KUNDAL PURE AND SAFE SANITIZING WIPE PLUS (BENZALKONIUM CHLORIDE)- benzalkonium chloride liquid
THESKINFACTORY Co., Ltd.

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.

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benzalkonium chloride

Water
Aloe Barbadensis Leaf Extract
Camellia Sinensis Leaf Extract
Chamomilla Recutita (Matricaria) Extract
Phenoxyethanol
Cetylpyridinium Chloride
Caprylyl Glycol
Cocamidopropyl PG-Dimonium Chloride Phosphate
Citric Acid

hand sanitizing tissue to help reduce bacteria on the skin

keep out of reach of the children

pull ouy wipes and sanitize hands when necessary

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

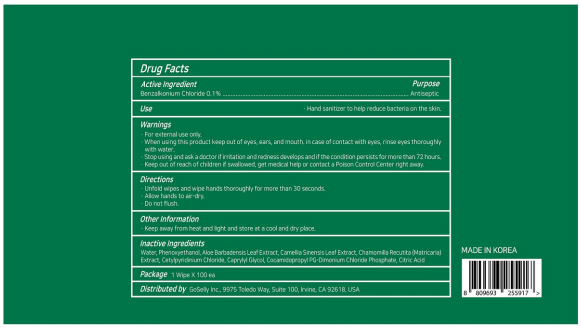
Do not use
- in children less than 2 months of age
- on open skin wounds

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



KUNDAL PURE AND SAFE SANITIZING WIPE PLUS (BENZALKONIUM CHLORIDE)

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74773-0026
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
MATRICARIA RECUTITA (UNII: G0R4UBI2ZZ)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74773-0026-1	7 g in 1 POUCH; 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 - THESKINFACTORY Co., Ltd. (694804099)

Registrant - THESKINFACTORY Co., Ltd. (694804099)

Establishment

Name	Address	ID/FEI	Business Operations
C&TECH CORPORATION		688204698	manufacture(74773-0026)

Establishment

Name	Address	ID/FEI	Business Operations
THESKINFACTORY Co., Ltd.		694804099	label(74773-0026)