ELLE ANTIBACTERIAL HAND WASH- benzalkonium chloride liquid
Alta Sales 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.

ELLE ANTIBACTERIAL HAND WASH LAVENDER

DRUG FACTS

Active Ingredient: Benzalkonium Chloride 0.24%

Purpose: Antiseptic

Use For handwashing to decrease bacteria on the skin

Warnings For external use only

When using this product
Avoid contact with eyes. In case of eye contact, flush 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 into DRY hands
• Lather vigorously for at least 15 seconds
• Rinse and dry thoroughly

Inactive Ingredients: Water, Lauramine Oxide, Cetrimonium Chloride, Peg-150 Distearate, Coco Glucoside, Glyceryl Oleate, Tetrasodium Edta, Glycerin, Parfum, Citric Acid, Benzyl Alcohol, Methylchloroisothiazolinone, Methylisothiazolinone.

Dermatologically Tested

Paraben Free

Hypoallergenic

Cruelty Free

ELLE™ is used by Alta Sales Inc under license from HACHETTE FILIPACCHI PRESSE SA, Paris, France.
Made in Turkey
www.ellepersonalcare.us

Packaging

ELLE ANTIBACTERIAL HAND WASH

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79165-112
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.24 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
EDETATE SODIUM (UNII: MP1J8420LU)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79165-112-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/03/2020	

Labeler - Alta Sales Inc (080087931)

Revised: 9/2020 Alta Sales Inc