

CREST PRO-HEALTH GUM DETOXIFY WINTER MINT- cetylpyridinium chloride rinse

The Procter & Gamble Manufacturing Company

Crest Pro-Health Gum Detoxify Winter Mint

Drug Facts

Active ingredient

Cetylpyridinium chloride 0.07%

Purpose

Antigingivitis/antiplaque

Uses

- helps prevent and reduce plaque and gingivitis
- helps control plaque bacteria that contribute to the development of gingivitis and bleeding gums

Warnings

Ask a dentist if symptoms persist or condition worsens after regular use.

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

Directions

- use after your normal brushing and flossing routine; rinse toothpaste from mouth prior to use
- adults and children 6 yrs. & older: Rinse for 30 seconds with 20 mL (4 teaspoonfuls) twice a day
 - do not swallow
 - children 6 years to under 12 years of age: supervise use
- children under 6 years of age: do not use

Inactive ingredients

water, glycerin, propylene glycol, flavor, poloxamer 407, sodium benzoate, sodium saccharin, sucralose, benzoic acid, blue 1, yellow 6

Questions?

1-800-285-9139

DISTR. BY PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 1 L Bottle Label

Crest®

Pro Health

CPC ANTIGINGIVITIS/ANTIPLAQUE ORAL RINSE

GUM

DETOXIFY

24 HR Protection

WITH ADVANCED

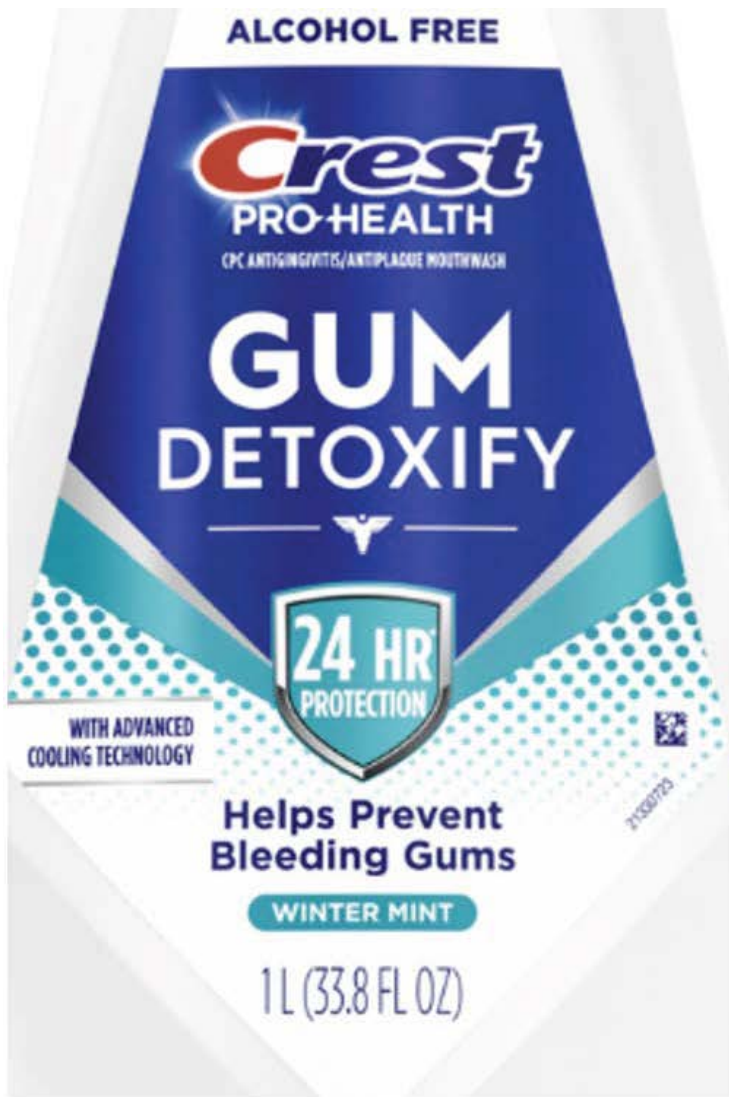
COOLING TECHNOLOGY

Helps Prevent

Bleeding Gums

WINTER MINT

1 L (33.8 FL OZ)



| CREST PRO-HEALTH GUM DETOXIFY WINTER MINT | | | |
|--|---|---------------------------|-----------------|
| cetylpyridinium chloride rinse | | | |
| Product Information | | | |
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:84126-648 |
| Route of Administration | ORAL | | |
| Active Ingredient/Active Moiety | | | |
| | Ingredient Name | Basis of Strength | Strength |
| | CETYLPIRIDINIUM CHLORIDE (UNII: D9OM4SK49P) (CETYLPIRIDINIUM - UNII: CUB7J10JV3) | CETYLPIRIDINIUM CHLORIDE | 0.07 g in 1 mL |
| Inactive Ingredients | | | |
| | Ingredient Name | | Strength |
| | SODIUM BENZOATE (UNII: OJ245FE5EU) | | |
| | SUCRALOSE (UNII: 96K6UQ3ZD4) | | |
| | BENZOIC ACID (UNII: 8SKN0B0MIM) | | |

| | |
|---|--|
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |
| POLOXAMER 407 (UNII: TUF2IVW3M2) | |
| WATER (UNII: 059QF0KO0R) | |
| GLYCERIN (UNII: PDC6A3C0OX) | |
| SACCHARIN SODIUM (UNII: SB8ZUX40TY) | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| FD&C YELLOW NO. 6 (UNII: H77VEI93A8) | |

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:84126-648-01 | 1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 01/01/2026 | |
| 2 | NDC:84126-648-05 | 500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 01/01/2026 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M022 | 01/01/2026 | |

Labeler - The Procter & Gamble Manufacturing Company (004238200)