ANTISPETIC- eucalyptol, menthol, methyl salicylate, thymol mouthwash
Meijer Distribution Inc.

Meijer Blue Mint Antiseptic Mouthrinse
664.003/664AT rev 2 - AU

TEP

SEALED WITH A PRINTED NECKBAND FOR YOUR PROTECTION

Active ingredients

Eucalyptol 0.092%

Menthol 0.042%

Methyl salicylate 0.060%

Thymol 0.064%

Purpose

Antigingivitis, Antiplaque

Use

helps control plaque that leads to gingivitis

Warnings

for this product

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, poloxamer 407, benzoic acid, sodium saccharin, sodium benzoate, flavor, green.3

Questions

1-888-593-0593

Disclaimer

*This product is not manufactured or distributed by Kenvue, Inc, distributor of Cool Mint ® Listerine ® Antiseptic Mouthwash.

Adverse reactions

DISTRIBUTED BY

MEIJER DISTRIBUTION, INC.

GRAND RAPIDS, MI 49544

www.meijer.com

OUR QUALITY GUARANTEE

The Meijer Family

www.meijer.com/satisfaction

how2recycle.info

Principal display panel

NDC 41250-664-86

Compare to Cool Mint ® Listerine ® Active Ingredients*

meijer

Mouth Rinse

Antigingivitis

Antiplaque

Blue Mint

Antiseptic

Kills germs that cause bad breath, plaque and gingivitis

ADA Accepted

American Dental Association

- Helps prevent and reduce plaque
- Helps prevent and reduce gingivitis

33.8 FL OZ (1.05 QT) 1L

ANTISPETIC

eucalyptol, menthol, methyl salicylate, thymol mouthwash

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41250-664
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	0.6 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: 8SKN0B0MIM)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-664-19	94 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	
2	NDC:41250-664-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	
3	NDC:41250-664-77	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	
4	NDC:41250-664-69	250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	11/11/2017
5	NDC:41250-664-44	532 mL in 1 PACKAGE; Type 0: Not a Combination Product	08/11/1993	01/01/2020
6	NDC:41250-664-12	1500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	
7	NDC:41250-664-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	
8	NDC:41250-664-13	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/1993	05/08/2016

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/11/1993	

Labeler - Meijer Distribution Inc. (006959555)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(41250-664)

Revised: 2/2024 Meijer Distribution Inc.