

CREST COMPLETE PLUS SCOPE OUTLAST ULTRA- sodium fluoride paste, dentifrice

The Procter & Gamble Manufacturing Company

Crest Complete Plus Scope Outlast Ultra

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, flavor, sodium lauryl sulfate, sodium hydroxide, alcohol (0.7%), sodium saccharin, xanthan gum, glycerin, poloxamer 407, carbomer, polysorbate 80, sodium benzoate, cetylpyridinium chloride, benzoic acid, titanium dioxide, blue 1, yellow 5

Questions?

1-800-492-7378

DISTR. BY PROCTER & GAMBLE,
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 184 g Tube Carton

Crest

complete plus

PREMIUM

NET WT 6.5 OZ (184 g)

ANTICAVITY FLUORIDE TOOTHPASTE

scope®

OUTLAST®

ULTRA

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 usual for brushing is accidentally swallowed, get medical help or contact a Poison Control Center right away.	
Directions • adults and children 7 yrs. & older: brush teeth thoroughly after meals or at least twice a day or less 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	
Drug Facts (continued)	
Inactive ingredients sorbitol, water, hydrated silica, disodium pyrophosphate, flavor, sodium lauryl sulfate, sodium hydroxide, alcohol (0.7%), sodium saccharin, xanthan gum, glycerin, potassium citrate, polyvinylpyrrolidone 40, sodium benzoate, cetylpyridinium chloride, benzal acid, titanium dioxide, blue 1, yellow 5	
Questions? 1-800-488-7777	
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For best results, brush your tongue as part of your routine.	



CREST COMPLETE PLUS SCOPE OUTLAST ULTRA

sodium fluoride paste, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69423-918
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	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)				
POLOXAMER 407 (UNII: TUF2IVW3M2)				
SORBITOL (UNII: 506T60A25R)				
WATER (UNII: 059QF0KO0R)				
HYDRATED SILICA (UNII: Y6O7T4G8P9)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
ALCOHOL (UNII: 3K9958V90M)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				
GLYCERIN (UNII: PDC6A3C0OX)				
POLYSORBATE 80 (UNII: 6OZP39ZG8H)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)				
BENZOIC ACID (UNII: 8SKN0B0MIM)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
SODIUM ACID PYROPHOSPHATE (UNII: H5WWD9LZUD)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
XANTHAN GUM (UNII: TTV12P4NEE)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
Product Characteristics				
Color	white	Score		
Shape		Size		
Flavor	MINT	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69423-918-65	1 in 1 CARTON	01/05/2021	
1		184 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69423-918-63	1 in 1 CARTON	01/05/2021	
2		178 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	01/05/2021	
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Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 2/2024

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