

PAIN RELIEF PATCHES- camphor, menthol, methyl salicylate patch

Harris Teeter

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.

Medicated Relief Patches

Active Ingredients

Active Ingredients Purpose

Camphor 3.1%Topical Analgesic

Menthol 6.0%Topical Analgesic

Methyl Salicylate 10.0%Topical Analgesic

For External Use Only

Allergy alert: If prone to allergic reaction from aspirin or salicylates, consult a doctor before use

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 eyes, mucous membranes or rashes
- Do not bandage tightly
- Do not use at the same time as other topical analgesics
- Dispose of used patch in manner that keeps product away from children and pets. Used patches still contain the drug product that can produce serious adverse effects if a child or pet chews or ingests this patch.

Stop use and consult a doctor

- If rash, itching or excessive skin irritation develops
- If condition worsens
- if symptoms last more than 7 days or clear up and occur again with a few days

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Uses

For temporary relief of minor aches and pains of muscles and joints associated with arthritis, simple backache, strains, sprains, and bruises.

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 physician.

Inactive Ingredients

Hydrogenated Poly, Pentaerythrityl Tetra-di-t-butyl Hydroxyhydrocinnamate, Petroleum, Styrene / Isoprene Copolymer

Indication and Usage

For temporary relief of minor aches and pains.

Pain Relief Patch Label

Box of 60

NDC:72036-907-60

Medicated Pain Relief Patch

NDC 72036-907-60

Compare to Saloquist® Pain Relieving Patch active ingredient*

Medicated Pain Relief Patch

Medicated Pain Relief Patch

Medicated Pain Relief Patch

Upper Back

Lower Back

*This product is not manufactured or distributed by Harris Teeter America, Inc., a subsidiary of Hainbow Pharmaceutical Co., Inc., distributor of Saloquist® Pain Relieving Patch.

Neck, Shoulder, Back & Joint

Topical Analgesic

Temporary Relief of Aches & Pains

Targeted Single-Use Patch

60 PATCHES
2.83 in. x 1.81 in.
(7.2 cm x 4.6 cm)

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Support your local schools. Together in Education programs details available at any Harris Teeter or harristeeter.com.

yourplusguarantee.
If you are not 100% satisfied with the product, simply return it with your MC card for a full refund. Plus we will replace it with a like item at your choice. For full details: 1-800-422-4111 at harristeeter.com.

Drug Facts

Active ingredient Purpose
Camphor 3.1% Topical analgesic
Menthol 6.03% Topical analgesic
Methyl Salicylate 10.0% Topical analgesic

Uses For temporary relief of minor aches and pains of muscles and joints associated with arthritis, simple backache, strains, sprains and bruises.

Warnings
For External Use Only
Allergy alert: If prone to allergic reaction from aspirin or salicylates, consult a doctor before use.
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 eyes, mucous membranes or noses ■ Do not bandage tightly ■ Do not use at the same time as other topical analgesics ■ Dispose of used patch in manner that always keeps product away from children and pets. Used patches still contain the drug product that can produce serious adverse effects if a child or pet chews or ingests this patch.
Stop use and consult a doctor if ■ Rash, itching or excessive skin irritation develops ■ Condition worsens ■ Symptoms last more than 7 days or 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.

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 physician.

Other information
Store in a clean, dry place outside of direct sunlight. Protect from excessive moisture.

Inactive ingredients
Hydrogenated Poly, Styrene/Isoprene Copolymer, Petroleum, Pentaerythrityl Tetra-di-t-butyl Hydroxyhydrocinnamate

Questions?
Customer Care Help Line
866-326-1313
Mon-Fri 8:30 am-4:30 pm CST

PROUDLY DISTRIBUTED BY:
HARRIS TEETER, LLC
MATTHEWS, NC 28105
MADE IN CHINA
#92-82818 05/18

0 72036 72992 7

LOT EXP

Other Information: Store in clean, dry place outside of direct sunlight. Protect from excessive moisture.

DIST. BY
HARRIS TEETER, LLC
MATTHEWS, NC 28105

PAIN RELIEF PATCHES

camphor, menthol, methyl salicylate patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72036-907
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 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	6 g in 100 g
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	10 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
HYDROGENATED POLYDECENE (550 MW) (UNII: U333RI6EB7)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	
LIQUID PETROLEUM (UNII: 6ZAE7X688J)	
STYRENE/ACRYLAMIDE COPOLYMER (MW 500000) (UNII: 5Z4DPO246A)	

Product Characteristics

Color		Score	
Shape	SQUARE	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72036-907-60	60 in 1 BOX	05/01/2018	
1		9 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	05/01/2018	

Labeler - Harris Teeter (047279351)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Aqua Gel Biotech Co.,Ltd.		529 128 763	manufacture(72036-907)

Revised: 5/2018

Harris Teeter