

ANTISPETIC- eucalyptol, menthol, methyl salicylate, thymol mouthwash

Target Corporation

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.

Drug Facts

Active ingredients

Eucalyptol 0.092%

Menthol 0.042%

Methyl salicylate 0.060%

Thymol 0.064%

Purpose

Antigingivitis, antiplaque

Use

help control plaque that leads to gingivitis

Warnings for this product

Warnings

Do not use

if you have painful or swollen gums, pus from the gum line, loose teeth or increased spacing between the teeth. See your dentist immediately. These may be signs of periodontitis, a serious form of gum disease.

Stop use

And ask a dentist if gingivitis, bleeding, or redness persists for more than 2 weeks.

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 12 years of age and older - vigorously swish 20 mL (2/3 FL OZ or 4 teaspoonfuls) between teeth for 30 seconds then spit out; do not swallow

children under 12 years of age - consult a dentist or doctor

- this rinse is not intended to replace brushing or flossing

other information

cold weather may cloud this product. Its antiseptic properties are not affected. Store at room temperature (59°-77°F).

Inactive ingredients

water, alcohol 21.6%, sorbitol solution, flavor, poloxamer 407, benzoic acid, sodium saccharin, sodium benzoate, FD&C green no.3

Disclaimer

This product is not manufactured or distributed by Johnson & Johnson Healthcare Products, distributor of Listerine.

ADA Council Statement

The ADA Council on Scientific Affairs Acceptance of UP & UP blue mint antiseptic mouthwash is based on its finding that the product is effective in helping to prevent and reduce gingivitis and plaque above the gumline, when used as directed.

Adverse Reactions

Distributed by Target Corporation

Minneapolis, MN 55403

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

shop Target.com

Guest Services 1-800-910-6874

principal display panel

SAFETY SEALED WITH PRINTED NECKBAND FOR YOUR PROTECTION

ADA

Accepted

American

Dental

Association

antiseptic

mouthwash

Kills germs that cause bad breath, plaque and the gum disease gingivitis

Compare to Listerine Antiseptic Cool Mint

BLUE MINT FLAVOR

up & up

50.7 FL OZ (1.5 L)

SAFETY SEALED WITH PRINTED NECKBAND FOR YOUR PROTECTION



antiseptic mouthwash

Kills germs that cause bad breath, plaque
and the gum disease gingivitis

Compare to Listerine® Antiseptic Cool Mint®*



50.7 FL OZ (1.5 L)

L0001691FG

ANTISPETIC

eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-664
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
-----------------	-------------------	----------

EUCALYPTOL (UNII: RV6J6604TK) (EUCALYPTOL - UNII:RV6J6604TK)	EUCALYPTOL	0.92 mg in 1 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.42 mg in 1 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	0.60 mg in 1 mL
THYMOL (UNII: 3J50XA376E) (THYMOL - UNII:3J50XA376E)	THYMOL	0.64 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
SORBITOL (UNII: 506T60A25R)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
BENZOIC ACID (UNII: 8SKN0B0MM)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C GREEN NO. 3 (UNII: 3P3ONR601S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-664-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	
2	NDC:11673-664-77	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	
3	NDC:11673-664-19	94 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	08/11/1993	

Labeler - Target Corporation (006961700)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		790752542	manufacture(11673-664)