DR.PAUHLS TOOTHPAST E VANILLE BLACK TEA FLAVOR- silicon dioxide, pyridoxine hydrochloride paste, dentifrice
CURESCRIPT

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

Active ingredients: SILICA 13.0%, PYRIDOXINE HYDROCHLORIDE 0.05%

INACTIVE INGREDIENT

Inactive ingredients:

SORBITOL, SODIUM PCA, GLYCERIN, SODIUM LAURYL SULFATE, FLAVOR, CELLULOSE GUM, MENTHOL, XYLITOL, CITRUSGRANDIS (GRAPEFRUIT) SEED EXTRACT, SODIUM SACCHARIN HYDROXYAPATITE, CAMELLIA SINENSIS LEAF EXTRACT, CALENDULA OFFICINALIS FLOWER EXTRACT, ANTHEMIS NOBILIS FLOWER EXTRACT, PROPOLIS EXTRACT

PURPOSE

Purpose: Anti-plaque, Anti-caries, Anti-periodontitis

WARNINGS

Warnings:

Keep out of the reach of children under 6 years of age.

If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

KEEP OUT OF REACH OF CHILDREN

Keep out of the reach of children under 6 years of age.

Uses

Uses:

- Helps protect against cavities
- reduce plaque
- gum health
- Helps protect against gingivitis, periodontitis
- Helps freshen breath
- brightening teeth

Directions

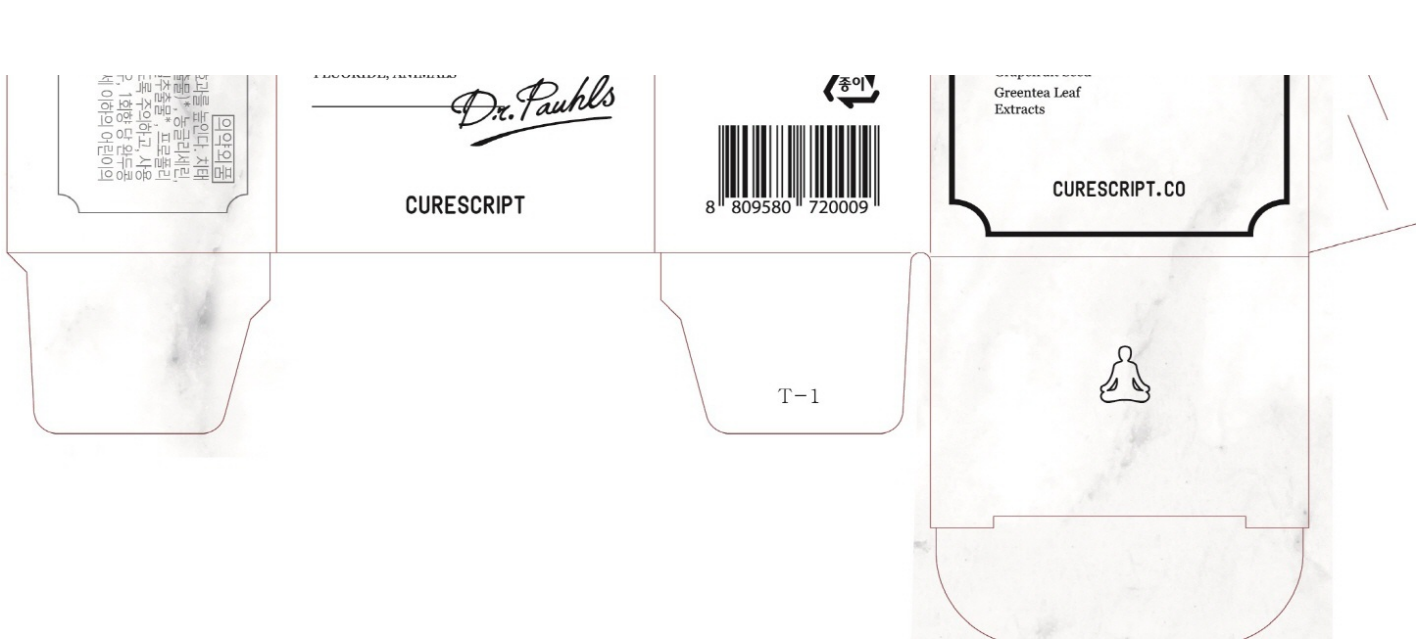
Directions:

Adults and children 2 years of age and older: Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or physician

Children 2 to 6 years: Use only a pea sized amount and supervise child's brushing and rinsing (to minimize swallowing)

Children under 2 years: Ask a dentist or physician

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



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silicon dioxide, pyridoxine hydrochloride paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72265-030
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	13.0 g in 100 g
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV) (PYRIDOXINE - UNII:KV2JZ1BI6Z)	PYRIDOXINE	0.05 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
SORBITOL (UNII: 506T60A25R)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72265-030-02	1 in 1 CARTON	07/01/2018	
1	NDC:72265-030-01	100 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		07/01/2018	

Labeler - CURESCRIPT (694894509)

Registrant - CURESCRIPT (694894509)

Establishment

Name	Address	ID/FEI	Business Operations
Kumho Dental Pharm. Co., Ltd.		631133766	manufacture(72265-030)

Revised: 7/2018 CURESCRIPT