

GIANT EAGLE REFRESHING CITRUS REFILL- benzalkonium chloride liquid
Apollo Health and Beauty Care 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 ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

helps eliminate bacteria on hands.

Warnings

For external use only

When using this product

avoid contact with eyes. In case of contact, rinse thoroughly with water.

Stop use and ask a doctor if
irritation or redness develops and lasts.

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

- Use to refill a pump bottle. From pump bottle, apply onto wet hands.
- Lather and rinse thoroughly.

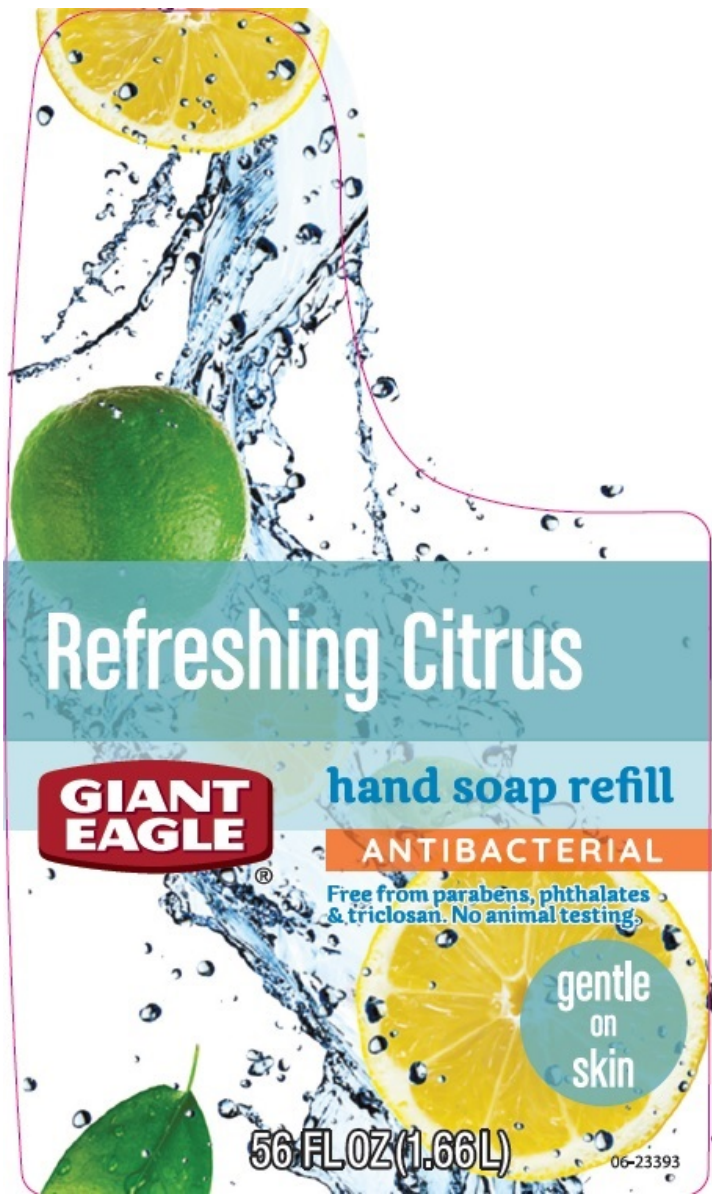
Other information

- store at room temperature

Inactive ingredients

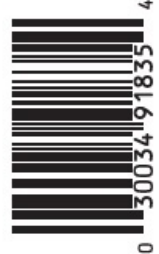
Water (Aqua), Cetrimonium Chloride, Glycerin, Lauramidopropylamine Oxide, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Citrus Nobilis (Mandarin Orange) Peel Extract, Camellia Sinensis Leaf Extract, Zingiber Officinale (Ginger) Root Extract, Methylchloroisothiazolinone, Methylisothiazolinone, Blue 1 (CI 42090), Yellow 5 (CI 19140).

Label Copy



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MADE IN CANADA



OUR QUALITY GUARANTEE

Giant Eagle's commitment to quality is always 100% satisfaction or double your money back.

06-23394

GIANT EAGLE REFRESHING CITRUS REFILL			
benzalkonium chloride liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63148-724
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL
Inactive Ingredients			
	Ingredient Name		Strength
	WATER (UNII: 059QF0KO0R)		

CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)
GLYCERIN (UNII: PDC6A3C0OX)
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
COCO MONOETHANOLAMIDE (UNII: C80684146D)
PEG-120 METHYL GLUCOSE DIOLATE (UNII: YM0K64F20V)
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)
EDETATE SODIUM (UNII: MP1J8420LU)
SODIUM SULFATE (UNII: 0YPR65R21J)
CITRUS NOBILIS (UNII: 8MFF77J91V)
GREEN TEA LEAF (UNII: W2ZU1RY8B0)
GINGER (UNII: C5529G5JPQ)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-724-56	1656 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/12/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/12/2018	

Labeler - Apollo Health and Beauty Care Inc. (201901209)

Registrant - Apollo Health and Beauty Care Inc. (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(63148-724)

Revised: 9/2018

Apollo Health and Beauty Care Inc.