KAZU HAND SANITIZING WIPES UNSCENTED- benzalkonium chloride cloth
N Bridge 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.

Benzalkonium Chloride

Water, glycerin, butylene glycol, aloe barbadensis leaf extract, 1,2-hexanediol, sodium hyaluronate, melaleuca alternifolia (tea tree) leaf extract, maltodextrin, camellia sinensis leaf extract

Antiseptic

keep out of reach of the children

wet hands thoroughly with product and allow to dry without wiping

for children under 6, use only under adult supervision

not recommended for infants

For external use only. Flammable. Keep away from heat or flame.

Do not use
- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

for external use only

KAZU HAND SANITIZING WIPES UNSCENTED

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76966-0020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.18 g in 50

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:76966-0020-1	50 in 1 PACKAGE; Type 0: Not a Combination Product	09/09/2020	

Marketing Information

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

Labeler - N Bridge Inc. (694892752)

Registrant - N Bridge Inc. (694892752)

Establishment

Name	Address	ID/FEI	Business Operations
Joy Life Co., Ltd.		689846233	manufacture(76966-0020)

Establishment

Name	Address	ID/FEI	Business Operations
N Bridge Inc.		694892752	label(76966-0020)