COMPANION- companion antibacterial hand soap liquid
Preserve International

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.

COMPANION
Foaming Antibacterial Hand Soap

DRUG FACTS
Active ingredients

Benzalkonium Chloride 0.13%.....Antibacterial

PURPOSE

Antimicrobial

USES

For hand washing to minimize bacteria on the skin.

WARNINGS

For External use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

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

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

- Pump foam into dry hands.
- Lather with water for atleast 20 seconds.
- Rinse and pat dry.

OTHER INFORMATION

Store below 110°F (43°C).

INACTIVE INGREDIENTS

Water, Xanthan Gum, Glycerin, Sodium Lauryl Eather Sulfate, Alkylpolyglycoside,

Cocamidopropyl Betaine, Citric Acid, Sodium Citrate, Tetrasodium Ethylenediamine Tetraacetate, Sodium Benzoate, Waterfall Mist, Blue #1, Yellow #5.

COMPANION

Foaming Antibacterial Hand Soap

- Kills 99.9% of common bacteria
- Moisturizing Formula
- Light Fresh Scent

NET CONTENTS: 16.9 oz. (750 ml)

Item No.: 4900912/6

Manufactured By:

Preserve International, a wholly-owned subsidiary of Neogen ®

944 Nandino Blvd. Lexington, KY 40511 USA

800-477-8201 (USA/Canada) or 859-254-1221 L7018-0920

COMPANION

companion antibacterial hand soap liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60648-7200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	2.9 mg in 1 L

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
EDETATE SODIUM (UNII: MP1J8420LU)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60648-7200-1	0.75 L in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	11/04/2020	
2	NDC:60648-7200-2	3.78 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/04/2020	

Labeler - Preserve International (808154199)

Registrant - Preserve International (808154199)

Establishment

Name	Address	ID/FEI	Business Operations
Preserve International		808154199	manufacture(60648-7200) , api manufacture(60648-7200)

Revised: 7/2023 Preserve International