

AUVON PAIN RELIEF- menthol patch
SHENZHEN YUWEN E-COMMERCE CO., LTD.

83391-002

MENTHOL AUVON Pain Relief Patch

Drug Facts

Active ingredients

Menthol 5%

Purposes

Topical Analgesic

Use

Temporary relief from minor aches and pains of sore muscles and joints

Warnings

For external use only

Do not use

- If you are allergic to the listed ingredients
- If you are pregnant or breast feeding
- If you are under 12 years of age
- On wounds, cuts, damaged/broken/irritated skin
- On eyes or mucous membranes
- With heating pads/devices or wrap with a bandage
- With other topical analgesics
- If the package arrives damaged or opened.

When using this product

- Use only as directed
- Read and follow all directions and warnings on this label

Stop use and ask a doctor if

You are experiencing pain, swelling or blistering

Redness is present or irritation develops

Symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children and pets.

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

Directions

- Clean and dry the patch application area (no wound or hair), pull to separate the film, peel off one side of the film, apply the exposed patch to the skin, peel off the other side of the film, and press the patch firmly to the skin.
- Do not repeatedly use or repeatedly reapply the patch as far as possible
- Cut first if used on joints
- Use in the affected area no more than 4 times daily
- Wash hands with cool water after use

Storage

- Avoid storing patches in direct sunlight
- Protect patches from excessive moisture

Inactive ingredients

Glycerin, sodium polyacrylate, dihydroxyaluminum aminoacetate, methylparaben, propylparaben, polyacrylic acid, Kaolin, polysorbate 80, propylene glycol, tartaric acid, PVP, titanium dioxide, water, CMC, mineral oil, petrolatum,



AUVON PAIN RELIEF

menthol patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83391-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	470 mg

Inactive Ingredients

Ingredient Name	Strength
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZ17B19X)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A218C7HI9T)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
TARTARIC ACID (UNII: W4888119H)
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)
WATER (UNII: 059QF0KO0R)
PROPYLPARABEN (UNII: Z8IX2SC1OH)
KAOLIN (UNII: 24H4NWX5CO)
POLYACRYLIC ACID (250000 MW) (UNII: 9G2MAD7J6W)
PETROLATUM (UNII: 4T6H12BN9U)
MINERAL OIL (UNII: T5L8T28FGP)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83391-002-01	1 in 1 POUCH; Type 0: Not a Combination Product	07/05/2023	
2	NDC:83391-002-02	8 in 1 BOX; Type 0: Not a Combination Product	07/05/2023	
3	NDC:83391-002-03	15 in 1 BOX; Type 0: Not a Combination Product	07/05/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/05/2023	

Labeler - SHENZHEN YUWEN E-COMMERCE CO., LTD. (544559614)

Revised: 12/2024

SHENZHEN YUWEN E-COMMERCE CO., LTD.