ARTHRITIS CREAM- trolamine salicylate 10% cream
CVS

CVS Arthritis Cream

Trolamine Salicylate 10%

Topical analgesic

Temporarily relieves minor pain associated with:

Arthritis

Simple backache

muscle strains

sprains

bruises

For external use only

Allergy Alert:

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

When using the product

use only as directed

do not bandage tightly 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 doctor if

condition worsens

symptoms persist for more than 7 days or clear up and occur again within few days

redness is present

irritation develops

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 right away. (1-800-222-1222)

Directions:

adults and children over 12 years:

apply generously to affected area

massage into painful area until thoroughly absorbed into skin

repeat as necessary, but not more than 4 times daily

Children 12 years and younger: ask a doctor

Store at 20° - 25°C (68° - 77°F)

Aloe barbadensis leaf juice, cetyl alcohol, glycerin, mineral oil, phenoxyethanol, stearic acid, triethanolamine, water, xanthan gum

ARTHRITIS CREAM

trolamine salicylate 10% cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51316-990
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
		Basis of

Ingredient Name	Basis of Strength	Strength
TROLAMINE SALICYLATE (UNII: H8O4040BHD) (SALICYLIC ACID - UNII:O414PZ4LPZ)	TROLAMINE SALICYLATE	10 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TROLAMINE (UNII: 9O3K93S3TK)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-990-10	85 g in 1 TUBE; Type 0: Not a Combination Product	09/15/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	09/15/2022	

Labeler - CVS (062312574)

Revised: 12/2024 CVS