UNIVERSAL ADVANCED WHITENING ANTI-CAVITY FLUORIDE- sodium monofluorophosphate paste
Universal Distribution Center LLC

UNIVERSAL ADVANCED WHITENING ANTI-CAVITY FLUORIDE TOOTHPASTE

Active Ingredient

Sodium Monofluorophosphate 0.76%

(1000 ppm)

Purpose

Anticavity toothpaste

Uses

- builds increasing protection against painful sensitivity of teeth due to cold, heat, acids, sweets or contact.
- aids in the prevention of dental cavities.

Warning

When using this product

- if pain\ sensitivity still persists after 4 weeks of use, please visit your dentist.

Stop and ask a dentist

- if the problem persists or worsens.

Sensitivity teeth may indicate a serious problem that may need prompt care by a dentist.

Keep out of reach of children

- If accidentally swallowed more than used for brushing, seek professional help or contact a Poison Control Center immediately.

Directions

- adults and children of 2 years and older: brush teeth thoroughly after meals or at least twice a day, or as directed by a dentist.
- do not swallow.
- to minimize swallowing, use a pea-sized amount in children under 6 years old.
- supervise children's brushing until good habits are established.
- children under 2 years: ask a dentist before use.

Children under 12 years of age: Consult a dentist or doctor

Other information

- store in a cool, dry place.

Inactive Ingredients

Calcium carbonate, Water, Glycerine, Silica, Sodium Lauryl sulphate, Sorbitol, Xanthan gum, Sodium saccharine, Sodium benzoate, Sodium carboxy methyl cellulose, Flavor and FD&C blue #1

PRINCIPAL DISPLAY PANEL

UNIVERSAL ADVANCED WHITENING ANTI-CAVITY FLUORIDE TOOTHPASTE

UNIVERSAL ADVANCED WHITENING ANTI-CAVITY FLUORIDE

sodium monofluorophosphate paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-112
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.76 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
CALCIUM CARBONATE (UNII: H0G9379FGK)	
WATER (UNII: 059QF0KO0R)	

GLYCERIN (UNII: PDC6A3C0OX)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SORBITOL (UNII: 506T60A25R)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-112-01	181 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	05/30/2020	

Labeler - Universal Distribution Center LLC (019180459)

Revised: 11/2023 Universal Distribution Center LLC