

**CREST COMPLETE PLUS SCOPE OUTLAST- sodium fluoride gel, dentifrice**  
**The Procter & Gamble Manufacturing Company**

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**Crest Complete Plus Scope Outlast Liquid Gel**

***Drug Facts***

**Active ingredient**

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

**Purpose**

Anticavity toothpaste

**Use**

helps protect against cavities

**Warnings**

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

**Directions**

- adults and children 2 yrs. & older: brush teeth thoroughly after meals or at least twice a day or use as directed by a dentist
  - do not swallow
  - to minimize swallowing use a pea-sized amount in children under 6
  - supervise children's brushing until good habits are established
- children under 2 yrs.: ask a dentist

**Inactive ingredients**

sorbitol, water, hydrated silica, alcohol (1.4%), sodium lauryl sulfate, flavor, glycerin, cellulose gum, trisodium phosphate, sodium phosphate, sodium saccharin, carbomer, polysorbate 80, sodium benzoate, cetylpyridinium chloride, blue 1, yellow 5

**Questions?**

**1-800-492-7378**

Dist. by Procter & Gamble, Cincinnati, OH 45202

## PRINCIPAL DISPLAY PANEL - 130 g Bottle Label

scope®

OUTLAST®

Crest® plus  
complete

ANTICAVITY FLUORIDE TOOTHPASTE  
minty fresh  
liquid gel

NET WT 4.6 OZ (130 g)



0 37000 71161 2

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<b>Directions</b> • adults and children 2 yrs. & older: brush teeth thoroughly after meals or at least twice a day or use as directed by a dentist • do not swallow • to minimize swallowing use a pea-sized amount in children under 6 • supervise children's brushing until good habits are established • children under 2 yrs.: ask a dentist	
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<b>Questions?</b> 1-800-492-7378	91411989
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## CREST COMPLETE PLUS SCOPE OUTLAST

sodium fluoride gel, dentifrice

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:37000-983
<b>Route of Administration</b>	DENTAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>SODIUM FLUORIDE</b> (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.5 mg in 1 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>HYDRATED SILICA</b> (UNII: Y6O7T4G8P9)	
<b>SODIUM PHOSPHATE, TRIBASIC, ANHYDROUS</b> (UNII: SX01TZO3QZ)	
<b>SODIUM PHOSPHATE</b> (UNII: SE337SVY37)	
<b>CARBOXYPOLYMETHYLENE</b> (UNII: 0A5MM307FC)	
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>CARBOXYMETHYLCELLULOSE SODIUM</b> (UNII: K679OBS311)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>CETYLPIRIDINIUM CHLORIDE</b> (UNII: D9OM4SK49P)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	

**Product Characteristics**

<b>Color</b>	green	<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:37000-983-04	130 g in 1 BOTTLE; Type 0: Not a Combination Product	12/31/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	12/31/2020	

Revised: 10/2023

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