CREST COMPLETE MULTI BENEFIT WHITENING PLUS SCOPE MINTY FRESH GEL-
sodium fluoride paste, dentifrice
Procter & Gamble Manufacturing Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Crest ®
Complete Multi-Benefit
Whitening + Scope® Minty Fresh 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, disodium pyrophosphate, sodium lauryl sulfate, flavor, sodium hydroxide, alcohol (0.7%), xanthan gum, sodium saccharin, glycerin, carbomer, cellulose gum, 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 - 119 g pump label

Crest®

complete

MULTI-BENEFIT

FLUORIDE ANTICAVITY TOOTHPASTE

WHITENING+

scope®

KILLS MILLIONS OF

BAD BREATH GERMS*

MINTY

FRESH GEL

NET WT 4.2 OZ (119 g)

CREST COMPLETE MULTI BENEFIT WHITENING PLUS SCOPE MINTY FRESH GEL

sodium fluoride paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37000-494
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
SODIUM ACID PYROPHOSPHATE (UNII: H5WVD9LZUD)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALCOHOL (UNII: 3K9958V90M)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	

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:37000-494-42	119 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part355	08/01/2017	

Labeler - Procter & Gamble Manufacturing Company (004238200)

Revised: 5/2023 Procter & Gamble Manufacturing Company