PAIN RELIEF PATCH- pain relief patch patch QUEST USA CORP.

78691-141

Active Ingredient

CAMPHOR (SYNTHETIC) 80mg MENTHOL 24mg NONIVAMIDE 16mg

Purpose

Topical analgesic

Use

For temporary relief of pains

Warnings

For external use only

Do not use

Do not get into eyes or on mucous membranes
Do not apply to wounds, damaged or irritated skin
Do not bandage or cover with any type of wrap except clothing
Do not use with a heating pad or apply external heat
Do not use 1 hour prior to bathing or within 30 minutes after bathing

When Using

When using this product Use only as directed

Stop Use

Stop use and ask a doctor if ·Condition worsens ·Severe skin irritation occurs ·Pain persists for more than 7 days ·Symptoms clear up and occur again within a few days

Ask Doctor

If pregnant or breast-feeding, or if you have sensitive skin, ask a healthcare professional before use. Children under 12 years, consult doctor before use

Keep Out Of Reach Of Children

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

Directions

Directions •Adults and children 12 years and over apply to affected area; change patch 1 to 2 times daily •Children under 12 years, consult doctor before use

How to apply:

·Clean and dry affected area

•Cut open pouch and remove patch

Remove protective film and apply directly to area of pain

Apply to affected area not more than 3 times daily

·Wash hands with soap after applying patch

·Reseal pouch containing unused patches

Other information

Other information

This product may cause allergic reaction on some individuals.Store in cool dry place away from direct sunlight.

Inactive ingredients

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Aluminum Glycinate, Ethylhexylglycerin, Eucalyptus Oil, Glycerin, Kaolin, Mentha Oil, Phenoxyethanol, Polyvinylpyrrolidone,Propylene Glycol, Sodium Carboxymethylcellulose, Sodium Polyacrylate, Sorbitol Solution, Tartaric Acid, Water

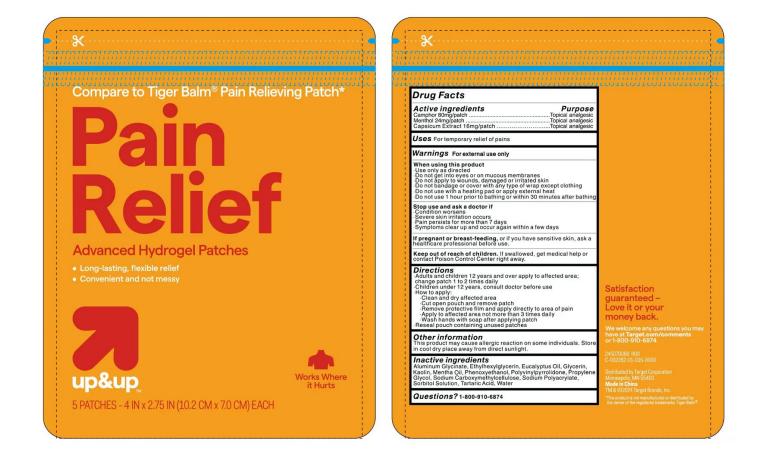
Questions

Questions?

1-800-910-6874

PRINCIPAL DISPLAY PANEL





PAIN RELIEF PATCH pain relief patch patch						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Sou	ltem Code (Source)		NDC:78691-141	
Route of Administration	TOPICAL					
Active Ingredient/Active	Moiety					
Ingredient Name			Basis of Strength		Strength	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII: 5TJD82A1ET)			CAMPHOR (SYNTHETIC)		80 mg in 100	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)			MENTHOL		24 mg in 100	
NONIVAMIDE (UNII: S846B8910R) (NONIVAMIDE - UNII:S846B8910R) NONIVAMIDE					16 mg in 100	
Inactive Ingredients						
Ingredient Name					Strength	
EUCALYPTUS OIL (UNII: 2R04ON	1662)					

PHENOXYETHANOL (UNII: HIE492ZZ3T)					
GLYCERIN (UNII: PDC6A3C0OX)					
SODIUM ACRYLATE (UNII: 7C98FKB43H)					
WATER (UNII: 059QF0K00R)					
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)					
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)					
KAOLIN (UNII: 24H4NWX5CO)					
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K6790BS311)					
SORBITOL SOLUTION (UNII: 8KW3E20702)					
TARTARIC ACID (UNII: W48881119H)					
POVIDONE (UNII: FZ989GH94E)					
Packaging					
# Item Code Package Description Marketing Start Marketi Date Da	-				
NDC:78691-141- 5 in 1 BOX; Type 0: Not a Combination 06/14/2024					
Marketing Information					
Marketing Application Number or Monograph Marketing Start Market	ting End				
	ate				

Labeler - QUEST USA CORP. (079869689)

Revised: 6/2024

QUEST USA CORP.