KOSETTE CARE HAND SANITIZER ALCOHOL-FREE- benzalkonium chloride gel
Innovation Labs, 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.

Kosette CARE Hand Sanitizer 500 mL

Active Ingredient

Benzalkonium Chloride 0.11%

Purpose

Antimicrobial

Uses

- To decrease bacteria on skin when soap and water is not available.
- Recommended for repeated use.

Warnings

- **For external use only.**
- **When using this product** avoid contact with eyes. In case of contact with eyes, flush thoroughly with water.
- **Stop use and ask a doctor if** skin irritation or redness develops and lasts.
- **Keep out of reach of children**. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough product in your palm to thoroughly cover your hands. Rub hands together until dry.
- Children under 6 years should be supervised when using this product.

Other information

- Read the directions and warnings before use.
- Store in a cool, dry and well-ventilated place.

Inactive ingredients

water(aqua), butylene glycol, propanediol, acrylates/c10-30 alkyl acrylate crosspolymer, caprylhydroxamic acid, caprylyl glyceryl ether, polyglyceryl-4 caprate, polyglyceryl-6 caprylate, potassium hydroxide, maltodextrin, moringa oleifera seed extract, frangrance(parfum)

Package Label - Principal Display Panel

Kosette

CARE

HAND SANITIZER

KILLS 99.9% OF GERMS

ALCOHOL-FREE

Safe for all, our gentle formula effectively disinfects hands while retaining skin's natural moisture leaving it soft and smooth.

16.9 fl.oz / 500 ml

KOSETTE CARE HAND SANITIZER ALCOHOL-FREE

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73284-420
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0011 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
MORINGA OLEIFERA SEED OIL (UNII: REM6A5QMC0)	
CAPRYLYL GLYCERYL ETHER (UNII: MI97BW74XZ)	
POLYGLYCERIN-6 (UNII: M51422LRAM)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
WATER (UNII: 059QF0KO0R)	
PROPANEDIOL (UNII: 5965N8W85T)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
POLYGLYCERYL-4 CAPRATE (UNII: 3N873UN885)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73284-420-40	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/01/2020	

Labeler - Innovation Labs, Inc. (117109069)

Establishment

Name	Address	ID/FEI	Business Operations
Innovation Labs, Inc.		117109069	manufacture(73284-420)

Revised: 7/2020 Innovation Labs, Inc.