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, Sorbitol, Glycerin, Sodium Lauryl Sulfate, Hydrated Silica, Cellulose Gum, Flavor, Sodium Saccharin, Tetrasodium Pyrophosphate, Sodium Benzoate

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

PRINCIPAL DISPLAY PANEL - 24 GRAM Tube Label

 $\text{Nature Mint}_{\text{\mathbb{R}}}$

ANTICAVITY FLUORIDE TOOTHPASTE

SODIUM MONOFLUOROPHOSPHATE - 0.76%

FRESH MINT FLAVOR

NET WT 0.85 OZ (24 GRAMS)



NATURE MINT ANTICAVITY

sodium monofluorophosphate paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53247-123
Route of Administration	DENTAL		

Active Ingredient/Active Moiety

Ingredient Name	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)				
Sorbitol (UNII: 506T60A25R)				
Glycerin (UNII: PDC6A3C0OX)				
Sodium Lauryl Sulfate (UNII: 368GB5141J)				
Hydrated Silica (UNII: Y6O7T4G8P9)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K6790BS311)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SODIUM PYROPHOSPHATE (UNII: O352864B8Z)				
Sodium Benzoate (UNII: OJ245FE5EU)				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53247-123- 01	7.94 g in 1 PACKET; Type 0: Not a Combination Product	01/01/2008	
2	NDC:53247-123- 02	17 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
3	NDC:53247-123- 03	24 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
4	NDC:53247-123- 04	43 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
5	NDC:53247-123- 05	78 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
6	NDC:53247-123- 06	1 in 1 CARTON	01/01/2008	
6		130 g in 1 TUBE; Type 0: Not a Combination Product		
7	NDC:53247-123- 07	1 in 1 CARTON	01/01/2008	
7		181 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC monograph drug	M021	01/01/2008	

Labeler - Bob Barker Company Inc. (058525536)