

SANITIZER- sanitizer liquid

KANGSHENG MEDICAL TECHNOLOGY (WENZHOU) 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](#).

Hexamine

Benzalkonium bromide

uses

To decrease bacteria on the skin that could cause disease

Recommended for repeat use

use without water

DIRECTIONS

Wet hands throughly with product and rub until dry without wiping

For children under 6 years old , use only under adult supervision

For exlernal use only

Keep out of reach of children, in case of accidental ingestion,seek professional assistance

Discontinue if skin becomes irritaled and ask a doctor

Don' t use it together with peroxide and detergent

Keep in a cool and dry place

Deionized water

Keep out of reach of children

Disinfection

Sterilization

No Rinseing



NON TOXIC DISINFECTANT

Use Without Water
Disinfectant

500ML

ACTIVE INGREDIENT:

- Hexamine : 0.23%–0.27%
- Benzalkonium bromide : 0.08%–0.1%

USESS:

- To decrease bacteria on the skin that could cause disease
- Recommended for repeat use
- Use without water

DIRECTIONS :

- Wet hands thoroughly with product and rub until dry without wiping
- For children under 6 years old , use only under adult supervision

WARNING :

- For external use only
- Keep out of reach of children, in case of accidental ingestion, seek professional assistance .
- Discontinue if skin becomes irritated and ask a doctor
- Don' t use it together with peroxide and detergent
- Keep in a cool and dry place

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SANITIZER

sanitizer liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:93068-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM BROMIDE (UNII: 151T1GQ42D) (BENZALKONIUM BROMIDE - UNII:151T1GQ42D)	BENZALKONIUM BROMIDE	0.5 g in 500 mL
METHENAMINE (UNII: J50OIX95QV) (METHENAMINE - UNII:J50OIX95QV)	METHENAMINE	1.35 g in 500 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:93068-001-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/30/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		11/30/2020		

Labeler - KANGSHENG MEDICAL TECHNOLOGY (WENZHOU) CO. , LTD (554543825)

Establishment			
Name	Address	ID/FEI	Business Operations
KANGSHENG MEDICAL TECHNOLOGY (WENZHOU) CO. , LTD		554543825	manufacture(93068-001)

Revised: 11/2020

KANGSHENG MEDICAL TECHNOLOGY (WENZHOU) CO. , LTD