CREST GUM DETOXIFY- stannous fluoride paste, dentifrice The 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®

Gum Detoxify ®

Drug Facts

Active ingredient
Stannous fluoride 0.454%
(0.14% w/v fluoride ion)

Purposes

Anticavity, antigingivitis toothpaste

Uses

- aids in the prevention of cavities
- helps prevent gingivitis
- helps interfere with the harmful effects of plaque associated with gingivitis
- helps control plaque bacteria that contribute to the development of gingivitis

Warning

Keep out of reach of children under 12 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 12 yrs. 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.
- do not swallow
- children under 12 yrs. of age: ask a dentist

Other information

- products containing stannous fluoride may produce surface staining of the teeth
- adequate toothbrushing may prevent these stains which are not harmful or permanent and may be

removed by your dentist

- this Crest is specially formulated to help prevent staining
- see your dentist regularly

Inactive ingredients

water, sorbitol, hydrated silica, sodium lauryl sulfate, carrageenan, sodium gluconate, flavor, xanthan gum, zinc citrate, stannous chloride, sodium saccharin, sodium hydroxide, sucralose, titanium dioxide

Questions?

1-800-594-4158 DISTR. BY PROCTER & GAMBLE, CINCINNATI, OH 45202

Principal Display Panel - 24 g tube in carton

Crest®

Gum

Detoxify®

FLUORIDE TOOTHPASTE FOR ANTICAVITY AND ANTIGINGIVITIS

Neutralizes plaque bacteria, even around the gum line

DEEP CLEAN

NET WT 0.85 OZ (24 g)



stannous fluoride paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37000-481
Route of Administration	DENTAL		

l	Active Ingredient/Active Moiety			
l	Ingredient Name	Basis of Strength	Strength	
l	STANNOUS FLUORIDE (UNII: 3FTR44B32Q) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1.4 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
SUCRALOSE (UNII: 96K6UQ3ZD4)		
STANNOUS CHLORIDE (UNII: 1BQV3749L5)		
WATER (UNII: 059QF0KO0R)		
HYDRATED SILICA (UNII: Y6O7T4G8P9)		
CARRAGEENAN (UNII: 5C69 YCD2YJ)		
SODIUM GLUCONATE (UNII: R6Q3791S76)		
XANTHAN GUM (UNII: TTV12P4NEE)		
ZINC CITRATE (UNII: K72I3DEX9B)		
SODIUM LAURYL SULFATE (UNII: 368 GB5141J)		
SORBITOL (UNII: 506T60A25R)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
SACCHARIN SO DIUM (UNII: SB8 ZUX40 TY)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor	SPEARMINT	Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-481-01	1 in 1 CARTON	08/01/2017	
1		24 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:37000-481-41	1 in 1 CARTON	0 1/29 /20 18	
2		116 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 final	part355	08/01/2017	

${f Labeler}$ - The Procter & Gamble Manufacturing Company (004238200)

Revised: 1/2018

The Procter & Gamble Manufacturing Company