

**CREST PRO HEALTH DENSIFY RINSE- sodium fluoride rinse**  
**The Procter & Gamble Manufacturing Company**

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**Crest Pro Health Densify Rinse**

***Drug Facts***

**Active ingredient**

Sodium Fluoride 0.02% (0.01% w/v fluoride ion)

**Purpose**

Anticavity

**Use**

Aids in the prevention of dental cavities

**Warning**

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

**Directions**

- Adults and children 6 years & older:
  - Use twice a day after brushing your teeth with a toothpaste.
  - Vigorously swish 10 mL (2 teaspoonfuls) of rinse between your teeth for 1 minute and then spit out.
  - Do not swallow the rinse.
  - Do not eat or drink for 30 minutes after rinsing.
  - Instruct children under 12 years of age in good rinsing habits (to minimize swallowing).
  - Supervise children as necessary until capable of using without supervision.
- Children under 6 years of age: consult a dentist or doctor.

**Inactive ingredients**

water, propylene glycol, poloxamer 407, polysorbate 80, flavor, benzoic acid, sucralose, sodium benzoate, phosphoric acid, disodium phosphate

**Questions?**

1-800-285-9139

DISTR. BY PROCTER & GAMBLE,

CINCINNATI, OH 45202

## PRINCIPAL DISPLAY PANEL - 946 mL Bottle Label

**Crest®**

PRO HEALTH™

ANTICAVITY FLUORIDE RINSE

DENSIFY™

ACTIVELY REBUILDS TOOTH DENSITY

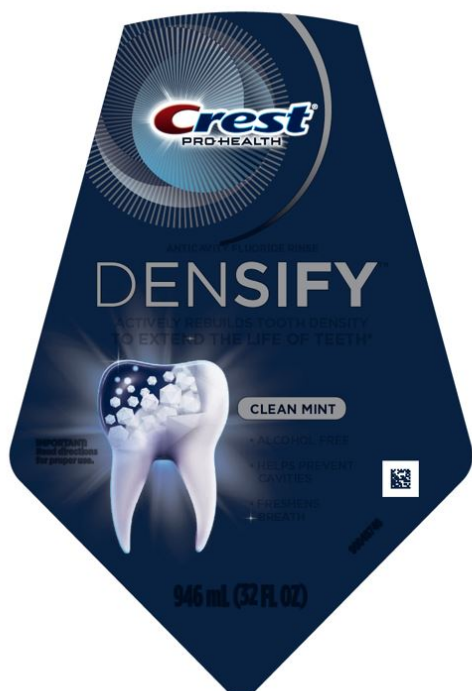
TO EXTEND THE LIFE OF TEETH\*

CLEAN MINT

- ALCOHOL FREE
- HELPS PREVENT CAVITIES
- FRESHENS BREATH

IMPORTANT: Read directions for proper use.

946 mL (32 FL OZ)



## CREST PRO HEALTH DENSIFY RINSE

sodium fluoride rinse

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69423-882
<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:Q80VPU4080)	FLUORIDE ION	0.1 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>BENZOIC ACID</b> (UNII: 8SKN0B0MIM)	
<b>PHOSPHORIC ACID</b> (UNII: E4GA8884NN)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>POLOXAMER 407</b> (UNII: TUF2IWW3M2)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS</b> (UNII: 22ADO53M6F)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	

### 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:69423-882-32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2022	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	01/01/2022	

**Labeler** - The Procter & Gamble Manufacturing Company (004238200)