

TETRACYTE TOPICAL TETRACYCLINE HYDROCHLORIDE- tetracycline hydrochloride spray
VIADERMA DISTRIBUTION INC

Tetracyte Topical Spray 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 claims 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:

Tetracycline
Hydrochloride HCl 3%
(active ingredient)

Topical Spray

First Aid & Antibiotic

TETRACYTE™

VIA DERMA®
Pharma

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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.

VIA DERMA®
Pharma

TETRACYTE™

First Aid & Antibiotic

Topical Spray

Tetracycline
Hydrochloride HCl 3%
(active ingredient)



NDC-71262-010-01

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1.85 FL OZ (55ml)

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

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

NDC: 71262-010-01

VIADERMA[®]
Pharma

TETRACYTE[™]

First Aid & Antibiotic

Topical Spray

Tetracycline Hydrochloride HCl 3%
(active ingredient)

1.85 FL OZ (55ml)



TETRACYTE TOPICAL TETRACYCLINE HYDROCHLORIDE

tetracycline hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71262-010
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
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
ACETIC ACID (UNII: Q40Q9N063P)	
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: 70097M6130)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

SORBIC ACID (UNII: X045WJ989B)

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71262-010-01	1 in 1 CARTON	05/01/2024	
1		55 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 DISTRIBUTION INC (081113521)

Revised: 8/2024

VIADERMA DISTRIBUTION INC