ANTICAVITY FLUORIDE RINSE- sodium fluoride mouthwash
HEB

Drug Facts HEB 163

TEP

SEALED WITH PRINTED NECKBAND FOR YOUR PROTECTION

Active ingredient

Sodium fluoride 0.0221% (0.01% w/v fluoride ion)

Purpose

Anticavity

Use

aids in the prevention of dental cavities

Warnings

For this product

Keep out of reach of children

If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 6 years of age and older:

- use twice daily after brushing your teeth with toothpaste
- vigorously swish 10 mL (2 teaspoonfuls) of rinse between your teeth for 1 minute and then spit out
- do not swallow the rinse
- do not eat or drink for 30 minutes after rinsing
- instruct children under 12 years of age in good rinsing habits (to minimize swallowing)
- supervise children as necessary until capable of using without supervision
- children under 6 years of age: consult a dentist or doctor

Other information

- store at controlled room temperature 20º -25º C (68º - 77º F)
- cold weather may temporarily cloud this product

Inactive ingredients

water, alcohol (21.6 %v/v), sorbitol, poloxamer 407, eucalyptol, flavor, methyl salicylate, menthol, phosphoric acid, sodium saccharin, thymol, disodium phosphate, sucralose, red 40, blue 1

Questions?

Call 1-888-593-0593

This product is not manufacturd or distributed by Johnson & Johnson Healthcare Products, distributor of Listerine Total Care Anticavity Mouthwash

MADE WITH PRIDE AND CARE FOR H-E-B, SAN ANTONIO, TX 78204

principal display panel

Compare to Listerine Total Care

Anticavity Mouthwash

H-E-B

Mint

ANTICAVITY MOUTHWASH

SODIUM FLUORIDE AND ACIDULATED PHOSPHATE TOPICAL SOLUTION
- Kills Germs That Cause Bad Breath
- Helps Prevent Cavities

IMPORTANT: read directions for proper use.

33.8 FL OZ (1 QT) 1L

ANTICAVITY FLUORIDE RINSE

sodium fluoride mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-163
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
SORBITOL (UNII: 506T60A25R)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
EUCALYPTOL (UNII: RV6J6604TK)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
MENTHOL (UNII: L7T10EIP3A)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
THYMOL (UNII: 3J50XA376E)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-163-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/29/2010	
2	NDC:37808-163-77	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/29/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	09/29/2010	

Labeler - HEB (007924756)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(37808-163)

Establishment

Name	Address	ID/FEI	Business Operations

Vi-Jon, LLC		088520668	manufacture(37808-163)