

**QUALITY CHOICE BLUE MINT- eucalyptol, menthol, methyl salicylate, thymol liquid
CHAIN DRUG MARKETING ASSOCIATION INC**

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

Antiplaque/Antigingivitis

Uses

To help reduce and prevent plaque and gingivitis

Warnings

☐**Do not use**☐for children under 12 years of age.

Keep out of reach of children

If more than used for rinsing is accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- Rinse full strength for 30 seconds with 20 mL (2/3 fluid ounce or 4 teaspoonfuls) morning and night.
- Do not swallow

Other information

- Store at room temperature.
- Cold weather may cloud this product. Its antiseptic properties are not affected.

Inactive ingredients


Water (Aqua), Alcohol (21.6%), Sorbitol, Flavor, Poloxamer 407, Benzoic Acid, Sodium Saccharin, Sodium Benzoate, Green 3 (CI 42053)

Questions or comments?

1-248-449-9300

Label Copy

SEALED WITH PRINTED NECK BAND FOR YOUR PROTECTION



Compare to
Listerine® Cool Mint
Antiseptic Mouthwash*

Antiseptic Mouth Rinse

Antigingivitis | Antiplaque

Kills Germs That Cause:
Bad Breath
Plaque & Gingivitis
Gum Disease

Blue Mint
1 Liter (33.8 FL OZ)

06-20801

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MADE IN CANADA

DO NOT USE IF PRINTED BAND AROUND CAP IS BROKEN OR MISSING.

TO OPEN: SQUEEZE CAP AND TURN.

TO CLOSE: TURN CAP UNTIL IT LOCKS.

*This product is not manufactured or distributed by Johnson & Johnson Healthcare Products, division of McNeil-PPC, Inc., distributor of Cool Mint Listerine® Antiseptic Mouthwash.



Distributed by C.D.M.A., Inc.®
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Novi, MI 48376-0995
www.qualitychoice.com
Questions: 248-449-9300



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06-20802

QUALITY CHOICE BLUE MINT

eucalyptol, menthol, methyl salicylate, thymol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-557
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: 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:63868-557-33	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part356	09/13/2015	

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)**Registrant** - APOLLO HEALTH AND BEAUTY CARE (201901209)**Establishment**

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(63868-557)

Revised: 9/2015

CHAIN DRUG MARKETING ASSOCIATION INC