ZORARI WATERLESS DISINFECTION- benzalkonium chloride gel
Zorin Pharmaceutical Technology (Hangzhou) 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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hand wash

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

USE

Disinfection gel helps decrease bacteria on skin. When water, soap and towel are not available.

Recommended for repeated use.

Warnings

For external use only.

When using this product

Do not apply around eyes. Do not use in ears and mouth. In case of contact with eyes, mouth, flush with water.

Stop use and ask a doctor if

- irritation and redness develops and persists for more than 72 hours.

Keep out of reach of children.

Children must be supervised in use of this product.

Directions

Squeeze small amount to your palms and thoroughly spread on both hands. Rub into skin until dry.

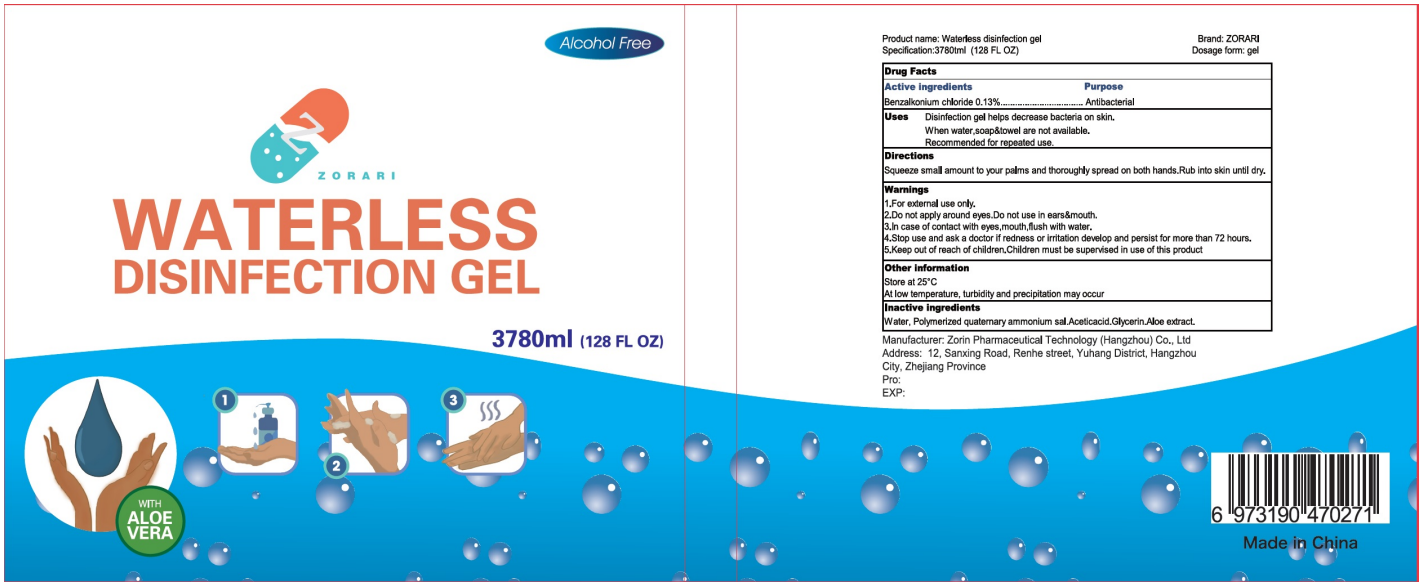
Other information:

Store at 25□□ At low temperature, turbidity and precipitation may occur.

Inactive ingredients

Water (Aqua), Polymerized quatermary ammonium sal.Aceticacid. Glycerin, Aloe extract.

package



ZORARI WATERLESS DISINFECTION

benzalkonium chloride gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75183-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75183-010-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/23/2020	
2	NDC:75183-010-02	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/23/2020	
3	NDC:75183-010-03	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/23/2020	

Labeler - Zorin Pharmaceutical Technology (Hangzhou) Co Ltd. (554529819)

Establishment

Name	Address	ID/FEI	Business Operations
Zorin Pharmaceutical Technology (Hangzhou) Co Ltd.		554529819	manufacture(75183-010)

Revised: 7/2020 Zorin Pharmaceutical Technology (Hangzhou) Co Ltd.