

CREST CAVITY PROTECTION COOL MINT - sodium fluoride gel, dentifrice
A-S Medication Solutions

CREST CAVITY PROTECTION COOL MINT

Drug Facts

Active ingredient

Sodium fluoride 0.243% (0.15% w/v fluoride ion)

Purpose

Anticavity toothpaste

Use

helps protect teeth and roots 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, sodium lauryl sulfate, trisodium phosphate, cellulose gum, flavor, sodium phosphate, sodium saccharin, carbomer, blue 1

Questions?

1-800-492-7378

Dist. by Procter & Gamble, Cincinnati OH 45202

www.crest.com

HOW SUPPLIED

Product: 50090-6233

NDC: 50090-6233-0 161 g in a TUBE / 24 in a CASE

Sodium Fluoride



CREST CAVITY PROTECTION COOL MINT

sodium fluoride gel, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-6233(NDC:37000-025)
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
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM PHOSPHATE, TRIBASIC (UNII: A752Q30A6X)	
SORBITOL (UNII: 506T60A25R)	
WATER (UNII: 059QF0KO0R)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	

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:50090-6233-0	24 in 1 CASE	11/18/2022	
1		161 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	06/16/1995	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-6233)

Revised: 6/2024

A-S Medication Solutions