AMP RELIEF PAIN RELIEF CREAM- menthol cream
COASTAL FORMULAS LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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 thsi 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
- symptoms 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-feeing,

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 excessive heat or open flame.

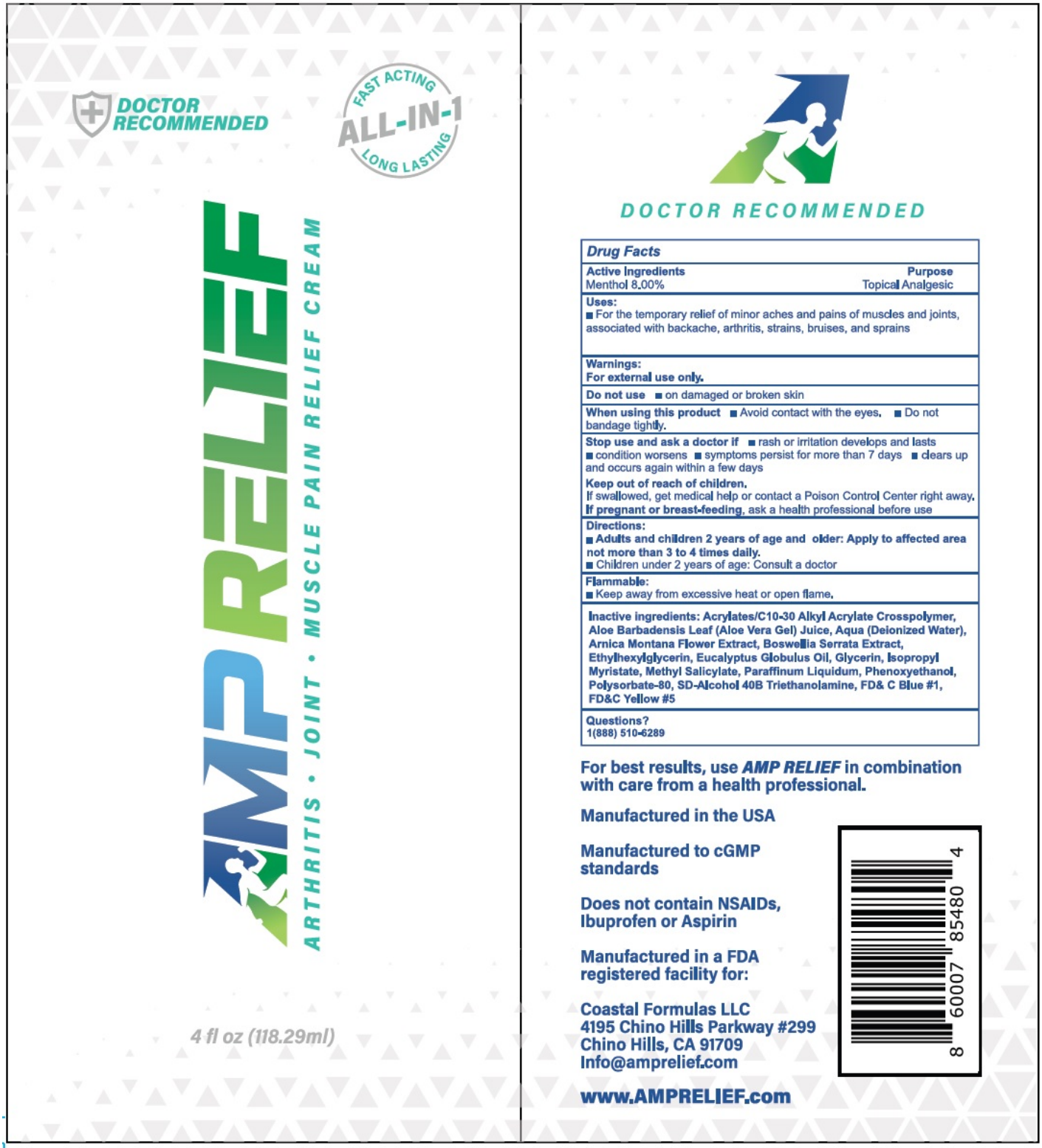
Inactive ingredients:

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:82560-273
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:82560-273-00	118.29 mL in 1 TUBE; Type 0: Not a Combination Product	02/15/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/15/2022	

Labeler - COASTAL FORMULAS LLC (085954368)

Revised: 2/2022 COASTAL FORMULAS LLC