T3 WOUND WASH- benzalkonium chloride solution
Patient Focused Telehealth

T3 Wound Wash

Active Ingredient

Benzalkonium Chloride (0.1% by volume)

purpose

Antiseptic

Use

First aid to help prevent infection in minor diabetic ulcers, pressure ulcers, cuts, scrapes and burns.

Warnings

For external use only

When using this product

Avoid contact with eyes

stop use and ask a doctor if

the condition persists or gets worse. Do not use longer than 1 week unless directed by a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact a poison control center right away.

Directions

- Clean the affected area
- Apply a small amount of this product on the area 1 to 3 times daily
- Allow to dry
- Cover the affected area with sterile bandage if needed
- When bandaged, let dry first

Other Information

Store at 40 to 120 F

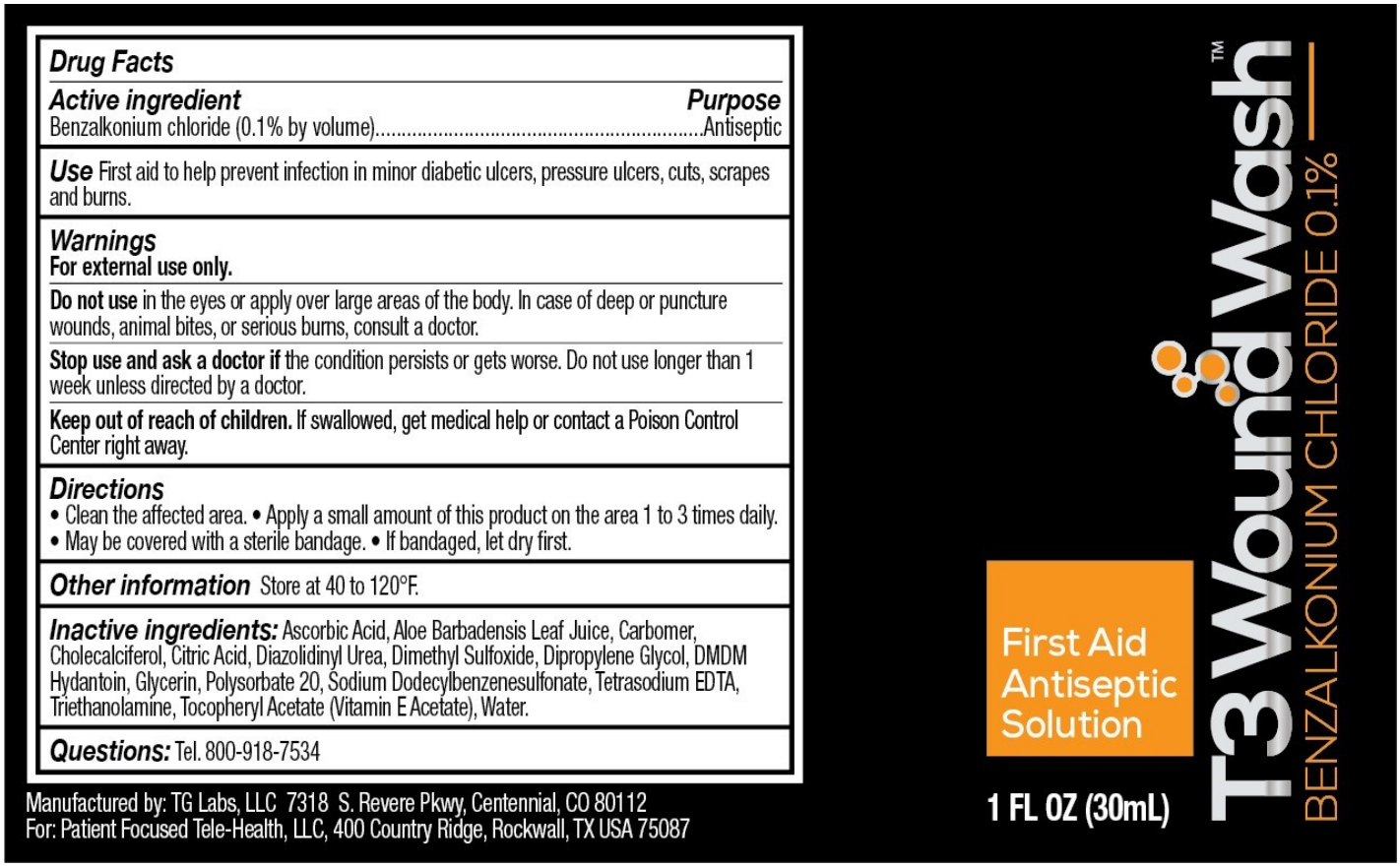
Inactive Ingredients

ascorbic acid, aloe barbadensis leaf juice, carbomer, cholecalciferol, citric acid, dimethyl sulfoxide, diazolidinyl urea, dipropylene glycol, dmdm hydantoin, glycerin, polysorbate 20, sodium dodecylbenzene sulfonate, tetrasodium EDTA, triethanolamine, tocopheryl acetate (vitamin E acetate), water

Questions

Tel. 800-918-7534 Email Address: info@patientfocusedtelehealth.com Manufactured by: TG Labs, LLC 7318 S Revere Pkwy, Centennial CO 80112 For: Patient focused tele-health LLC, 400 Country Ridge, Rockwall, TX USA 75087

Principal Display Panel



T3 Wound Wash

Benzalkonium Chloride 0.1%

First Aid Antiseptic Solution

1 FL OZ 30 mL

T3 WOUND WASH
benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72053-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.03 g in 30 mL

Inactive Ingredients

Ingredient Name	Strength
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM DODECYLBENZENESULFONATE (UNII: 554127163Y)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
EDETATE SODIUM (UNII: MP1J8420LU)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72053-004-04	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/25/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	07/25/2022	

Labeler - Patient Focused Telehealth (081008911)