SAVON DE PROVENCE- benzalkonium chloride liquid
HOME & BODY COMPANY

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.

Savon De Provence Antibacterial Hand Soap 17 fl.oz

Active ingredient

0.13% Benzalkonium Chloride

PURPOSE

Antibacterial

Use

Helps reduce bacteria on hands

WARNING

For external use only.
When using this product do not use in or near the eyes.
In case of contact, rinse eyes throughly with water.

Keep out of reach of children

Keep out of reach of children, except under adult supervision. If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

Wash hands and rinse

INACTIVE INGREDIENTS

Water (Aqua), Cocamidopropyl Betaine, Lauramine Oxide, Cellulose Gum, DMDM Hydantoin, Polysorbate 20, Fragrance, Glycerin

PRODUCT LABEL

SAVON DE PROVENCE

LAVENDER

ANTIBACTERIAL

HAND SOAP

KILLS 99.9% OF GERMS

With the power of **0.13 percent Benzalkonium Chloride**, this hand soap reduces the bacteria on the skin, gently cleans and moisturizes, and leaves the skin smelling great.

474 ml ℮ 16 fl.oz

SAVON DE PROVENCE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73746-021
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
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0KO0R)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73746-021-01	502 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/05/2020	

Marketing Information

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

Labeler - HOME & BODY COMPANY (081290720)

Registrant - HOME & BODY COMPANY (081290720)

Establishment

Name	Address	ID/FEI	Business Operations
HOME & BODY COMPANY		081290720	manufacture(73746-021)

Revised: 5/2020 HOME & BODY COMPANY