YOUKE AN- benzalkonium chloride disinfectant spray liquid HUNAN ZONWE PHARMACEUTICAL 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 chloride

keep out of children

Kills 99.99% of germs

a little

It shall be directly sprayed onto skin, wounds, mucosa and infected parts. It can be applied several times a day to bring continuous antiseptic effects on skin surface for lasting protection.

Propylene glycol

Sodium dihydrogen phosphate dihydrate

Disodium hydrogen phosphate

Disodium edetate, dihydrate

water

- 1. This product is one kind of solution for external use, and is never for internal usage.
- 2. Activity of this product may be affected by anion surfactant or soap, and cannot be used together.
- 3.It is forbidden for people who are allergic to this product.
- 4. Turbidity or deposits may appear at a low temperature. It can be placed in warm waterfor heating, or used after it is dissolved through shaking.
- 5.It shall be sealed and kept in a dark and dry place.
- 6. Mucosa disinfection is only used for diagnosis and treatment by a medical institution.

佑可安® BENZALKONIUM CHLORIDE DISINFECTANT SPRAY 8mm **₩ 中原制薬** | 佑可安® **V 中原制薬**|佑可安® [Warnings] Active ingredient and its content 1. This product is one kind of solution for external use, and The primary active ingredient in this product is benzalkonium is never for internal usage. 2. Activity of this product may chloride, its content is be affected by anion surfactant or soap, and cannot be used **BENZALKONIUM CHLORIDE** BENZALKONIUM CHLORIDE 0.12%-0.14%. together. 3.It is forbidden for people who are allergic to this product. **DISINFECTANT SPRAY DISINFECTANT SPRAY** Type of targeted microorganism 4. Turbidity or deposits may appear at a low temperature. It can be placed in warm water for heating, or used after it is dissolved through shaking. 5.It shall be sealed and kept in a dark and dry place. 6.Mucosa disinfection is only **Directions** Hygienic hand disinfection; Skin and mucosa disinfection; Wound cleaning, disinfection and care; Disinfection after a minimally invasive facelit; Disinfection against bacteria or fungal infection; Skin disinfection through injection. used for diagnosis and treatment by a medical institution. [Hygiene Permit no.] (Hunan) Hygiene and disinfection permit No. (2015) 0007 [Reference standard] Q/CBCH001 [Size] 60ml/bottle [Form of drug] Spray [Term of validity] 24 months [Manufacturer] Hunan Zonwe Uses Uses It shall be directly sprayed onto skin, wounds, mucosa and infected parts. It can be applied several times a day to bring continuous antiseptic effects on skin surface for lasting protection. [Address of manufacturer] No.1 Jiangshan Road, Tianyuan District, Zhuzhou City, Hunan Province [Tel. No.] 0731-22841333 Features **Features** Kills 99.99% of germs Kills 99.99% of germs Alcohol-free & Non-irritating Applicable for both pregnancies and children Alcohol-free & Non-irritating Applicable for both pregnancies and children Filmed once applied. Essential for family and Hunan Zonwe Pharm Co., Ltd. Hunan Zonwe Pharm Co., Ltd. [Batch No] [Date of Manufacture] [Date of Expiry] 40mm 40mm 50mm 50mm

benzalkonium chloride disinfectant spray liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54749-001
Route of Administration	EXTRACORPOREAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKO NIUM (UNII: 7N6 JUD5X6 Y) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM	0.072 mg in 60 mL	

Inactive Ingredients			
Ingredient Name	Strength		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
SODIUM PHO SPHATE, MO NO BASIC, DIHYDRATE (UNII: 5QWK665956)			
SODIUM PHO SPHATE DIBASIC DIHYDRATE (UNII: 9425516 E2T)			
WATER (UNII: 059QF0KO0R)			
EDETATE DISO DIUM (UNII: 7FLD91C86K)			

Product Characteristics			
Color	white (transparent)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:54749-001-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		04/06/2020	

Labeler - Hunan zonwe pharmaceutical co., Ltd. (547493962)

Establishment				
Name	Address	ID/FEI	Business Operations	
HUNAN ZONWE PHARMACEUTICAL CO., LTD.		547493962	manufacture (54749-001)	