# 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.

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#### **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.

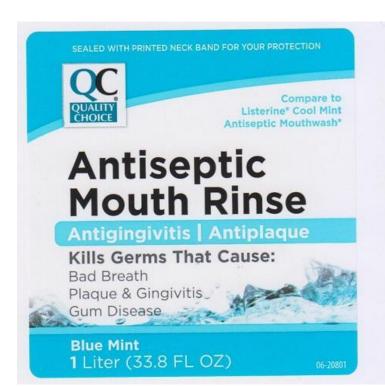
#### **Inactive ingredients**

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**



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Questions or comments? 1-	248-449-9300			

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 Listering\* Antiseptic Mouthwesh





06-20802

## **QUALITY CHOICE BLUE MINT**

**Route of Administration** 

eucalyptol, menthol, methyl salicylate, thymol liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-557	

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: 3J50 XA376E) (THYMOL - UNII:3J50 XA376E)	THYMOL	0.64 mg in 1 mL		

Inactive Ingredients			
Strength			

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		
_	F 1 T			

# 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