

**MEDICHOICE PREMIUM ANTICAVITY FLUORIDE- sodium fluoride paste,
dentifrice
O&M HALYARD, INC.**

MediChoice Premium Anticavity Fluoride Toothpaste

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Drug Facts

ACTIVE INGREDIENT

Sodium Fluoride - 0.243% (0.15% w/v fluoride ion)

PURPOSE

Anticavity

USE

Aids in the prevention of dental cavities

WARNINGS

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

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

sorbitol, silica, water, glycerin, polythelene glycol 1500, sodium lauryl sulfate, flavor, titanium dioxide, cellulose gum, carrageenan, trisodium orthophosphate, sodium saccharin, sodium benzoate, tetrasodium pyrophosphate, methylparaben, propylparaben, FD & C blue #1

Made in India, Distributed by Owens & Minor, 9120 Lockwood Boulevard, Mechanicsville, VA 23116

MediChoice Premium Anticavity Fluoride Toothpaste 24g (39892-0603-2)

NDC 39892-0603-2

MEDI CHOICE®

Premium Anticavity Fluoride Toothpaste

cool mint
 fights cavities
smooth texture
gentle formula

Reorder No. PC7385
NET WT. 0.85 OZ.
(24 GRAMS)

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

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MEDICHOICE PREMIUM ANTICAVITY FLUORIDE

sodium fluoride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:39892-0603
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	2.43 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
SORBITOL (UNII: 506T60A25R)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
CARRAGEENAN (UNII: 5C69YCD2YJ)	
SODIUM PHOSPHATE, TRIBASIC, DODECAHYDRATE (UNII: B70850QPHR)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:39892-0603-2	24 g in 1 TUBE; Type 0: Not a Combination Product	03/01/2013	

Marketing Information

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

Labeler - O&M HALYARD, INC. (081057389)**Registrant** - Owens & Minor, Inc (847412269)

Establishment

Name	Address	ID/FEI	Business Operations
Dabur India Limited		650599231	manufacture(39892-0603)

Revised: 10/2025

O&M HALYARD, INC.