## ARTHRITIS RELIEF MAXIMUM STRENGTH- trolamin salicylate 10% cream Geiss, Destin & Dunn, Inc

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### Arthritis Relief, 10% Trolamine Salicylate

Trolamine Salicylate 10%

Topical Analgesic.

Temporarily relieves minor pain associated with arthritis, simple backache, muscle strains, sprains, and cramps.

### For external use only.

**Allergy Alert:** if prone to allergic reaction from aspirin or salicylates, consult a doctor before use.

When using this product use only as directed, avoid taking a bath or shower within 1 hour before or after applying, do not bandage or use with a heating pad, avoid contact with eyes or mucous membranes, do not apply to wounds or damaged skin.

**Stop use and ask a doctor** if condition worsens or symptoms persist for more than 7 days, if symptoms clear up and occur again within a few days, or if irritation develops.

Keep out of reach of children.

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

**Adults and children 12 years of age**: apply generously to the affected area, massage into painful area until thoroughly absorbed into the skin, repeat as necessary, but not more than 3 to 4 times daily. **Children under 12 years of age**: ask a doctor.

Aloe Barbadensis Leaf Juice, Cetyl Alcohol, Glycerin, Methylparaben, Mineral Oil, Potassium Phosphate, Propylparaben, Stearic Acid, Triethanolamine, Water.



10% Trolamine Salicylate

Pain Relieving Cream with Aloe

### GOODSENSE<sub>®</sub>

NDC 50804-207-03

**Odorless** 

# ritis Relief

10% Trolamine Salicylate

NET WT 3 OZ (85 g)

Pain Relieving Cream

\*Compare to active ingredient of Aspercreme®



with Aloe

EOLD

Distributed by: Geiss, Destin & Dunn, Inc. Peachtree City, GA 30269 www.valuelabels.com 1-866-696-0957



\*This product is not manufactured or distributed by the owners of the registered trademark Aspercreme".

Warning: This product contains chemicals known to the state of California to cause cancer.

**Inactive ingredients** Aloe Barbadensis Leaf Juice, Cetyl Alcohol, Glycerin, Methylparaben, Mineral Oil, Potassium Phosphate, Propylparaben, Stearic Acid, Triethanolamine, Water.

 repeat as necessary, but no more than 4 times daily Children 12 years or younger: ask a doctor. Adults and children over 12 years: • Apply generously to affected area • massage into painful area until thoroughly absorbed into skin

> Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately. It pregnant or breast-feeding, ask a health care professional before use.

> > · redness is present · irritation develops

Stop use and ask a doctor if • condition worsens • symptoms persist for more than 7 days or clear up and occur again within a few days

use with a heating pad • avoid contact with eyes or mucous membranes. • do not apply to wounds or damaged skin

When using this product • use only as directed • avoid taking a bath or shower within 1 hour before or after applying • do not bandage tightly or

Allergy alert: If prone to allergic reaction from aspirin or salicylates, consult a doctor before use.

For external use only. **Warnings** 

USES temporarily relieves minor pain associated with • arthritis • simple backache • muscle strains • bruises • cramps

Trolamine Salicylate 10%

Topical Analgesic Purpose Active ingredient

Drug Facts

### **ARTHRITIS RELIEF MAXIMUM STRENGTH**

trolamin salicylate 10% cream

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:50804-214

Route of Administration TOPICAL

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength TROLAMINE SALICYLATE (UNII: H804040BHD) (SALICYLIC ACID - UNII: 0414PZ4LPZ) TROLAMINE SALICYLATE TROLAMINE SALICYLATE

Inactive Ingredients				
Ingredient Name	Strength			
STEARIC ACID (UNII: 4ELV7Z65AP)				
CETYL ALCOHOL (UNII: 936JST6JCN)				
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
TROLAMINE (UNII: 903K93S3TK)				
METHYLPARABEN (UNII: A218C7H19T)				
MINERAL OIL (UNII: T5L8T28FGP)				
POTASSIUM PHOSPHATE, MONOBASIC (UNII: 4J9FJ0HL51)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:50804-214- 03	1 in 1 CARTON	06/27/2014			
1		85 g in 1 TUBE; Type 0: Not a Combination Product				

Marketing In	Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M017	06/27/2014			

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

Revised: 1/2024 Geiss, Destin & Dunn, Inc