DECURE- sodium monofluorophosphate paste, dentifrice Scope medical devices private limited

DeCure Toothpaste

Drug Facts

Active Ingredients:

Sodium Monofluorophosphate 0.76%

Purpose

Anticavity (Total Fluoride Content - 1000 ppm Approx.)

Uses:

Regular brushing with Fluoride toothpaste helps protect teeth roots against cavities.

Warnings:

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

In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

Directions:

Adults and children 2 years and older.	Brush teeth thoroughly after meals or at least twice a day or use as directed by a dentist or physician
Children under 6 yrs.	To minimize swallowing use a pea - sized amount and supervise brushing until good habits are established.
Children under 2 yrs.	Ask a dentist or physician.

Inactive Ingredients:

Calcium Carbonate, Water, Sorbitol, Silica, Sodium Lauryl Sulphate, Flavor, Sodium Silicate, Tetra Sodium Pyrophosphate, Cellulose Gum, Xanthan Gum, Benzyl Alcohol, Sodium Saccharin, Sodium Benzoate

With Fluoride Anticavity Protection

Manufactured For: Scope Medical Devices Pvt. Ltd. 11/10, Mandhour Industrial Area, Ambala -134003, India

Factory Neutral Code DNH/COS/DNH/52 Ref #:330139 Made in India

Exp. Date & Batch No. on crimp.

Packaging

Manufactured For: Scope Medical Devices Pvt. Ltd. DNH/COS/DNH/52 11/10, Mandhour Industrial Area, Ambala -134003, India

Factory Neutral Code Ref #:330139 Made In India





Net Weight 10gms / 0.35oz

Toothpaste

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DECURE

sodium monofluorophosphate paste, dentifrice

Product	Inform	ation
Product	11110111	Ialion

Inactive Ingredients

Product Type HUMAN OTC DRUG Item Code (Source) NDC:84132-100

Route of Administration DENTAL

Active Ingredient/Active Moiety

SACCHARIN SODIUM (UNII: SB8ZUX40TY) **SODIUM BENZOATE** (UNII: OJ245FE5EU)

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	Ingredient Name	Basis of Strength	Strength
	SODIUM MONOFLUOROPHOSPHATE (UNII: C810JCZ56Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	7.6 mg in 1 g

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Ingredient Name	Strength	
CALCIUM CARBONATE (UNII: H0G9379FGK)		
WATER (UNII: 059QF0KO0R)		
SORBITOL (UNII: 506T60A25R)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
SODIUM SILICATE (UNII: IJF18F77L3)		
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)		
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)		
XANTHAN GUM (UNII: TTV12P4NEE)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		

l	Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
		NDC:84132-100- 10	10 g in 1 TUBE; Type 0: Not a Combination Product	05/11/2024			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M021	05/11/2024			