ZEP PEAR ANTI-BACTERIAL- benzalkonium chloride liquid
ZEP PEAR AB HS- benzalkonium chloride liquid
Zep 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.

66949-393/539 Pear AB

Benzalkonium Chloride 0.13%

Antiseptic Handwash

Uses

For hand washing to decrease bacteria on skin.

For external use only.

Do not use in the eyes; if in eyes, rinse promptly and thoroughly with water.

- Do not swallow.
- If swallowed, do not induce vomiting and if individual is conscious, give large quantities of water to drink and consult a physician immediately.

Stop use and ask doctor if skin irritation or redness persists for more than 72 hours.

Keep out of reach of children and pets. Children must be supervised in use of this product.

Directions

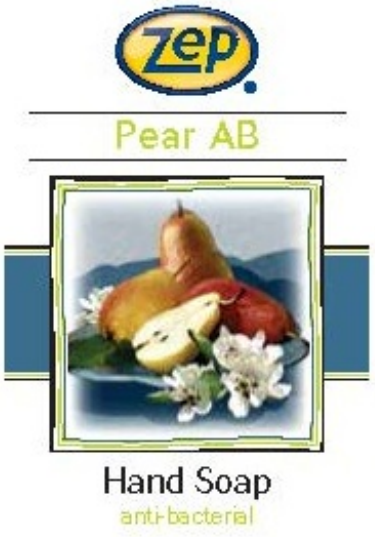
- Wet hands with water.
- Press pump to dispense product into hands.
- Massage soap into hands and wrists, emphasizing back of hands, knuckles and cuticles.
- Rinse hands thoroughly and dry.

Store at 20 to 25°C (68 to 77°F).
Do not freeze.
Dispose in accordance with all applicable federal, state and local regulations.

Water, Cetrimonium Chloride, Lauryl/Myristyl Amidopropyl Amine Oxide, Glycerin, Cocamide DIPA, PEG-120 Methyl Glucose Dioleate, Sodium Chloride, Fragrance, Citric Acid, Methylchloroisothiazolinone, Methylisothiazolinone, Tetrasodium EDTA, Red 4, Yellow 5

Questions or comments?

Call 1-877-I-BUY-ZEP (1-877-428-9937)



ZEP PEAR ANTI-BACTERIAL

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-539
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
WATER (UNII: 059QF0KO0R)	
LAUROYL/MYRISTOYL AMIDOPROPYL AMINE OXIDE (UNII: HY9O6ZW9CY)	
COCO DIISOPROPANOLAMIDE (UNII: S485AM948Q)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-539-24	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/15/2017	

ZEP PEAR AB HS

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-393
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
COCO DIISOPROPANOLAMIDE (UNII: S485AM948Q)	
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
LAUROYL/MYRISTOYL AMIDOPROPYL AMINE OXIDE (UNII: HY9O6ZW9CY)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-393-01	6000 mL in 1 CASE; Type 0: Not a Combination Product	03/03/2017	
2	NDC:66949-393-11	6000 mL in 1 CASE; Type 0: Not a Combination Product	03/03/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/03/2017	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-393, 66949-539)

Revised: 12/2019 Zep Inc.