UN CARD SANITIZER- benzalkonium liquid
Shelko Electronics Co.,Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

---

benzalkonium

Water

Antibacterial, anti-fungal, indoor air purification, humidity control and insect repellent, sick house syndrome, formaldehyde neutralization, forest bathing effect and stress relief, sterilization deodorization, hand sanitizer, super bacteria inhibition, sleep effect, house mite avoidance, allergy and atopy improvement

Do not drink.
Beware of fire.
Keep out of reach of children.
Avoid direct sunlight.
Do not use directly on the skin

Keep Out of Reach of Children

for topical use only

Dilute with 5g of oil in 500ml of water

UN CARD sanitizer

Drug Facts

Active Ingredient

benzalkonium

Uses

Antiviral, Antibacterial virus, Antibacterial, anti-fungal, indoor air purification, humidity control and insect repellent, sick house syndrome, formaldehyde neutralization, forest bathing effect and stress relief, sterilization deodorization, hand sanitizer, super bacteria inhibition, sleep effect, house mite avoidance, allergy and atopy improvement

Warnings

Do not drink.
Beware of fire.
Keep out of reach of children.
Avoid direct sunlight.
Do not use directly on the skin
Keep out of reach of children
If swallowed, get medical help or contact a person control center immediately

Directions

Apply where needed

Other Information

store between 20-25 °C (68-77 °F)
avoid freezing and excessive heat above 40 °C (104 °F)

Inactive Ingredients

Amino acid, water

UN CARD SANITIZER benzalkonium liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82419-0002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM (UNII: 7N6JUD5X6Y) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM	0.015 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:82419-0002-1	5 mL in 1 POUCH; Type 0: Not a Combination Product	11/28/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/28/2022	

Labeler - Shelko Electronics Co.,Ltd. (688446459)

Registrant - Shelko Electronics Co.,Ltd. (688446459)

Establishment

Name	Address	ID/FEI	Business Operations
Shelko Electronics Co.,Ltd.		688446459	manufacture(82419-0002)

Revised: 11/2022 Shelko Electronics Co.,Ltd.