

HONEST-PASTE KIDS- dental type silica paste
BOONCO 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.

80925-201

Active ingredients

Dental type silica 8%

Purpose

Anticavity

Use

helps protect against cavities

Warnings

When using this product, avoid contact with eyes and lips. If contact occurs, rinse with water.

Stop use and ask a dentist if

irritation or redness develops.

Keep out of reach of children.

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

Directions

- children under 1 year: ask a dentist or physician
- children under 6 years: instruct in good brushing and rinsing habits (to minimize swallowing)

Inactive ingredients

glycerin, cellulose gum, ascorbic acid, rosemary extract, scutellaria root extract, green tea extract, xylitol, calendula extract, chamomile extract, enzyme treatment stevia, l-menthol, raspberry flavor, sorbitol, purified water

Package Label



dental type silica paste

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80925-201
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	8 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
LEVOMENTHOL (UNII: BZ1R15MTK7)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
ROSEMARY (UNII: IJ67X351P9)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
SCUTELLARIA BAICALENSIS ROOT (UNII: 7J95K7ID2S)	
XYLITOL (UNII: VCQ006KQ1E)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
CHAMOMILE (UNII: FGL3685T2X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80925-201-01	60 g in 1 TUBE; Type 0: Not a Combination Product	10/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/28/2020	

Labeler - BOONCO Co., Ltd (694871408)

Registrant - BOONCO Co., Ltd (694871408)

Establishment

Name	Address	ID/FEI	Business Operations
BOONCO Co., Ltd		694871408	manufacture(80925-201)

Revised: 1/2022

BOONCO Co., Ltd