

COLGATE CAVITY PROTECTION- sodium fluoride gel, dentifrice
Mission Hills S.A de C.V

Colgate® Cavity Protection Gel

Drug Facts

Active ingredient

Sodium fluoride 0.24% (0.15% w/v fluoride ion)

Purpose

Anticavity

Use

helps protect against cavities

Warnings

Keep out of reach of children under 6 years 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 years of age and older	brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or physician
children 2 to 6 years	use only a pea sized amount and supervise child's brushing and rinsing (to minimize swallowing)
children under 2 years	ask a dentist or physician

Inactive ingredients

sorbitol, water, hydrated silica, sodium lauryl sulfate, flavor, PEG-12, cellulose gum, sodium saccharin, FD&C blue no. 1, FD&C yellow no. 5

Questions?

1-800-468-6502

Dist. by:

COLGATE-PALMOLIVE COMPANY
New York, NY 10022

PRINCIPAL DISPLAY PANEL - 119 g Tube Label

***Cavity
Protection
Gel***

*Helps
Strengthen
Teeth*

Colgate®
Fluoride Toothpaste

ADA
*Accepted
American
Dental
Association*

Toothpaste

*NET WT
4.2 OZ
(119 g)*

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TWIST OFF CAP AND REMOVE FOIL SEAL. DO NOT USE IF SEAL IS NOT INTACT.

ADA Accepted
The ADA Council on Scientific Affairs' Acceptance of Colgate Cavity Protection toothpaste is based on its finding that the product is effective in helping to prevent and reduce tooth decay, when used as directed.

NO ANIMAL BASED INGREDIENTS FOR INSTITUTIONAL USE

Dist. by: COLGATE-PALMOLIVE COMPANY
New York, NY 10022 Made in Mexico
www.colgate.com

0 35000 78000 3 479952 951256 6499

COLGATE CAVITY PROTECTION

sodium fluoride gel, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65954-591
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080)		FLUORIDE ION	1.1 mg in 1 g	
Inactive Ingredients				
Ingredient Name			Strength	
SORBITOL (UNII: 506T60A25R)			680 mg in 1 g	
WATER (UNII: 059QF0KO0R)				
HYDRATED SILICA (UNII: Y6O7T4G8P9)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
POLYETHYLENE GLYCOL 600 (UNII: NL4J9F21N9)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				
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:65954-591-71	119 g in 1 TUBE; Type 0: Not a Combination Product	04/06/2014	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH DRUG	M021	04/06/2014		

Labeler - Mission Hills S.A de C.V (812312122)

Revised: 4/2019

Mission Hills S.A de C.V