DR. ZENNI GUM PROJECT PROPOLI TOOTH FOR GUM DISEASE- sodium monofluorophosphate paste, dentifrice
Zeniton Co.,Ltd.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ACTIVE INGREDIENT

Sodium Monofluorophosphate 0.75%

PURPOSE

Anticavity

Keep out of reach of children

Keep out of reach of children

Uses

■ Helps protect against cavities
■ Removal of plaque

WARNINGS

Keep out or reach of children under 6 yrs. of age. If more than used for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

INACTIVE INGREDIENTS

D-Sorbitol Solution, Silicon Dioxide, Water, Concentrated Glycerin, Propolis Extract, Sodium Chloride, Sodium Cocoyl Glutamate, Xanthangum, Peppermint Oil, Aminocaproic Acid, Aluminum Chlorohydroxy Allantoinate, Grapefruit Seed Extract, Xylitol, Tocopherol Acetate, Calendula Extract, Eucalyptus Extract, Green Tea Extract, Sage Extract, Aloe Extract, Mastic Oil, Myrrh, Chitosan

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 doctor.
■ Children under 6 years: To minimize swallowing, use a pea-sized amount and supervise brushing until good habits are established.
■ Children under 2 years: Consult a dentist or doctor.

Other Information

■ Do not store this product in an inappropriate place such as high or low temperatures or under direct sun light (1~30℃)

QUESTIONS

■ www.zeniton.com

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

DR. ZENNI GUM PROJECT PROPOLI TOOTH FOR GUM DISEASE

sodium monofluorophosphate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73029-0006
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.7581 g in 100 g

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73029-0006-1	100 g in 1 TUBE; Type 0: Not a Combination Product	02/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part355	02/01/2019	

Labeler - Zeniton Co.,Ltd. (688416831)

Registrant - Zeniton Co.,Ltd. (688416831)

Establishment

Name	Address	ID/FEI	Business Operations
DONG IL PHARMS CO., LTD.		557810721	manufacture(73029-0006)

Revised: 12/2021 Zeniton Co.,Ltd.