MAXIXUM SECURITY GEL TOOTHPASTE- fluoride toothpaste paste, dentifrice
Dabur India Limited

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.

MAXIMUM SECURITY® GEL TOOTHPASTE

NET WT 0.15 OZ

DRUG FACTS:

ACTIVE INGREDIENT

Sodium fluoride- 0.22% (0.1% w/v fluoride ion)

PURPOSE

Anticavity.

USE

Helps protect against cavities.

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 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 physician.

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

Purified Water, Sorbitol, Carbopol, Sodium Lauryl Sulphate, Flavor, Poly Ethylene Glycol 1500, Sodium Saccharin, Precipitated Silica, Sodium Carboxy Methyl Cellulose, Methyl Paraben, Propyl Paraben.

Product of India,
Exclusive Distributor:
Bob Barker Co. Inc.,
Phone: 1-800-334-9880.

Expires: 3 years from the Date of Mfg.

C.No. DNH/COS/DNH/52
Mfg. Date & Batch No. on crimp.

24.06.2009

MAXIMUM SECURITY GEL TOOTHPASTE
ACTIVE INGREDIENT: SODIUM FLUORIDE - 0.22 %
(0.1% w/v fluoride Ion)
SR. NO. INGREDIENTS

1. Treated water
2. Sorbitol
3. Carbopol
4. Sodium lauryl sulphate
5. Flavor
6. Polyethylene glycol 1500
7. Sodium saccharin
8. Precipitated Silica
9. Sodium Carboxy Methyl Cellulose
10. Methyl Paraben
11. Propyl Paraben

Principal Display Panel

Packet Label
TEAR HERE
MAXIMUM SECURITY®
GEL TOOTHPASTE
NET WT 0.15 OZ

MAXIXUM SECURITY GEL TOOTHPASTE

fluoride toothpaste paste, dentifrice

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68747-6028
Route of Administration	DENTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	SODIUM FLUORIDE	2.2 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
POLYETHYLENE GLYCOL 1500 (UNII: 1212Z7S33A)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	MINT (MINT)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68747-6028-1	4.2 g in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part355	08/25/2009	

Labeler - Dabur India Limited (650319218)

Establishment

Name	Address	ID/FEI	Business Operations
Dabur India Limited		650319218	MANUFACTURE

Revised: 8/2008 Dabur India Limited