URIFLEX FAST ACTING ANALGESIC PAIN RELIEF- menthol cream
Rejuvica LLC

URIFLEX Fast Acting Analgesic Pain Relief Cream

Drug Facts

Active Ingredients

Menthol 4.50%

Purposes

External Analgesic

Uses:

For temporary relief of minor aches and pains associated with simple backache, arthritis, bruises, sprains, and strains.

Warnings:

For External Use Only. Avoid Contact With Eyes.

DO NOT Apply

to Open Wounds or Damaged Skin. If symptoms persist for more than seven days, discontinue use and consult physician.

KEEP OUT OF REACH OF CHILDREN.

If swallowed, consult physician. Do not bandage tightly.

If pregnant or breast feeding,

contact physician prior to use.

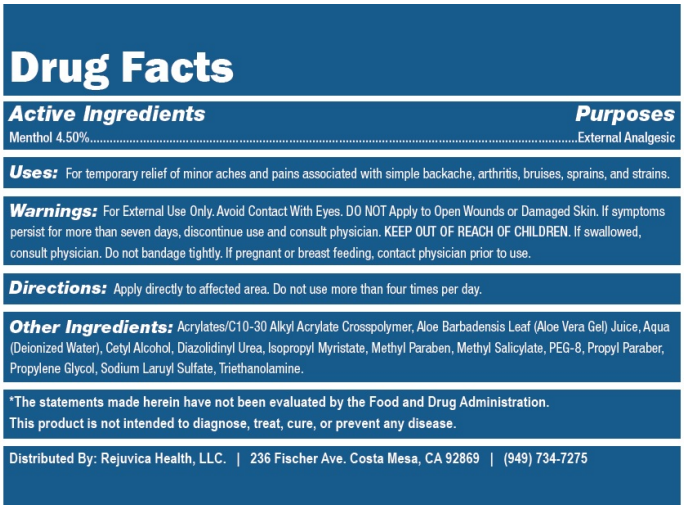
Directions:

Apply directly to affected area. Do not use more than four times per day.

Other Ingredients:

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Cetyl Alcohol, Diazolidinyl Urea, Isopropyl Myristate, Methyl Paraben, Methyl Salicylate, PEG-8, Propyl Paraber, Propylene Glycol, Sodium Laruyl Sulfate, Triethanolamine.

Package Labeling:



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*The statements made herein have not been evaluated by the Food and Drug Administration.
This product is not intended to diagnose, treat, cure, or prevent any disease.

Distributed By: Rejuvica Health, LLC. | 236 Fischer Ave. Costa Mesa, CA 92869 | (949) 734-7275

URIFLEX FAST ACTING ANALGESIC PAIN RELIEF

menthol cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73168-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	45 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73168-004-00	118 mL in 1 JAR; Type 0: Not a Combination Product	05/20/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/20/2019	

Labeler - Rejuvica LLC (062848492)

Revised: 11/2023 Rejuvica LLC