

**CREST COMPLETE MULTI-BENEFIT PLUS SCOPE- sodium fluoride gel,
dentifrice**

The Procter & Gamble Manufacturing Company

Crest®

complete

MULTI-BENEFIT™

Drug Facts

Active ingredient

Sodium fluoride 0.243%

Purpose

Anticavity toothpaste (0.15% w/v fluoride ion)

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, trisodium phosphate, flavor, glycerin, cellulose gum, sodium phosphate, sodium saccharin, carbomer, polysorbate 80, sodium benzoate, cetylpyridinium chloride, benzoic acid, blue 1, yellow 5

Questions?

1-800-492-7378

Dist. by Procter & Gamble, Cincinnati, OH 45202

PRINCIPAL DISPLAY PANEL - 130 g Bottle Label

scope

+ whitening

Crest

Complete plus

ANTICAVITY FLUORIDE TOOTHPASTE

minty fresh

liquid gel

91378648

NET WT 4.6 OZ (130 g)



CREST COMPLETE MULTI-BENEFIT PLUS SCOPE

sodium fluoride gel, dentifrice

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:37000-822 |
| Route of Administration | DENTAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|---------------|
| SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080) | FLUORIDE ION | 1.1 mg in 1 g |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| SORBITOL (UNII: 506T60A25R) | |
| WATER (UNII: 059QF0KO0R) | |
| HYDRATED SILICA (UNII: Y6O7T4G8P9) | |
| SODIUM LAURYL SULFATE (UNII: 368GB5141J) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) | |
| SODIUM PHOSPHATE (UNII: SE337SVY37) | |
| SACCHARIN SODIUM (UNII: SB8ZUX40TY) | |
| POLYSORBATE 80 (UNII: 6OZP39ZG8H) | |
| SODIUM BENZOATE (UNII: OJ245FE5EU) | |
| CETYLPIRIDINIUM CHLORIDE (UNII: D9OM4SK49P) | |
| BENZOIC ACID (UNII: 8SKN0B0MIM) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| FD&C YELLOW NO. 5 (UNII: I753WB2F1M) | |
| ALCOHOL (UNII: 3K9958V90M) | |

Product Characteristics

| | | | |
|-----------------|------------|---------------------|--|
| Color | white | Score | |
| Shape | | Size | |
| Flavor | PEPPERMINT | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:37000-822-04 | 130 g in 1 BOTTLE; Type 0: Not a Combination Product | 07/01/2011 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M021 | 07/01/2011 | |

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 1/2026

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