

**MEDICHOICE FLUORIDE- sodium monofluorophosphate paste, dentifrice
O&M HALYARD, INC.**

MediChoice Fluoride Toothpaste

Drug Facts

ACTIVE INGREDIENT

Sodium Monofluorophosphate - 0.76% (0.15% w/v fluoride ion)

PURPOSE

Anticavity

USE

Aids in the prevention of dental cavities

WARNINGS

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

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

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 2 to 6 years of age should use only a pea-sized amount in order to minimize swallowing. Supervise children and help them create good brushing and rinsing habits until capable of using without supervision.
- Children under 2 years of age: Consult a dentist or a doctor.

INACTIVE INGREDIENTS

CALCIUM CARBONATE, WATER, SORBITOL, PRECIPITATED SILICA, SODIUM LAURYL SULFATE, SODIUM CARBOXY METHYL CELLULOSE, SODIUM SILICATE TETRA SODIUM PYROPHOSPHATE, SODIUM SACCHARIN, METHYL PARABEN, TITANIUM DIOXIDE POLYETHYLENE GLYCOL 400, PROPYL PARABEN

NDC 39892-0602-1



FLUORIDE TOOTHPASTE MINT



Net wt. 0.6 oz. (17g) Reorder: PC7306

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INACTIVE INGREDIENTS calcium carbonate, water, sorbitol, precipitated silica, sodium lauryl sulfate, flavor, sodium carboxy methyl cellulose, sodium silicate, tetra sodium pyrophosphate, sodium saccharin, methylparaben, titanium dioxide, polyethylene glycol 400, propylparaben	
Distributed by Owens & Minor, 9120 Lockwood Boulevard, Mechanicsville, VA 23116 Made in India	

1222101395 Rev. B

C.No. DNH/COS/DNH/52 Exp. Date & Lot No. on Crimp



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MEDICHOICE FLUORIDE

sodium monofluorophosphate paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:39892-0602
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)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
SODIUM SILICATE (UNII: IJF18F77L3)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	MINT (Mint)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:39892-0602-1	17 g in 1 TUBE; Type 0: Not a Combination Product	03/01/2013	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M021	03/01/2013	

Labeler - O&M HALYARD, INC. (081057389)

Registrant - Owens & Minor Inc. (847412269)

Revised: 10/2025

O&M HALYARD, INC.