TETRACYTE TOPICAL TETRACYCLINE HYDROCHLORIDE- tetracycline hydrochloride ointment VIADERMA DISTRIBITION INC

Tetracyte Topical Ointment Tetracycline Hydrochloride

Drug Facts

Active Ingredient (in each gram)

Tetracycline-HCl 30mg

Purpose

First Aid/Antibiotic

Indications

First aid to help prevent the risk of skin infection in minor cuts, scrapes, or 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 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 trauma

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

Stop use if product is misused

- this product is an OTC antibiotic for human use
- contains no alcohol, no animal ingredients
- blended for typical skin color
- may stain cloth
- no clamis regarding stem cell healing are implied for this product

Other Information

Keep refrigerated

Inactive Ingredients

acetic acid, ascorbic acid, chlorhexidine gluconate, cholecalciferol, dimethyl sulfoxide, dipropylene glycol, glucono delta lactone, glycerin, histidine, hydroxethyl-cellulose, magnesium stearate, methylparaben, sodium hydroxide, sorbic acid, water

Package Labeling:



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VIADERMA®

TETRACYTE

First Aid & Antibiotic

Topical Ointment

Tetracycline Hydrochloride HCI 3% (active ingredient)



Keep this and all drugs away from children, In case of accidental ingestion, seek medical assistance or contact a poison control immediately. **Side effect**: same as other tetracycline products.

NDC-79790-000-00



TETRACYTE

First Aid & Antibiotic

Topical Ointment

Tetracycline Hydrochloride HCI 3% (active ingredient)



0.5 FLOZ (15ml)

Licensed to ViaDerma Distribution by: ViaDerma, Inc. Distributed by: ViaDerma Distribution. Report any side effects to: ViaDerma Distribution 1050 E Flamingo Rd Suite 107-1360 Las Vegas, NV 89119 Tel: 1-800-585-8685 www.viadermainc.com

PATENT PENDING





TETRACYTE

First Aid & Antibiotic

Topical Ointment

Tetracycline Hydrochloride HCl 3% (active ingredient)



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TETRACYTE
First Aid & Antibiotic

Topical Ointment

Tetracycline
Hydrochloride HCl 3%
(active ingredient)
0.5 FLOZ (15ml)

KEEP THIS AND ALL DRUGS AWAY FROM CHILDREN, IN CASE OF ACCIDENTAL INGESTION SEEK MEDICAL ASSISTANCE OR CONTACT A POSION CONTROL IMMEDIATELY.

TETRACYTE TOPICAL TETRACYCLINE HYDROCHLORIDE

tetracycline hydrochloride ointment

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71262-009

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
TETRACYCLINE HYDROCHLORIDE (UNII: P6R62377KV) (TETRACYCLINE - UNII:F8VB5M810T)

Basis of Strength
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)				
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				

SORBIC ACID (UNII: X045WJ989B)	
WATER (UNII: 059QF0KO0R)	

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:71262-009- 01	1 in 1 CARTON	05/01/2024				
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product					

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M004	05/01/2024			

Labeler - VIADERMA DISTRIBITION INC (081113521)

Revised: 7/2024 VIADERMA DISTRIBITION INC