

VIABECLINE FIRST AID ANTIBIOTIC- tetracycline hydrochloride ointment
VIADERMA II, INC.

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.

Viabecline First Aid Antibiotic

Drug Facts

Active ingredient (in each gram)

Tetracycline hydrochloride 30 mg

Purpose

First Aid Antibiotic

Use

First aid to help prevent skin infection in minor cuts, scrapes, and burns.

Warnings

For external use only. May be harmful if swallowed.

Allergy alert: Do not use if allergic to any ingredient listed on this label.

Do not use

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

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor

if condition persists or gets worse.

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 (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily.
- May be covered with a sterile bandage.

Other information

- Keep product refrigerated to preserve its effectiveness and color
- Stop use if product is misused: If the bottle is left open and/or if not refrigerated, the liquid will tend to turn black over time. Discard the product if the liquid turns black due to misuse.
- This product is an OTC antibiotic for human use.
- Contains no alcohol, no animal ingredients.
- Blended for typical skin color.
- May stain cloth.
- No claims regarding stem cell healing are implied for this product.

Inactive ingredients

ACETIC ACID, ASCORBIC ACID, CHLORHEXIDINE GLUCONATE, CHOLECALCIFEROL, DIMETHYL SULFOXIDE, DIPROPYLENE GLYCOL, GLUCONO DELTA LACTONE, GLYCERIN, HISTIDINE, HYDROXYETHYL CELLULOSE, MAGNESIUM STEARATE, METHYLPARABEN, SODIUM HYDROXIDE, SORBIC ACID, STEARIC ACID, WATER

Package Labeling:

Viabecline Topical Ointment
Patent pending
Alcohol free, no biological products
viadermalicensing.com NDC Code: 69006-001



0 04307 30111 2

Remove top of plastic box to read full contents of the enclosed Drug Facts label. Plastic box top is not sealed.

Viabecline

Topical Ointment

Refrigerate

First Aid
Antibiotic
Tetracycline
Hycholride, 3g

Drug Facts

Active ingredient (in each gram)	Purpose
Tetracycline hydrochloride 30 mg	First Aid Antibiotic

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Warnings

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Do not use ■ in the eyes ■ over large areas of the body ■ longer than 1 week unless directed by doctor
Ask a doctor before use if you have ■ deep or puncture wounds ■ animal bites ■ serious burns
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NET WT 28 ml (0.9 oz)

Side effects

Same as other tetracycline products
Report any side effects to ViaDerma, Inc.

4640 Admiralty Way, Suite 500, Marina Del Rey, CA 90292
Phone: (310) 496-5744 Fax: (310) 943-1457
Email:drotiko@yahoo.com

VIABECLINE FIRST AID ANTIBIOTIC

tetracycline hydrochloride ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69006-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TETRACYCLINE HYDROCHLORIDE (UNII: P6R62377KV) (TETRACYCLINE - UNII:F8VB5M810T)	TETRACYCLINE HYDROCHLORIDE	30 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ACETIC ACID (UNII: Q40Q9N063P)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
GLYCERIN (UNII: PDC6A3C0OX)	
HISTIDINE (UNII: 4QD397987E)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBIC ACID (UNII: X045WJ989B)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69006-005-00	1 in 1 BOX	05/13/2017	
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:69006-005-01	28 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	05/13/2017	

Labeler - VIADERMA II, INC. (079387584)

