

NOXYDERM- benzalkonium chloride liquid
NOxy Health Products

Active Ingredient

Active Ingredient	Purpose
Benzalkonium Chloride 0.13%	First Aid Antiseptic

Purpose

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First aid antiseptic

Uses: first aid to help prevent infection in minor:

- cuts
- scrapes
- burns

Do not use

Do not use

- in the eyes
- over large areas of the body
- longer than 1 week unless directed by a doctor

Ask a doctor before use if you have

- deep or puncture wounds, animal bites, or serious burns

Warnings

For external use only

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For external use only

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact Poison Control right away

Instructions for use

Directions

- Clean the affected area
- Apply a small amount of this product on the area 1 to 3 times daily
- May be covered with a sterile bandage
- If bandaged, let dry first

Other safety information

Store at room temperature

Dosage

- Apply a small amount of this product on the area 1 to 3 times daily

Inactive ingredients

Sodium Nitrite

CoCo Betaine

Deionized Water

Citric Acid

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Questions

Questions? call 1-888-585-4120

NOxyDERM

Drug Facts

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Distributed by: Wasatch Product Development Inc.
427 W 11950
Draper, UT 84020

Learn more: noxyderm.com

NOxyDERM™
ADVANCED FIRST AID
HEALING FOAM

With Benzalkonium Chloride
Cleanses, Disinfects
and Promotes Healing
of Persistent Cuts,
Scrapes and Burns

Formulated with
Nitric Oxide,
nature's super
molecule

Alcohol Free • 3.4 fl oz.

NOXYDERM

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83823-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
COCO-BETAINE (UNII: 03DH2IZ3FY)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM NITRITE (UNII: M0KG633D4F)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83823-001-01	100 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/05/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/05/2024	

