

**EVO SENSITIVE WHITENING ANTI-CAVITY FLUORIDE- sodium  
fluoride mouthwash  
Ashtel Studios, Inc.**

-----  
**evo<sup>®</sup> SENSITIVE WHITENING ANTI-CAVITY FLUORIDE MOUTHWASH**

***Drug Facts***

***Active ingredient***

Sodium Fluoride 0.05%

***Purpose***

Anticavity

***Uses***

- Helps reduce Plaque-associated Gingivitis
- Helps control gum infections

***Warnings***

**Stop use and ask dentist if**

- Gingivitis, bleeding, or redness persists for more than 2 weeks.
- You have painful or swollen gums, pus, loose teeth, or increased spacing between teeth. These may be signs of periodontitis.

**Keep out of reach of children under 6 years of age.** If more than the rinsing amount is swallowed, seek medical help or contact a Poison Control Center right away.

***Directions***

**• Adults and children 12 years and older:**

Adults and children 12 years and older: Use 10 mL (2 teaspoons) undiluted. Swish vigorously for 1 minute, then spit out. Use once a day.

**• Children 6 years to under 12 years:**

Should be supervised when using this product.

**• Children under 6 years of age:**

Not recommended for use.

- Do not swallow.

***Other information***

- Do not exceed the recommended dosage.
- This rinse is not a substitute for brushing or flossing.
- Do not use if safety seal is broken or missing.

### ***Inactive ingredients***

Water, Sorbitol, Glycerin, Xylitol, Sodium Benzoate, Potassium Sorbate, Peg-40 Hydrogenated Castor Oil, Mentha Haplocalyx Oil, Sodium Saccharin, Cetylpyridinium Chloride, Methyl Diisopropyl Propionamide, Sucralose, Citric Acid

### ***Questions or comments?***

Call toll free **1-877-274-8358**

### **COOL MINT**

HELPS PREVENT CAVITIES

STRENGTHS ENAMEL

CLEANS TEETH

FRESHENS BREATH

**MADE IN CHINA**

**GETEVO.NET**

1-909-434-0911 INTERNATIONAL

PATENTS, COPYRIGHTS AND TRADEMARKS GRANTED OR PENDING WORLDWIDE

DISTRIBUTED BY ASHTEL STUDIOS INC., ONTARIO, CALIFORNIA 91761

### **Packaging**



## EVO SENSITIVE WHITENING ANTI-CAVITY FLUORIDE

sodium fluoride mouthwash

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70108-210
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>SODIUM FLUORIDE</b> (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	SODIUM FLUORIDE	0.05 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>XYLITOL</b> (UNII: VCQ006KQ1E)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	
<b>POLYOXYL 40 HYDROGENATED CASTOR OIL</b> (UNII: 7YC686GQ8F)	
<b>MENTHA CANADENSIS TOP OIL</b> (UNII: N2285224W8)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>CETYLPYRIDINIUM CHLORIDE</b> (UNII: D9OM4SK49P)	
<b>METHYL DIISOPROPYL PROPIONAMIDE</b> (UNII: 6QOP5A9489)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	

## Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	MINT	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70108-210-01	296 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/10/2025	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	07/10/2025	

**Labeler -** Ashtel Studios, Inc. (148689180)

Revised: 7/2025

Ashtel Studios, Inc.