MEDPRIDE- sodium monofluorophosphate paste, dentifrice Shield Line LLC

Drug Facts

Active Ingredient

Sodium monofluorophosphate 0.76% (0.1% w/v fluoride ion)

Purpose

Anticavity toothpaste

Uses

aids in the prevention of dental cavities

Warning

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

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

Directions

■ do not swallow ■ supervise children as necessary until capable of using without supervision ■ instruct children under 12 years of age in good brushing and rinsing habits (to minimize swallowing) ■ adults and children 6 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 of age: do not use unless directed by a doctor or dentist

Inactive ingredients

calcium carbonate, water, sorbitol, hydrated silica, sodium lauryl sulfate, sodium carboxymethyl cellulose, sodium saccharin, sodium methylparaben, sodium propylparaben

Package Label



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- without supervision. instruct children under 12 years of age in good brushing and
- rinsing habits (to minimize swallowing).

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 children under 6 years of age: do not use unless directed
- by a doctor or dentist

Inactive ingredients

Calcium Carbonate, Sorbitol, Treated Water, Silica, Sodium Lauryl Sulphate, Flavour, Sodium Carboxy Methyl Cellulose, Sodium Silicate, Tetra Sodium pyrophosphate, Benzyl Alcohol, Sodium Saccharin, Propyl Paraben



MEDPRIDE

sodium monofluorophosphate paste, dentifrice

Product Information

HUMAN OTC DRUG NDC:52410-9005 **Product Type Item Code (Source)**

Route of Administration DENTAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CARBONATE (UNII: H0G9379FGK)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

METHYLPARABEN SODIUM (UNII: CR6K9C2NHK)	
PROPYLPARABEN (UNII: Z8IX2SC10H)	

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor	MINT	Imprint Code	
Contains			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:52410- 9005-3	42.5 g in 1 TUBE; Type 0: Not a Combination Product	05/27/2013			
2	NDC:52410- 9005-4	78 g in 1 TUBE; Type 0: Not a Combination Product	05/27/2013			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M021	05/27/2013		

Labeler - Shield Line LLC (078518916)

Revised: 3/2024 Shield Line LLC