GOODSENSE ANTIBIOTIC PLUS PAIN RELIEF- neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride cream
Geiss, Destin & Dunn, 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.

GoodSense®
Antibiotic Cream + Pain Relief

Drug Facts

Active ingredients (in each gram)	*Purpose*
Neomycin sulfate 3.5 mg	First aid antibiotic
Polymyxin B sulfate 10,000 units	First aid antibiotic
Pramoxine hydrochloride 10 mg	External analgesic

Uses

First aid to help prevent infection and temporary relief of pain or discomfort in minor:

- cuts
- scrapes
- burns

Warnings

For external use only.

Not for prolonged use.

Do Not Use

- in the eyes, nose, or over large areas of the body
- if you are allergic to any of the ingredients
- longer than 1 week unless directed by a 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 for more than 7 days, gets worse, or clears up and occurs again within a few days
- a rash or other allergic reaction develops
- redness, irritation, swelling, or pain persists or increases

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away: 800-222-1222.

Directions

Adults and children 2 years of age and older:

- 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
- the affected area may be covered with a sterile bandage

Children under 2 years of age: ask a doctor

Other information

- store at 15 to 30°C (59 to 86°F)
- contents filled by weight, not volume
- **Tamper Evident:** DO NOT USE IF FLAPS ON CARTON ARE NOT SEALED OR MISSING

Inactive ingredients

cetearyl alcohol, methylparaben, mineral oil, petrolatum, polysorbate 60, propylene glycol, purified water

Questions or comments?

866-323-0107

Distributed by
Geiss, Destin & Dunn, Inc
Peachtree City, GA 30269

PRINCIPAL DISPLAY PANEL - 14.2 g Tube Carton

GOODSENSE®
NDC 50804-076-01

Maximum Strength

Antibiotic Cream + Pain Relief
Neomycin Sulfate • Polymyxin B Sulfate • Pramoxine HCl
First Aid Antibiotic Pain Relieving Cream

Compare to active ingredients in
Neosporin® + Pain Relief*

100%
SATISFACTION
GUARANTEED

NET WT 0.5 oz (14.2 g)


GOODSENSE®
Maximum Strength
Antibiotic Cream
+ Pain Relief
Neomycin Sulfate
Polymyxin B Sulfate • Pramoxine HCl
First Aid Antibiotic
Pain Relieving Cream
Antibiotic Cream + Pain Relief
Neomycin Sulfate • Polymyxin B Sulfate • Pramoxine HCl
First Aid Antibiotic Pain Relieving Cream
Helps Prevent Infection in Minor Cuts, Scrapes and Burns
Plus Maximum Strength Pain Relief.
PL9600B
V02
GOODSENSE®
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Antibiotic Cream
+ Pain Relief
*This product is not manufactured by or distributed by Johnson & Johnson Corporation, owner of the registered trademark Neosporin®+ Pain Relief.
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MADE IN THE USA
Distributed by
Geiss, Destin & Dunn, Inc
Peachtree City, GA 30269
www.valuelabels.com
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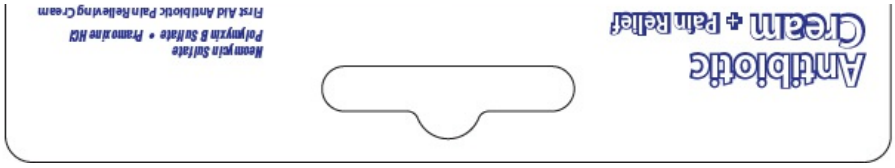

GOODSENSE ANTIBIOTIC PLUS PAIN RELIEF

neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-076
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
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**
methylparaben (UNII: A2I8C7HI9T)	
mineral oil (UNII: T5L8T28FGP)	
propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	
petrolatum (UNII: 4T6H12BN9U)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50804-076-01	1 in 1 CARTON	03/21/2012	
1		14.2 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part333B	03/21/2012	

Labeler - Geiss, Destin & Dunn, Inc. (076059836)

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

Establishment

Name	Address	ID/FEI	Business Operations
Natureplex LLC		062808196	MANUFACTURE(50804-076)