# EQUALINE TRIPLE ANTIBIOTIC PLUS- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride ointment Supervalu 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.

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equaline™ triple antibiotic plus

**Drug Facts** 

Active ingredients (each gram contains)	Purpose
Bacitracin zinc 500 units	First aid antibiotic
Neomycin sulfate 3.5 mg	First aid antibiotic
Polymyxin B sulfate 10,000 units	First aid antibiotic
Pramoxine hydrochloride 10 mg	Topical analgesic

#### Uses

first aid to help prevent infection and for the temporary relief of pain or discomfort in

- minor cuts
- scrapes
- burns

#### Warnings

## For external use only

#### Do not use

- in the eyes
- over large areas of the body
- if you are allergic to any of the ingredients
- longer than 1 week

#### Ask a doctor before use

• on deep or puncture wounds, animal bites, or serious burns

# Stop use and ask a doctor if

- condition gets worse
- condition persists for more than 7 days
- condition clears up and occurs again within a few days
- a rash or other allergic reaction develops

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

#### **Directions**

Adults and children 2 years of age and older:

• clean the affected area and dry thoroughly

- 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

Children under 2 years of age: consult a doctor

#### Other information

- To open: unscrew cap, pull tab remove foil seal
- store at room temperature
- see carton or tube crimp for lot number and expiration date

## **Inactive ingredient**

white petrolatum

Distributed by Albertsons, Inc. Boise, Idaho 83726

# PRINCIPAL DISPLAY PANEL - 28.4g Tube Carton

compare to Neosporin® Plus active ingredients\*

egualine<sup>TM</sup>

maximum strength

## triple antibiotic plus ointment

polymyxin B sulfate ● bacitracin zinc neomycin sulfate ● pramoxine hydrochloride

first aid antibiotic/pain relief ointment

**NET WT 1 OZ (28.4g)** 



compare to Neosporin® Plus active ingredients"

maximum strength

# triple antibiotic plus ointment

polymyxin B sulfate · bacitracin zinc neomycin sulfate · pramoxine hydrochloride first aid antibiotic/pain relief ointment

first aid antibiotic plus topical pain reliever helps prevent infection in minor cuts, scrapes, and burns plus topical pain reliever

> compare to Neosporin® Plus active ingredients\*

> > maximum strength

# triple antibiotic plus ointment

neomycin sulfate · pramoxine hydrochloride first aid antibiotic/pain relief ointment

equaline polymyxin B sulfate · bacitracin zinc

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equaline"

NET WT 1 OZ (28.4g)





Children under 2 years of age: consult a doctor

- see carron or tube crimp for lot number and expiration date
  - Other information

Drug Facts (continued)

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Made in Canada.

Boise, Idaho 83726 Albertsons, Inc. Distributed by

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of the registered trademark Warner-Lambert Consumer Healthcare, owns or distributed by Si Jonpoud sitU.

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First aid antibiotic STOROLOUS DIE 1811esodind Active ingredients (each gram contains) Drug Facts

equaline

# triple antibiotic plus ointment

#### **EQUALINE** TRIPLE ANTIBIOTIC PLUS

bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-254
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Bacitracin zinc (UNII: 89 Y4M234ES) (Bacitracin - UNII:58 H6 RWO 52I)	Bacitracin	500 [iU] in 1 g	
Neomycin sulfate (UNII: 057Y626693) (Neomycin - UNII:I16QD7X297)	Neomycin	3.5 mg in 1 g	
Polymyxin B sulfate (UNII: 19371312D4) (Polymyxin B - UNII:J2VZ07J96K)	Polymyxin B	10000 [iU] in 1 g	
Pramoxine hydrochloride (UNII: 88AYB867L5) (Pramoxine - UNII:068X84E056)	Pramoxine hydrochloride	10 mg in 1 g	

Inactive Ingredients	
Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-254-20	1 in 1 CARTON		
1		28.4 g in 1 TUBE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	03/31/2012	

# Labeler - Supervalu Inc (006961411)

# Registrant - Taro Pharmaceuticals U.S.A., Inc. (145186370)

Establishment				
Name	Address	ID/FEI	Business Operations	
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(41163-254)	

Revised: 3/2013 Supervalu Inc