

QUALITY CHOICE MINT- cetylpyridinium chloride 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.

Active ingredients

Cetylpyridinium Chloride 0.07%

Purpose

Anti plaque/Antigingivitis

Uses

- Helps reduce and prevent plaque and gingivitis.
- Helps control plaque bacteria that contribute to the development of gingivitis and bleeding gums.

Warnings

Ask dentist if symptoms persist or condition worsens after regular use.

Keep out of reach of children under 6 years of age.

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

Directions

- Use after your normal brushing and flossing routine; rinse toothpaste from mouth prior to use
- **Adults and children 6 years and older:** Rinse for 30 seconds with 20 mL (4 teaspoonfuls) twice a day.
- Do not swallow
- Children 6 years to under 12 years of age: supervise use
- **Children under 6 years of age: do not use.**

Other information

Store at room temperature

Inactive ingredients

Water (Aqua), Glycerin, Poloxamer 407, Flavor, Sodium Saccharin, Methylparaben, Sucralose, Propylparaben, Blue 1 (CI 42090)

Questions or comments?

1-248-449-9300

Label Copy



Do not use if band around cap is broken or missing.
To open: Squeeze smooth areas on cap and turn.
To close: Turn cap until it locks.

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 Novi, MI 48376-0995
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 Made in Canada



06-20800

QUALITY CHOICE MINT

cetylpyridinium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-553
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETYL PYRIDINIUM CHLORIDE (UNII: D9OM4SK49P) (CETYL PYRIDINIUM - UNII:CUB7J10JV3)	CETYL PYRIDINIUM CHLORIDE	0.7 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

GLYCERIN (UNII: PDC6A3C0OX)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-553-33	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 MARKET ING ASSOCIATION INC (011920774)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(63868-553)