MENTHOL PATCH- menthol patch patch
Guangzhou Hanhai Trading Co., Ltd

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.

Active Ingredient

Menthol 19.0%

Purpose

Pain Relief

Use

Use

Temporarily relieves muscle soreness and minor joints pain related to arthritis, backache, strains, sprains

Warnings

For external use only. Do not take it internally.

Keep Out Of Reach Of Children
Sensitive Skin

Do not use

Product shall not be eat

When Using

Use only as directedAvoid contacting with eyes or mucous membranesDo not apply to wounds or damaged skinDo not use with other ointments, creams, sprays, or linimentsDo not bandage or use with heating pad or devictStore in a cool dry place away from direct sunlight

Stop Use

In case of adverse reactions, please stop using this product.If the symptoms.

Ask Doctor

ask doctor in case of Sensitive Skin

Keep Oot Of Reach Of Children

Keep Out Of Reach Of Children

Directions

Take one piece of this product when needed, remove the waxpaper,
stick to the desired area and press the surrounding, replace it every 8 hours.
Children should be used under adult supervision.

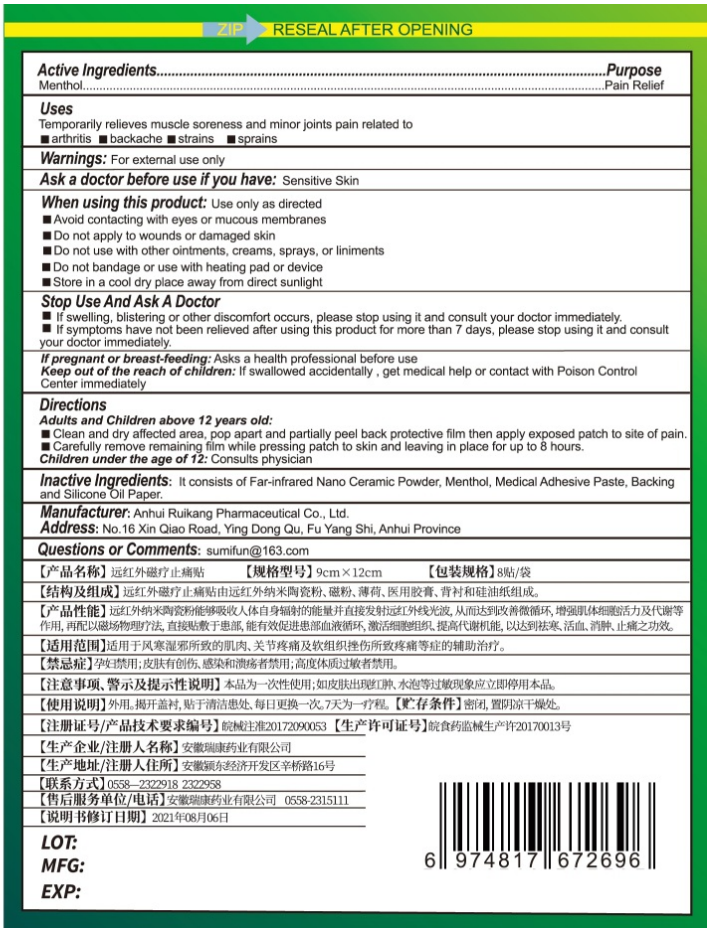
Other information

Store the product in a cool, dry and well-ventilated place
Avoid direct sunlight

Inactive ingredients

Far-infrared Nano Ceramic Powder,
Medical Adhesive Paste,
Backing
Silicone Oil Paper

PRINCIPAL DISPLAY PANEL



MENTHOL PATCH

menthol patch patch

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83675-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	19 g in 100

Inactive Ingredients

Ingredient Name	Strength
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
IVERMECTIN (UNII: 8883YP2R6D)	
DIMETHICONOL/TRIMETHYLSILOXYSILICATE CROSSPOLYMER (40/60 W/W; 1000000 PA.S) (UNII: 83D19O7250)	
BACKHOUSIA MYRTIFOLIA WHOLE (UNII: M5B93S0VJK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83675-010-01	8 in 1 BAG; Type 0: Not a Combination Product	09/19/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/19/2023	

Labeler - Guangzhou Hanhai Trading Co., Ltd (419707381)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Hanhai Trading Co., Ltd		419707381	label(83675-010) , manufacture(83675-010)

Revised: 9/2023 Guangzhou Hanhai Trading Co., Ltd