ASPERFLEX HOT- camphor, menthol, methyl salicylate patch
Akron Pharma Inc.

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.

AsperFlex
Tough on Pain
HOT Pain Relieving Patch

Active ingredients

Capsicum extract 0.025% as Capsaicin
Menthol 1.25%

Purpose

Topical analgesic

Uses

For temporary relief of minor aches and pains of muscles and joints associated with:

- arthritis
- simple backache
- strains
- bruises
- sprains

Warnings

For external use only

Do not use

- on wounds or damaged skin
- with a heating pad
- if you are allergic to any ingredients of this product

When using this product

- use only as directed
- avoid contact with the eyes, mucous membranes or rashes
- do not bandage tightly

Stop use and ask a doctor if

- rash, itching or excessive skin irritation develops
- contisions worsen
- symptoms persist for more than 7 days

- symptoms clear up and occur again within a few days

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222).

Directions

Adults and children 12 years of age and over:
- clean and dry affected area
- remove patch from film
- apply to affected area not more than 3 to 4 times daily
- remove patch from the skin after at most 8-hour application

Children under 12 years of age: consult a doctor

Other information
- store at room temperature 15°-30°C (59°-86°F).

Inactive ingredients

aluminum glycinate, propylene glycol, sodium acrylate/sodium acryloyldimethyl taurate copolymer, tartaric acid, 2,4-Imidazolidinedione, disodium edta, water, glycerin

Questions or comments?

Call toll-free 1-877-255-6999.

Manufactured for:
Akron Pharma, Inc.
Fairfield, NJ 07004

www.akronpharma.com

ASPERFLEX HOT

camphor, menthol, methyl salicylate patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71399-4439
Route of Administration	TOPICAL, PERCUTANEOUS, TRANSDERMAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CAPSICUM (UNII: 00UK7646FG) (CAPSICUM - UNII:00UK7646FG)	CAPSICUM	0.025 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.25 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM ACRYLATE (UNII: 7C98FKB43H)	

TARTARIC ACID (UNII: W4888I119H)	
HYDANTOIN (UNII: I6208298TA)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-4439-6	6 in 1 BOX	11/04/2021	
1		7 g in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/04/2021	

Labeler - Akron Pharma Inc. (067878881)

Revised: 2/2023 Akron Pharma Inc.