RENEW APOTHECARY LAVENDER SAGE- 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.

RENEW APOTHECARY LAVENDER SAGE

Active Ingredient

0.13% Benzalkonium Chloride

PURPOSE

Antibacterial

Use

Helps reduce bacterial 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.

DIRECTIONS

Wash hands and rinse

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.

INACTIVE INGREDIENTS

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

PRODUCT LABEL

RENEW APOTHECARY LAVENDER SAGE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73746-024
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**
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73746-024-01	1893 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
081290720		081290720	manufacture(73746-024)

Revised: 5/2020 HOME & BODY COMPANY