

ASPERFLEX ORIGINAL MAXIMUM STRENGTH- menthol 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
Maximum Strength
Tough on Pain
Menthol 7.5% Original

DRUG FACTS

Active Ingredient

Menthol 7.5%

Purpose

Topical analgesic

Uses

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

■ arthritis ■ simple backache ■ strains ■ sprains ■ bruises

Warnings

For external use only

Do not use

- on wounds, damaged, broken or irritated skin
- with a heating pad or apply local heat to the area of use

When using this product

- use only as directed. Read and follow all directions and warnings on this carton.
- do not allow contact with
- the eyes and mucous membranes
- rare cases of serious burns have been reported with products of this type
- do not apply to wounds or damaged, broken or irritated skin
- do not bandage tightly or apply local heat (such as heating pads) or a medicated patch to the area of use
- do not use at the same time as other topical analgesics

- a transient burning sensation may occur upon application but generally disappears in several days
- avoid applying into skin folds <do not use more than 4 patches at a time

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days
- severe burning sensation, redness or irritation develop
- you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied

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 older:

- gently fold the patch in half to remove center section of film backing. Apply the exposed adhesive portion to the site of pain.
- remove remaining film backing from both sides and finish applying to skin
- apply to affected area not more than 3 to 4 times daily

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.


18.00 mm

Maximum Strength
NDC 71399-4461-5



TOUGH ON PAIN

ORIGINAL

Menthol 7.5%

POWERFUL TARGETED PAIN RELIEF PATCH

INSTANT RELIEF

- ✓ Powerful Targeted Relief
- ✓ Icy to Dull, Hot to Relax
- ✓ Comfortable Fabric
- ✓ Stay in Place





LARGE

Contains 5 Patches

in 1 Resealable Pouch
3-15/16" X 7-13/16" (10 cm x 20 cm) each

AsperFlex™



Instructions to apply





Drug Facts	
Active ingredient	Purpose
Menthol 7.5%	Topical analgesic
Uses	
temporarily relieves minor pain associated with:	
■ arthritis ■ simple backache ■ muscle strains ■ sprains ■ bruises ■ cramps	
Warnings	
For external use only.	
Do not use	
■ on large areas of the body or on cut, irritated or swollen skin ■ on puncture wounds	
■ for more than one week without consulting a doctor	
When using this product	
■ use only as directed. Read and follow all directions and warnings on this carton. ■ do not allow contact with the eyes and mucous membranes ■ rare cases of serious burns have been reported with products of this type	
■ do not apply to wounds or damaged, broken or irritated skin ■ do not bandage tightly or apply local heat (such as heating pads) or a medicated patch to the area of use ■ do not use at the same time as other topical analgesics ■ a transient burning sensation may occur upon application but generally disappears in several days	
■ avoid applying into skin folds ■ do not use more than 4 patches at a time	
Stop use and ask a doctor if	
■ condition worsens ■ redness is present ■ irritation develops ■ symptoms persist for more than 7 days or clear up and occur again within a few days ■ you experience signs of skin injury, such as pain, swelling, or blistering where the product was applied	
If pregnant or breast-feeding, ask a health professional before use.	
Keep out of reach of children & pet 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 older:	
■ gently fold the patch in half to remove center section of film backing. Apply the exposed adhesive portion to the site of pain ■ remove remaining film backing from both sides and finish applying to skin	
■ use 1 patch for up to 8 hours, once a day	
children under 12 years of age: ask 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

Rev.:11/21



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ASPERFLEX ORIGINAL MAXIMUM STRENGTH

menthol patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-4461
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	210 mg

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:71399-4461-5	5 in 1 CARTON	11/04/2021	
1		1 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.