ALCOHOL-FREE ANTICAVITY- sodium fluoride liquid
Target Corporation

UP & UP 482.001/482AB
Anticavity Fluoride Rinse

Active ingredient

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

Purpose

Anticavity

Use

aids in the prevention of dental cavities

Warnings

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 a toothpaste
- vigorously swish 10 mL (2 teaspoonfuls) of rinse between your teeth for 1 minute 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

Inactive ingredients

water, glycerin, cetylpyridinium chloride, flavor, sodium saccharin, phosphoric acid, sodium benzoate, sucralose, poloxamer 407, benzoic acid, disodium phosphate, propylene glycol, red 33, green 3

Questions or comments

1-800-910-6874

This rinse may cause temporary staining to the surface of teeth. This is not harmful, and adequate burshing may prevent it occurence.

*This product is not manufactured or distributed by Procter & Gamble, distributor of Crest Pro-Health Complete Anticavity Fluoride Rinse

Distributed by Target Corporation

Minneapolis, MN 55403

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

principal display panel

SEALED WTIH PRINTED NECKBAND FOR YOUR PROTECTION

anticavity

alcohol-free fluoride mouth rinse

Compare to the active ingredient in Crest Advanced*

helps prevent cavities

helps make enamel stronger

helps clean the entire mouth

freshens breath

no alcohol burn

IMPORTANT: read directions for proper use

up & up

MINT FLAVOR

33.8 FL OZ (1 L)

ALCOHOL-FREE ANTICAVITY

sodium fluoride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-482
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM FLUORIDE (UNII: 8ZYQ1474W7) (FLUORIDE ION - UNII:Q80VPU408O)	FLUORIDE ION	0.1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-482-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/09/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M021	03/09/2012	

Labeler - Target Corporation (006961700)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(11673-482)

Revised: 7/2024 Target Corporation