NATURE MINT ANTICAVITY- sodium monofluorophosphate paste, dentifrice Bob Barker Company Inc.

Nature Mint® Anticavity

Drug Facts:

ACTIVE INGREDIENT

Sodium Monofluorophosphate - 0.76% (0.1% W/V fluoride ion)

PURPOSE

Anticavity toothpaste

Use

Helps protect against cavities.

Warnings

Keep out of the 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 immediately.

Directions

Adults &	Brush teeth thoroughly, preferably
	after each meal or at least twice a day,
of age & older:	or as directed by a dentist or doctor.
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

Calcium Carbonate, Water, Glycerin, Silica, Sodium Lauryl Sulfate, Xanthan Gum, Flavor, Sodium Saccharin, Titanium Dioxide, Sodium Methylparaben, Sodium Propylparaben

Dist. by Bob Barker Co. Inc. Fuquay-Varina, NC 27526

PRINCIPAL DISPLAY PANEL - 43 Gram Tube Label

Nature Mint_®
ANTICAVITY FLUORIDE TOOTHPASTE
SODIUM MONOFLUOROPHOSPHATE - 0.76%
FRESH MINT FLAVOR

NET WT 1.5 OZ (43 GRAMS)



NATURE MINT ANTICAVITY

sodium monofluorophosphate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53247-131
Route of Administration	DENTAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Sodium Monofluorophosphate (UNII: C810JCZ56Q) (Fluoride Ion - UNII:Q80VPU408O)	Fluoride Ion	7.6 mg in 1 g	

Inactive Ingredients			
Ingredient Name	Strength		
Calcium Carbonate (UNII: H0G9379FGK)			
Water (UNII: 059QF0KO0R)			
Glycerin (UNII: PDC6A3C0OX)			
Silicon Dioxide (UNII: ETJ7Z6XBU4)			
Sodium Lauryl Sulfate (UNII: 368GB5141J)			
Xanthan Gum (UNII: TTV12P4NEE)			
Saccharin Sodium (UNII: SB8ZUX40TY)			
Titanium Dioxide (UNII: 15FIX9V2JP)			
Methylparaben Sodium (UNII: CR6K9C2NHK)			
Propylparaben Sodium (UNII: 625NNB0G9N)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53247-131- 01	7.94 g in 1 PACKET; Type 0: Not a Combination Product	01/01/2023	
2	NDC:53247-131- 02	17 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2023	
3	NDC:53247-131- 03	24 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2023	
4	NDC:53247-131- 04	43 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2023	
5	NDC:53247-131- 05	78 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2023	
6	NDC:53247-131- 06	1 in 1 CARTON	01/01/2023	
6		130 g in 1 TUBE; Type 0: Not a Combination Product		
7	NDC:53247-131- 07	1 in 1 CARTON	01/01/2023	
7		181 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	01/01/2023	

Labeler - Bob Barker Company Inc. (058525536)

Revised: 4/2024 Bob Barker Company Inc.