

TRIDERGEL- benzalkonium chloride gel
Trifluent Pharma LLC

TRIDERGEL™

Drug Facts

Active Ingredients

Benzalkonium Chloride (0.1%)

Purpose

Antimicrobial

Indications for management of:

- Stage 1-2 open wounds
- Superficial stasis ulcers
- Superficial diabetic skin ulcers
- Post-surgical incisions
- First and second-degree burns
- Cuts
- Abrasions and skin irritations

Warnings

For external use only.

When using this product avoid contact with eyes; rinse immediately with cool tap water if contact occurs.

Do not use more than the recommended dose or longer than prescribed without checking with your physician. Talk with your physician before you use any other medicines or cleansers on your skin. Ask your physician before prolonged sun exposure. If your symptoms do not improve or they worsen, contact your physician

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- Clean the wound before each application
- Apply a thin layer of Tridergel™ wound care gel to the wound surface and skin immediately surrounding the wound
- Cover the wound as directed by your physician

Inactive Ingredients

Aloe Barbadensis Leaf Juice Extract, Aminomethyl Propanol, Butylparaben, Carbomer,

Decyl Glucoside, Ethylparaben, Glycerin, Hydrolyzed Eggshell Membrane, Isobutylparaben, Methylparaben, Phenoxyethanol, Propylparaben, Purified Water, Sodium Hyaluronate.

QUESTIONS OR COMMENTS

CALL (210) 944-6920

PRINCIPAL DISPLAY PANEL - 29.6 mL Bottle Label

TRIFLUENT
PHARMA®

TRIDERGEL™
WOUND CARE GEL

BENZALKONIUM CHLORIDE

IN A GEL CONTAINING
TRANSFORMING GROWTH
FACTOR - β

NDC 73352-520-01

1.0 FL OZ (29.6 mL)

PRODUCT SHOULD BE ADMINISTERED
UNDER THE SUPERVISION OF A
LICENSED MEDICAL PRACTITIONER

Drug Facts

ACTIVE INGREDIENT..... PURPOSE
Benzalkonium Chloride 0.1% ..Antimicrobial

INDICATIONS for management of:
• Stage 1-2 open wounds • Superficial stasis ulcers • Superficial diabetic skin ulcers • Post-surgical incisions • First and second-degree burns • Cuts • Abrasions and skin irritations

WARNINGS
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INACTIVE INGREDIENTS
Aloe Barbadosensis Leaf Juice Extract, Aminomethyl Propanol, Butylparaben, Carbomer, Decyl Glucoside, Ethylparaben, Glycerin, Hydrolyzed Eggshell Membrane, Isobutylparaben, Methylparaben, Phenoxyethanol, Propylparaben, Purified Water, Sodium Hyaluronate

KEEP OUT OF REACH OF CHILDREN
If swallowed, get medical help or contact a poison control center immediately.



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WOUND CARE GEL

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IN A GEL CONTAINING
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Drug Facts (Cont)

DIRECTIONS
Clean the wound before each application. Apply a thin layer of Tridergel™ wound care gel to the wound surface and skin immediately surrounding the wound. Cover the wound as directed by your physician.

Contains eggshell membrane powder - a non-allergenic organic source of solubilized protein tissue growth factors including collagen, elastin, hyaluronic acid, eight essential amino acids and transforming growth factor - β.

QUESTIONS OR COMMENTS:
CALL (210) 944-6920
MANUFACTURED FOR:
Trifluent Pharma, LLC
San Antonio, TX 78213
Made In The U.S.A.
520-01
REV01 06-08-2023

TRIDERGEL

benzalkonium chloride gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73352-520
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.001 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
EGG SHELL MEMBRANE (UNII: N7QBR4212V)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

METHYLPARABEN (UNII: A2I8C7HI9T)	
ETHYLPARABEN (UNII: 14255EXE39)	
ISOBUTYLPARABEN (UNII: 0QQJ25X58G)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73352-520-01	29.6 mL in 1 BOTTLE, WTH APPLICATOR; Type 0: Not a Combination Product	12/11/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M003	12/11/2023	

Labeler - Trifluent Pharma LLC (117167281)

Registrant - Blaine Labs, Inc (017314571)

Revised: 10/2025

Trifluent Pharma LLC