HAND WASH- benzalkonium chloride soap
ULINE

Uline 466.001/466AB
Antibacterial Foaming Hand Wash

Active ingredient

Benzalkonium chloride 0.13%

purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

warnings

For external use only: hands only

When using this product

- avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children

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

Directions

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

water, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, fragrance, citric acid, tetrasodium EDTA,

benzophenone-4, sodium benzoate, red 4, yellow 5

Adverse reactions

Distributed by: ULINE, 12575 Uline Drive

pleasant Prairie, WI 53158

1-800-295-5510

uline.com

principal Display Panel

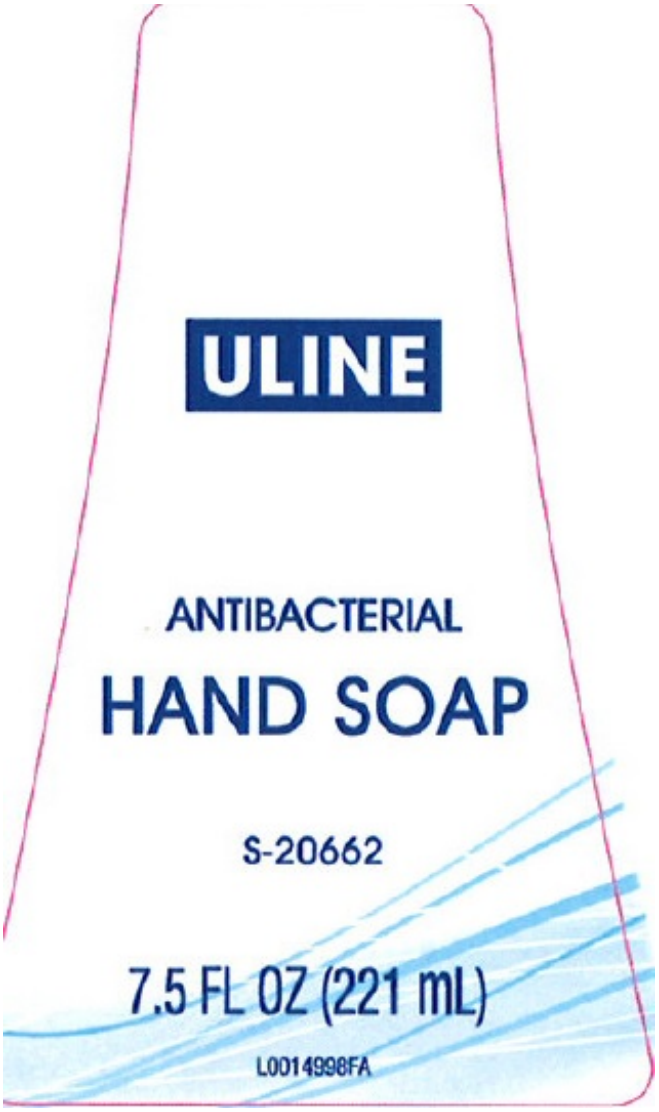
ULINE

ANTIBACTERIAL

HAND SOAP

S-20662

7.5 FL OZ (221 mL)



HAND WASH

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69790-466
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)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SULISOBENZONE (UNII: 1W6L629B4K)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69790-466-96	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/17/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/17/2016	

Labeler - ULINE (039612668)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(69790-466)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(69790-466)

Revised: 9/2024 ULINE