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 & Children 6 years of age & older:	Brush teeth thoroughly, preferably after each meal or at least twice a day, or as directed by a dentist or doctor.
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

Calcium Carbonate, Sorbitol, Water, Sodium Lauryl Sulfate, Silica, Glycerin, Cellulose Gum, Benzalkonium Chloride, Flavor, Sodium Saccharin, PEG 400, Sodium Chloride, Sodium Methylparaben, Menthol, 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:53247-129
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)	
Sorbitol (UNII: 506T60A25R)	
Water (UNII: 059QF0KO0R)	
Sodium Lauryl Sulfate (UNII: 368GB5141J)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
Glycerin (UNII: PDC6A3C0OX)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
Benzalkonium Chloride (UNII: F5UM2KM3W7)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
Sodium Chloride (UNII: 451W47IQ8X)	
METHYLPARABEN SODIUM (UNII: CR6K9C2NHK)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
PROPYLPARABEN SODIUM (UNII: 625NNB0G9N)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53247-129-01	7.94 g in 1 PACKET; Type 0: Not a Combination Product	01/01/2008	
2	NDC:53247-129-02	17 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
3	NDC:53247-129-03	24 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
4	NDC:53247-129-04	43 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
5	NDC:53247-129-05	78 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2008	
6	NDC:53247-129-06	1 in 1 CARTON	01/01/2008	
6		130 g in 1 TUBE; Type 0: Not a Combination Product		
7	NDC:53247-129-07	1 in 1 CARTON	01/01/2008	
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	01/01/2008	

Labeler - Bob Barker Company Inc. (058525536)

Revised: 2/2024 Bob Barker Company Inc.