PAIN RELIEF PATCH- camphor, menthol, methyl salicylate patch
Anhui Miao De Tang Pharmaceutical Co., Ltd.

Pain Relief Patch

Pain Relief Patch

Camphor 3.1%

Menthol 6.0%

Methyl Salicylate 10.0%

For the temporary minor aches and pains of muscles and joints associated with:
arthritis
simple backache
strains
bruises
sprains

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For external use only

if prone to allergic reaction from aspirin or salicylates, consult a doctor before use.

on wounds or damaged skin

with a heating pad

at the same time as other topical analgesics

if you are allergic to any ingredients of this product

use only as directed

avoid contact with the eyes, mucous membranes, or rashes

do not bandage tightly

rash, itching, or excessive skin irritation develops

conditions worsen

symptoms persist for more than 7 days

symptoms clear up and occur again within a few days

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

Adults and children 12 years of age and over:

clean and dry the affected area

remove the patch from the film

apply to affected area not more than 3 to 4 times daily for 7 days

remove patch from the skin after at most 8-hour application

Children under 12 years of age:

consult a doctor

store in a clean, dry place outside of direct sunlight

protect from excessive moisture

hydrogenated poly

Pentaerythrityl tetra-di-t-butyl

hydroxyhydrocinnamate

petroleum

styrene/isoprene copolymer

PAIN RELIEF PATCH

camphor, menthol, methyl salicylate patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81484-801
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	3.1 mg in 100
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	10 mg in 100
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	6 mg in 100

Inactive Ingredients	
Ingredient Name	**Strength**
HYDROGENATED POLYISOBUTENE 8 (UNII: 7YR4ZFS62E)	
NAPHTHA (UNII: O3L624621X)	
ETHYL P-HYDROXYHYDROCINNAMATE (UNII: 8R568DFF4T)	
STYRENE (UNII: 44LJ2U959V)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:81484-801-02	15 in 1 BOX	07/26/2023	
1	NDC:81484-801-01	1 in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/26/2023	

Labeler - Anhui Miao De Tang Pharmaceutical Co., Ltd. (405744102)

Registrant - Anhui Miao De Tang Pharmaceutical Co., Ltd. (405744102)

Establishment

Name	Address	ID/FEI	Business Operations
Anhui Miao De Tang Pharmaceutical Co., Ltd.		405744102	manufacture(81484-801)

Revised: 11/2023

Anhui Miao De Tang Pharmaceutical Co., Ltd.