

**CREST 3D WHITE LUXE DIAMOND STRONG- sodium fluoride rinse**  
**The Procter & Gamble Manufacturing Company**

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**Crest® 3D White Luxe™**

**Diamond Strong**

***Drug Facts***

**Active ingredient**

Sodium Fluoride 0.0219% (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 12 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 12 years of age: Consult a dentist or doctor.

**Inactive ingredients**

water, glycerin, propylene glycol, hydrogen peroxide, poloxamer 407, flavor, sodium saccharin, phosphoric acid, sucralose, disodium phosphate

**Questions?**

**1-800-285-9139**

**DISTR. BY PROCTER & GAMBLE,  
CINCINNATI, OH 45202**

**PRINCIPAL DISPLAY PANEL - 473 mL Bottle Label**

***WITH FLUORIDE  
ALCOHOL FREE***

***Crest***®

**3D WHITE**

ANTICAVITY FLUORIDE MOUTHWASH

**DIAMOND STRONG**

**FROM THE MAKERS OF WHITESTRIPS**®

- **STRENGTHENS** ENAMEL
- WHITENS TEETH
- HELPS PREVENT CAVITIES
- 91518226

**WINTERMINT**

IMPORTANT: Read directions for proper use.

946 mL

(32 FL OZ)



## CREST 3D WHITE LUXE DIAMOND STRONG

sodium fluoride rinse

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:37000-862
<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>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C00X)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>HYDROGEN PEROXIDE</b> (UNII: BBX060AN9V)	
<b>POLOXAMER 407</b> (UNII: TUF21VW3M2)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>PHOSPHORIC ACID</b> (UNII: E4GA8884NN)	
<b>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS</b> (UNII: 22ADO53M6F)	

**Product Characteristics**

<b>Color</b>	white (Clear)	<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:37000-862-01	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/19/2015	
2	NDC:37000-862-02	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/19/2015	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	02/19/2015	

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

Revised: 3/2025

The Procter &amp; Gamble Manufacturing Company