

CREST SCOPE ADVANCED- sodium fluoride rinse
The Procter & Gamble Manufacturing Company

Crest® Scope® Advanced™

Drug Facts

Active ingredient

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

Purpose

Anticavity

Use

Aids in the prevention of dental cavities

Warnings

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 (2teaspoonfuls) 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, alcohol (15 wt%), propylene glycol, flavor, benzoic acid, poloxamer 407, cetylpyridinium chloride, sucralose, phosphoric acid, disodium phosphate, yellow 5, blue 1

Questions?

1-800-862-7442

DISTR. BY PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 1000mL Bottle Label

Crest®

ANTICAVITY FLUORIDE MOUTHWASH

scope®

ADVANCED™

MULTI-ACTION FORMULA

+FLUORIDE

- HELPS PREVENT CAVITIES
- FORTIFIES ENAMEL
- FRESHENS BREATH
- CLEANS WHOLE MOUTH

IMPORTANT: Read directions for proper use.

SEE BACK FOR IMPORTANT INGREDIENT INFORMATION.

1L (33.8 FL OZ)



CREST SCOPE ADVANCED

sodium fluoride rinse

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:37000-208 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|--------------------------|-----------------|
| SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O) | FLUORIDE ION | 0.1 mg in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|--|-----------------|
| FD&C YELLOW NO. 5 (UNII: I753WB2F1M) | |
| CETYLPIRIDINIUM CHLORIDE (UNII: D9OM4SK49P) | |
| SUCRALOSE (UNII: 96K6UQ3ZD4) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| PHOSPHORIC ACID (UNII: E4GA8884NN) | |
| ALCOHOL (UNII: 3K9958V90M) | |
| BENZOIC ACID (UNII: 8SKN0B0MIM) | |
| POLOXAMER 407 (UNII: TUF2IVW3M2) | |
| WATER (UNII: 059QF0KO0R) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F) | |

Product Characteristics

| | | | |
|-----------------|-------|---------------------|--|
| Color | green | Score | |
| Shape | | Size | |
| Flavor | MINT | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:37000-208-05 | 500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 01/01/2019 | |
| 2 | NDC:37000-208-10 | 1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 01/01/2019 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------------|---|-----------------------------|---------------------------|
| OTC Monograph Drug | M021 | 01/01/2019 | |

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 10/2023

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