

CAREONE AMBER- benzalkonium chloride soap
Retail Business Services, LLC

72476-004- Careone Amber

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 Poison Control Center immediately.

Directions

- apply onto wet hands (only for 222 ml and 333 ml)
- Lather and rinse thoroughly.(For all packages)
- From a pump bottle, apply onto wet hands (only for 1.66 L bottle)
- Use to refill a pump bottle (only for 1.66 L bottle)

Other Information

Store at Room Temperature

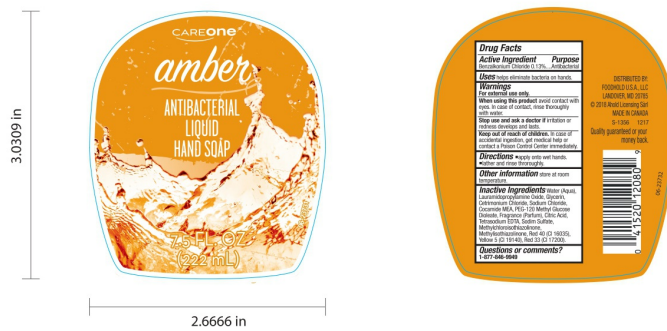
Inactive Ingredients

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?

1-877-846-9949

PDP2



PDP3

Front Label



Back Label



ART #:01596 DESIGNER #: KR **NOTE:**

CLIENT: APOLLO JOB DESCRIPTION: 06-24094-95 CAREONE AMBER DATE: JUNE 13, 2019 PROOF# 1



LABEL SIZE
4" X 2.4"
DIE# NEW
ROLL DIRECTION
3 FRONT 4 BACK

COLOURS
C M Y K
PMS 145

THE CLIENT IS RESPONSIBLE FOR APPROVING THE FOLLOWING:
1. ALL COPY
2. LABEL DIMENSIONS
3. COLOURS USED
4. REWIND COPY DIRECTION
5. LOCAL, PROVINCIAL OR NATIONAL REGULATIONS ASSOCIATED WITH THIS LABEL(S)

MATERIAL:
WHITE BOPP

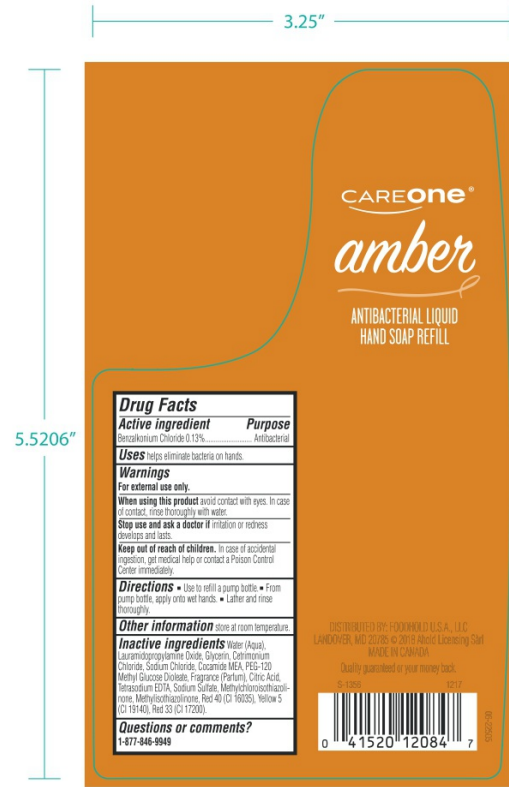
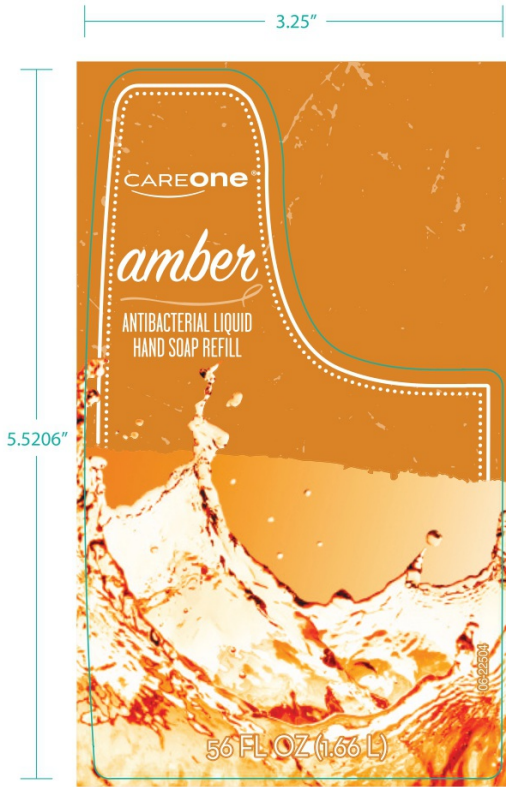
COATING:
GLOSS LAMINATE

THIS IS NOT A COLOUR ACCURATE PROOF.
Please refer to process and pantone colour charts for accurate colour representation. Every effort has been made to ensure the accuracy of this proof. However, the client is responsible for ensuring that label size, copy, graphics and colour separations are accurate, and to notify Perflex Label Inc. of any discrepancies prior to film, plate or label production. The client indemnifies Perflex Label Inc. against any liability related to costs incurred due to errors on a customer signed proof.

DIELINE
DOES NOT PRINT

APPROVED AS IS : _____
APPROVED WITH CHANGES INDICATED : _____
RE-PROOF DATE : _____

PLEASE NOTE THAT THERE WILL BE NO LABEL PRODUCTION WITHOUT A SIGNED PROOF. FAX APPROVALS TO 416 321-2267. THANK YOU.



	Northern label group inc.	Customer: Apollo DKT#: 7489-A
	# OF COLOURS: 4 Material: White PP <small>IMPORTANT: PLEASE READ!</small> <small>These colours are designated for reference only. Refer to actual ink samples for exact match.</small> LABEL WILL NOT PROCEED UNTIL ALL ITEMS BELOW HAVE BEEN CHECKED	Job Name: 56oz Care One Hand Soap Refill Description: Amber 06-22504 / 06-22505 Date: December 11, 2017 PO: 774509 NEW FILE REV# UPC Client Supplied <small>Please indicate any changes that may be needed. An approval of this file is limited to text placement, spelling accuracy and agreement on layout and screen represented colour.</small>
<input type="checkbox"/> Size of Label <input type="checkbox"/> Die Line <input type="checkbox"/> Spelling <input type="checkbox"/> UPC Code <input type="checkbox"/> Copy Layout <input type="checkbox"/> Copy Position <input type="checkbox"/> Colour Break <input type="checkbox"/> Product Code	Client Approved: _____	Changes YES <input type="checkbox"/> NO <input type="checkbox"/>

CAREONE AMBER

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72476-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	130 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

SODIUM SULFATE (UNII: 0YPR65R21J)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
FD&C RED NO. 40 (UNII: WZB9127XOA)
WATER (UNII: 059QF0KO0R)
GLYCERIN (UNII: PDC6A3C0OX)
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)
EDETATE SODIUM (UNII: MP1J8420LU)
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:72476-004-02	222 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2021	
2	NDC:72476-004-03	333 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2021	
3	NDC:72476-004-04	1660 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/01/2021	

Labeler - Retail Business Services, LLC (967989935)

Registrant - Apollo Health and Beauty Care (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care		201901209	manufacture(72476-004)

Revised: 10/2024

Retail Business Services, LLC