

CREST SENSITIVITY WHITENING PLUS SCOPE- sodium fluoride and potassium nitrate paste, dentifrice
The Procter & Gamble Manufacturing Company

Crest®
Sensitivity
Whitening Plus Scope

Drug Facts

| Active ingredients | Purpose |
|--|----------------------------------|
| Potassium nitrate 5% | Toothpaste for sensitive teeth |
| Sodium fluoride 0.243% (0.14% w/v fluoride ion) | Toothpaste for cavity prevention |

Uses

- when used regularly, builds increasing protection against painful sensitivity of the teeth to cold, heat, acids, sweets or contact
- aids in the prevention of cavities

Warnings

When using this product do not use longer than four weeks unless recommended by a dentist.

Stop use and ask a dentist if problem lasts or gets worse. Sensitive teeth may indicate a serious problem that may need prompt care.

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

Directions

- do not swallow
- 12 yrs. & older: Apply at least a 1-inch strip of the product onto a soft bristle toothbrush. Brush teeth thoroughly for at least 1 minute twice a day (morning and evening) or as recommended by a dentist. Make sure to brush all sensitive areas of the teeth.
- do not use in children under 12 yrs.

Inactive ingredients

water, hydrated silica, glycerin, sorbitol, trisodium phosphate, sodium lauryl sulfate, flavor, cellulose gum, alcohol (1.09%), xanthan gum, sodium saccharin, polysorbate 80, sodium benzoate, cetylpyridinium chloride, benzoic acid, titanium dioxide, blue 1

Questions?

1-800-492-7378

Dist. by Procter & Gamble, Cincinnati, OH 45202

PRINCIPAL DISPLAY PANEL - 170 g Tube Carton

Crest®

PRO-HEALTH

SENSITIVITY

TOOTHPASTE FOR SENSITIVE TEETH AND CAVITY PREVENTION

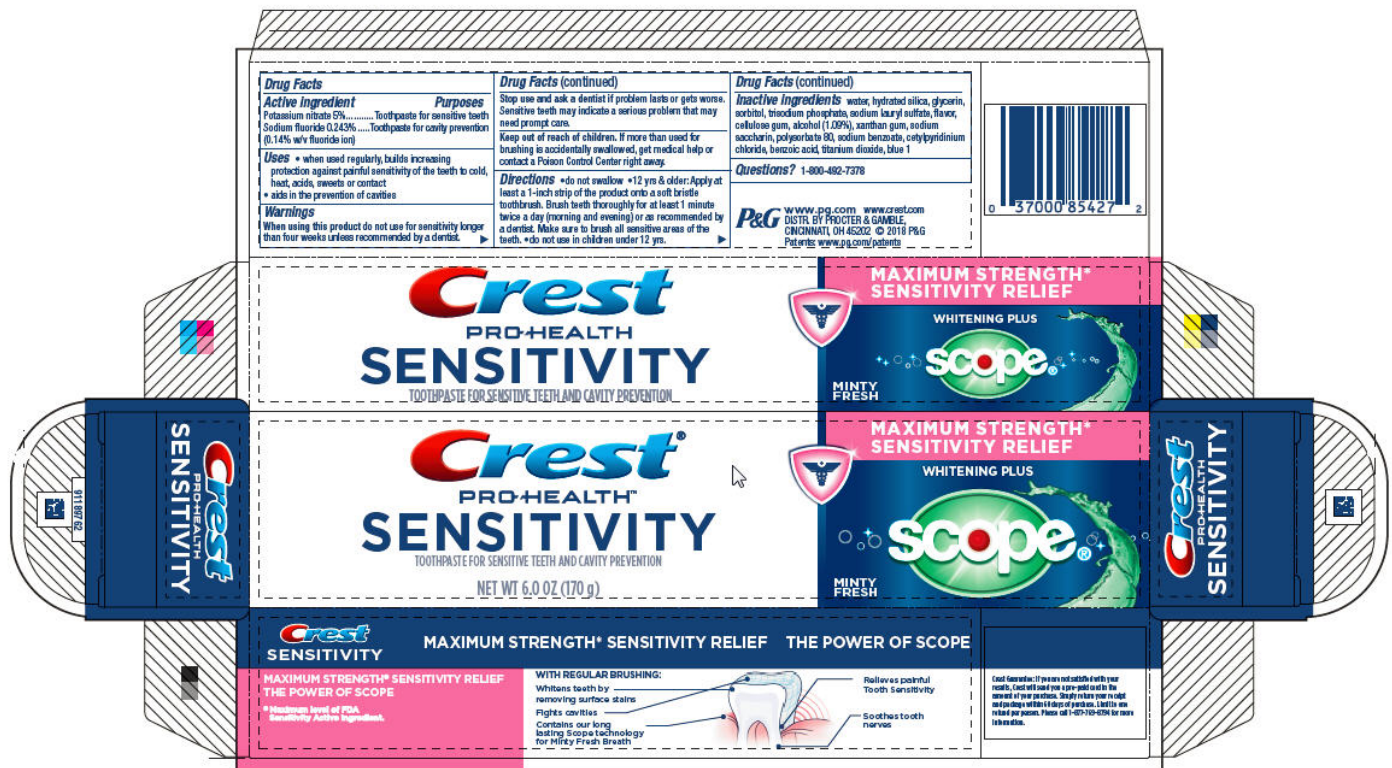
NET WT 6.0 OZ (170 g)

