DONE HAND ANTIBACTERIAL SWEET CITRIC- benzalkonium chloride liquid
Olein Recovery Corporation

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.

Done Hand Soap Antibacterial Sweet Citric

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on ths skin.

Warnings

For external use only

When using this product,

avoid contact with eyes. In case of contact flush with water.

Stop use and ask a doctor if

irritation and redness develops.

Keep out of reach of children.

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

Directions

wash skin, rinse and dry thoroughly

Inactive ingredients

purified water, sodium laureth sulfate, sodium C14-16 olefin sulfonate, cocamidorpyopyl betaine, sodium chloride, glycerine, fragrance, tetrasodium EDTA, methylchloroisothiazolinone, Methylisothiazolinone, D&C orange No.4

Package Labeling:

DONE HAND ANTIBACTERIAL SWEET CITRIC

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77142-006
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)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
D&C ORANGE NO. 4 (UNII: Q1LIY3BO0U)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77142-006-00	296 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/15/2020	

Labeler - Olein Recovery Corporation (188543446)

Revised: 5/2020 Olein Recovery Corporation