# MEDICHOICE ANTICAVITY FLUORIDE - sodium fluoride paste, dentifrice Owens and Minor

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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### MEDICHOICE ANTICAVITY FLUORIDE TOOTHPASTE

### **Active Ingredient**

Sodium Fluoride 0.22%

### **Purpose**

Anticavity

### Use:

Aids in the prevention of dental cavities.

### **Directions**

Adults and children 2 yrs 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 years: Ask a dentist or physician.

### Warning

If you accidentally swallow more than used for brushing seek professional assistance or contact a Poison Control Center Immediately.

### Keep out of reach of children

under 6 years of age.

### **Inactive Ingredients**

**Inactive Ingredients:** 

Calcium Carbonate, Water, Sorbitol, Sodium Lauryl Sulfate, Silica, Carboxymethyl Cellulose, Flavor, Potassium Nitrate, Sodium benzoate, Sodium saccharin, menthol

Expiration Date: Expiration Date and Batch No. on crimp of Tube

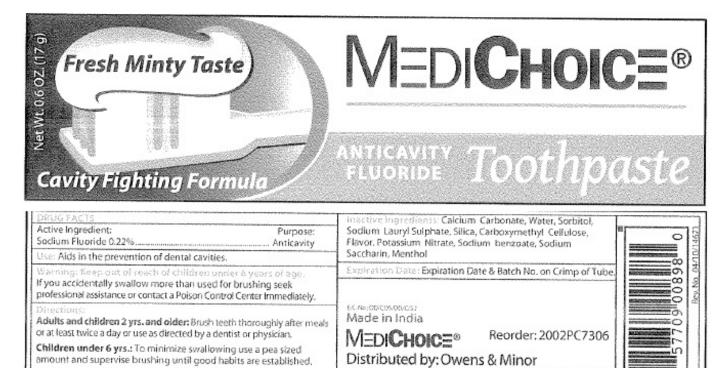
E.c No.: DD/COS/DD/C/52

Made in India

MediChoice Reorder: 2002PC7306

Distributed by: Owens and Minor

### **Product Tube Label**



Mechanicsville, VA 23116 • 804.723.7000

**Product Outer Packaging 144ct** 

Children Under 2 years: Ask a dentist or physician.

# NEDICHOICE® Reorder: 2002PC7306

Anticavity Fluoride Toothpaste 0.6 oz

Contents: 144/bx

Distributed by:

Owens & Minor Mechanicsville, VA 23116 Made in India



20757709008984

E.C. No.: DD/COS/DD/C/52 Rev No. 4/10/14672

### MEDICHOICE ANTICAVITY FLUORIDE

sodium fluoride paste, dentifrice

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:39892-0601

Route of Administration DENTAL

**Active Ingredient/Active Moiety** 

Ingredient Name Basis of Strength Strength

Inactive Ingredients				
Ingredient Name	Strength			
CALCIUM CARBONATE (UNII: H0 G9 379 FGK)				
WATER (UNII: 059QF0KO0R)				
SORBITOL (UNII: 506T60A25R)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)				
CARBO XYMETHYLCELLULO SE (UNII: 05JZI7B19X)				
PO TASSIUM NITRATE (UNII: RU45X2JN0 Z)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
MENTHOL (UNII: L7T10EIP3A)				

Packaging						
# Item C	o de	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:39892-06	501-1 5 in	1 CASE				
1	144	in 1 BOX				
1	17 ε	in 1 TUBE				
2 NDC:39892-06	501-2 5 in	1 CASE				
2	144	in 1 BOX				
2	24 ફ	g in 1 TUBE				

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part355	05/18/2010				

## Labeler - Owens and Minor (847412269)

Revised: 10/2011 Owens and Minor