AMP RELIEF PAIN RELIEF CREAM- menthol cream
COASTAL FORMULAS LLC

AMP RELIEF Pain Relief Cream

Drug Facts

Active Ingredients

Menthol 8.00%

Purpose

Topical Analgesic

Uses:

- For the temporary relief of minor aches and pains of muscles and joints, associated with backache, arthritis, strains, bruises, and sprains

Warnings:

For external use only.

Do not use

- on damaged or broken skin

When using this product

- Avoid contact with the eyes.
- Do not bandage tightly.

Stop use and ask a doctor if

- rash or irritation develops and lasts
- condition worsens
- sysmptoms persist for more than 7 days
- clears up and occurs again within a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

If pregnant or breast-feeding,

ask a health professional before use

Directions:

- Adults and children 2 years of age and older: Apply to affected area not more than 3

to 4 times daily.
- Children under 2 years of age: Consult a doctor

Flammable:
- Keep away from exessive heat or open flame.

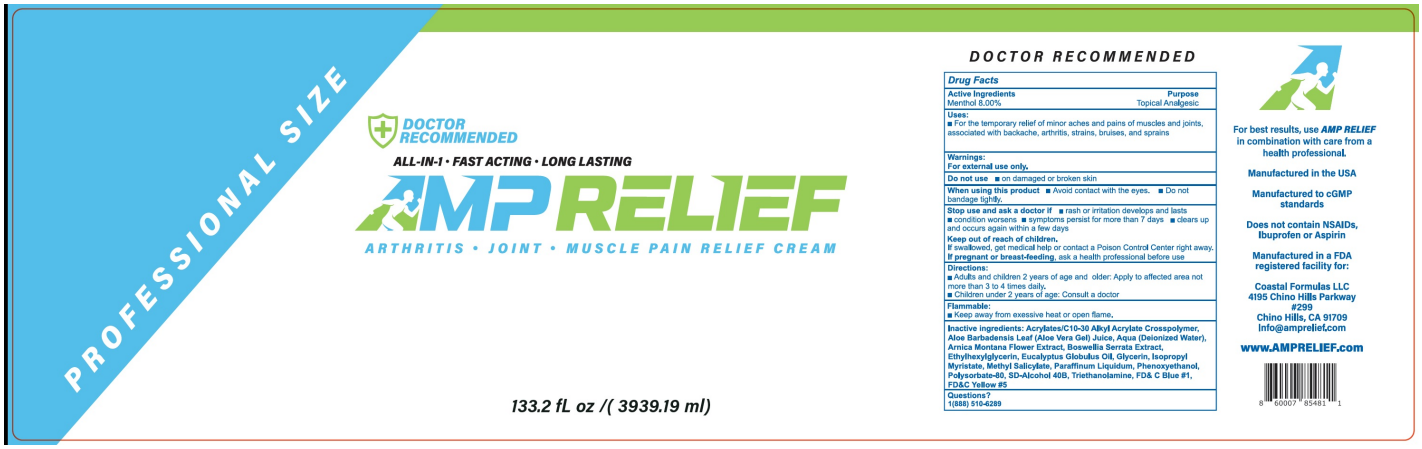
Inactive ingredients:

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Ethylhexylglycerin, Eucalyptus Globulus Oil, Glycerin, Isopropyl Myristate, Methyl Salicylate, Paraffinum Liquidum, Phenoxyethanol, Polysorbate-80, SD-Alcohol 40B, Triethanolamine, FD& C Blue #1, FD&C Yellow #5

Questions?

1(888) 510-6289

Package Labeling:



AMP RELIEF PAIN RELIEF CREAM

menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82560-283
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	80 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
MINERAL OIL (UNII: T5L8T28FGP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82560-283-00	3939.19 mL in 1 JUG; Type 0: Not a Combination Product	03/25/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/25/2022	

Labeler - COASTAL FORMULAS LLC (085954368)

Revised: 11/2023 COASTAL FORMULAS LLC