AIM CAVITY PROTECTION MULTI BENEFIT- sodium fluoride gel, dentifrice Church & Dwight Co., Inc.

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.

AIM CAVITY PROTECTION Multi Benefit

Active ingredients

Sodium fluoride (0.24%)

Purpose

Anticavity toothpaste

Use aids in the prevention of dental decay

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 do not swallow supervise children as necessary until capable of using without supervision

adults and children 2 years and older brush teeth thoroughly after meals or at least twice a day, or use as directed by a dentist or physician

children under 6 years instruct in good brushing and rinsing habits (to minimize swallowing)

children under 2 years ask a dentist or physician

Inactive ingredients sorbitol, water, hydrated silica, PEG-8, sodium lauryl sulfate, SD alcohol 38-B, flavor, cellulose gum, sodium saccharin, blue 1, yellow 10.

Questions or comments? Call 1-800-786-5135 Monday-Friday 9am-5pm ET

Principal Display Panel

Value Pack

20%

MORF

Product

on sizes up to 4.6 oz.

ADA

Accepted

American

Dental

Association

TAKE AIM AGAINST CAVITIES

Aim

CAVITY PROTECTION

Ultra Mint

Gel

MULTI-BENEFIT

Cleans

Freshens

Protects

Anticavity Fluoride Gel Toothpaste

NET WT. 5.5 OZ. (156g)



AIM CAVITY PROTECTION MULTI BENEFIT

sodium fluoride gel, dentifrice

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10237-633
Route of Administration	DENTAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080)	FLUORIDE ION	2.4 mg in 1 g		

Inactive Ingredients				
Ingredient Name	Strength			
SORBITOL (UNII: 506T60A25R)				
WATER (UNII: 059QF0KO0R)				
HYDRATED SILICA (UNII: Y6O7T4G8P9)				
PEG-8 STEARATE (UNII: 2P9L47VI5E)				
SODIUM LAURYL SULFATE (UNII: 368GB5141J)				
ALCOHOL (UNII: 3K9958V90M)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				

Product Characteristics			
Color	GREEN	Score	
Shape		Size	
Flavor	MINT (Ultra Mint)	Imprint Code	
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:10237-633- 60	1 in 1 CARTON	08/01/2015	09/26/2019	
1		170 g in 1 TUBE; Type 0: Not a Combination Product			
2	NDC:10237-633- 72	1 in 1 CARTON	08/01/2015	09/26/2019	
2		204 g in 1 TUBE; Type 0: Not a Combination Product			
3	NDC:10237-633- 22	1 in 1 CARTON	08/01/2015	09/26/2019	
3		62 g in 1 TUBE; Type 0: Not a Combination Product			
4	NDC:10237-633- 55	1 in 1 CARTON	08/01/2015		
4		156 g in 1 TUBE; Type 0: Not a Combination Product			
5	NDC:10237-633- 66	1 in 1 CARTON	08/01/2015		
		107 a in 1 TIDE. Time O. Not a Combination			

5	187 g in 1 TUBE; Type U: NOT a Combination Product			
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL	part355	03/01/2004		

Labeler - Church & Dwight Co., Inc. (001211952)

Establishment				
Name	Address	ID/FEI	Business Operations	
Church & Dwight Co., Inc.		043690812	MANUFACTURE(10237-633)	

Revised: 1/2023 Church & Dwight Co., Inc.