

P.O.V. LIGHT MOISTURIZING ANTIBACTERIAL HAND- benzalkonium chloride soap
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 cast of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

- apply onto dry hadns
- lather and rinse thoroughly.

Other information

Store at room temperature.

Inactive ingredients

Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Red 40 (CI 16035), Yellow 5 (CI 19140), Red 33 (CI 17200).

Questions or comments?

1-866-695-3030

Label Copy

P.O.V.TM

LIGHT
MOISTURIZING
ANTIBACTERIAL
**HAND
SOAP**

- NOURISHING MOISTURE
- ENHANCED HYDRATION WITH NATURAL INGREDIENTS

**7.5 FL OZ
(222 mL)**

06-22774

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 ■ apply onto dry hands ■ lather and rinse thoroughly.	
Other information store at room temperature.	
Inactive ingredients Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Red 40 (CI 16035), Yellow 5 (CI 19140), Red 33 (CI 17200).	
Questions or comments? 1-866-695-3030	

Distributed by:
Apollo Health & Beauty Care Inc.
1 Apollo Place, Toronto, ON M3J 0H2
06-22775

MADE IN CANADA



0 67153 94847 4



LIGHT
MOISTURIZING
ANTIBACTERIAL
**HAND
SOAP**

LIGHT
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**HAND
SOAP**

- NOURISHING MOISTURE
- ENHANCED HYDRATION WITH NATURAL INGREDIENTS

**56 FL OZ
(1.65 L)**

06-22782

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 ■ apply onto dry hands. ■ lather and rinse thoroughly.	
Other information store at room temperature.	
Inactive ingredients Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Red 40 (CI 16035), Yellow 5 (CI 19140), Red 33 (CI 17200).	
Questions or comments? 1-866-695-3030	

06-22783

MADE IN CANADA

Distributed by:

Apollo Health & Beauty Care Inc.

1 Apollo Place, Toronto, ON M3J 0H2



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P.O.V. LIGHT MOISTURIZING ANTIBACTERIAL HAND

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63148-199
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)	

LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)
GLYCERIN (UNII: PDC6A3C0OX)
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)
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)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
FD&C RED NO. 40 (UNII: WZB9127XOA)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)
D&C RED NO. 33 (UNII: 9DBA0SBB0L)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-199-08	222 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2017	
2	NDC:63148-199-56	1656 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/10/2017	

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(63148-199)

Revised: 11/2017

Apollo Health and Beauty Care Inc.