ANTICAVITY FLUORIDE RINSE- sodium fluoride mouthwash Target Corporation

Anticavity Fluoride Rinse 163.005 163AF-AG

Tamper Evident Statement

Sealed with printed neckband for your protecction

Active ingredient

Sodium fluoride 0.02% w/v (0.01% w/v fluoride ion)

Purpose

Anticavity

Use

aids in the prevention of dental cavities

Warning

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 20°-25° C (68° 77° 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 1-800-910-6874

Advrse reaction

Satisfaction guaranteed - Love it or your money back.

Distributed by

Target Corporation

Minneapolis, MN 55403

Made in the U.S. with U.S. and foreign components

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Target Brands, Inc.

DSP-TN-21091

DSP-M)-20087

how2recycle.info

Principal Panel Display

Multi-Action Mouthwash

Anticavity Fluoride Rinse

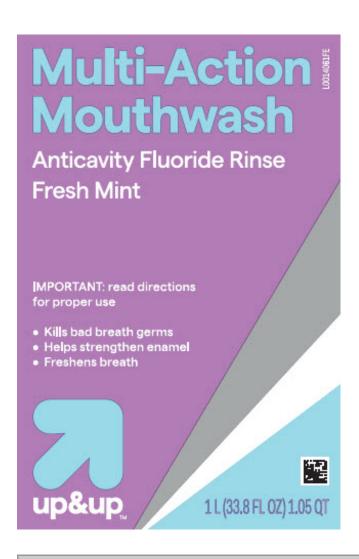
Fresh Mint

IMPORTANT: read directions for proper use

- Kills bad breath germs
- Helps strengthen enamel
- Freshens breath

up&up™

1L (33.8 FL OZ) 1.05 QT



ANTICAVITY FLUORIDE RINSE

sodium fluoride mouthwash

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11673-163

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU4080) FLUORIDE ION

0.1 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:11673- 163-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/06/1994	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	04/06/1994	

Labeler - Target Corporation (006961700)

Registrant - Vi-Jon, LLC (790752542)

Establishment				
Name	Address	ID/FEI	Business Operations	
Vi-Jon, LLC		790752542	manufacture(11673-163)	

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11673-163)

Revised: 12/2023 Target Corporation