Flavor, Gardenia Red, Water

KILL FIRETOOTHPASTE

sodium fluoride paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24765-133
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	SODIUM FLUORIDE	0.02 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
SORBITOL (UNII: 506T60A25R)	
XYLITOL (UNII: VCQ006KQ1E)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
MATRICARIA CHAMOMILLA WHOLE (UNII: G0R4UBI2ZZ)	
GINGER (UNII: C5529G5JPQ)	
TEA TREE OIL (UNII: VIF565UC2G)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24765-133-01	100 g in 1 TUBE; Type 0: Not a Combination Product	06/15/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/15/2021	

Labeler - Pharmacal-International. Co., Ltd. (557805060)

Registrant - Pharmacal-International. Co., Ltd. (557805060)

Establishment

Name	Address	ID/FEI	Business Operations
KMPHARMACEUTICAL Co.,Ltd.		689850153	manufacture(24765-133)

Revised: 10/2024 Pharmacal-International. Co., Ltd.