**MAXIMUM STRENGTH*
SENSITIVITY RELIEF**

WHITENING PLUS

scope®

MINTY FRESH



CREST SENSITIVITY WHITENING PLUS SCOPE

sodium fluoride and potassium nitrate paste, dentifrice

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:37000-898 |
| Route of Administration | DENTAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|---------------|
| POTASSIUM NITRATE (UNII: RU45X2JN0Z) (NITRATE ION - UNII:T93E9Y2844) | POTASSIUM NITRATE | 50 mg in 1 g |
| SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080) | FLUORIDE ION | 1.4 mg in 1 g |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| CETYLPIRIDINIUM CHLORIDE (UNII: D9OM4SK49P) | |
| BENZOIC ACID (UNII: 8SKN0B0MIM) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| SODIUM PHOSPHATE, TRIBASIC, ANHYDROUS (UNII: SX01TZ03QZ) | |
| WATER (UNII: 059QF0KO0R) | |
| HYDRATED SILICA (UNII: Y6O7T4G8P9) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| SORBITOL (UNII: 506T60A25R) | |
| SODIUM LAURYL SULFATE (UNII: 368GB5141J) | |
| CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) | |
| ALCOHOL (UNII: 3K9958V90M) | |
| XANTHAN GUM (UNII: TTV12P4NEE) | |
| SACCHARIN SODIUM (UNII: SB8ZUX40TY) | |
| POLYSORBATE 80 (UNII: 6OZP39ZG8H) | |
| SODIUM BENZOATE (UNII: OJ245FE5EU) | |

Product Characteristics

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

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:37000-898-41 | 1 in 1 CARTON | 01/01/2018 | 06/18/2019 |
| 1 | | 116 g in 1 TUBE; Type 0: Not a Combination Product | | |
| 2 | NDC:37000-898-06 | 1 in 1 CARTON | 01/01/2018 | |
| 2 | | 170 g in 1 TUBE; Type 0: Not a Combination Product | | |
| 3 | NDC:37000-898-12 | 2 in 1 CARTON | 01/01/2018 | 12/17/2024 |

| 3 | 170 g in 1 TUBE; Type 0: Not a Combination Product | | |
|------------------------------|--|----------------------|--------------------|
| Marketing Information | | | |
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC Monograph Drug | M021 | 01/01/2018 | |

Labeler - The Procter & Gamble Manufacturing Company (004238200)

| Establishment | | | |
|----------------------|---------|-----------|----------------------------|
| Name | Address | ID/FEI | Business Operations |
| Jost Chemical Co. | | 147882294 | api manufacture(37000-898) |

Revised: 1/2025

The Procter & Gamble Manufacturing Company