GOODSENSE EXTRA STRENGTH COLD AND HOT MEDICATED- menthol patch
Geiss, Destin and Dunn, Inc.

GoodSense Cold & Hot Medicated Patches 5ct Small 125A

Active ingredient **Purpose**

Menthol 5%.....Topical analgesic

Uses

temporariliy relieves minor aches and pains of muscles and joints associated with:
- simple backache
- arthritis
- strains
- burises
- sprains

Warnings

For external use only

Do not use
- on wonunds or damaged skin
- with a heating pad
- on a child under 12 years of age with arthritis-like conditions

Ask a doctor before use if you have
- redness over the affected area

When using this product:
- use only as directed
- avoid contact with eyes or mucous membranes
- do not bandage tightly
- discontinue use at least 1 hour before a bath or shower
- do not use immediately after a bath or shower

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
- excessive skin irritation occurs

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 (1-800-222-1222) right away.

Directions
- open pouch and remove patch
- carefully peel off protective backing and apply sticky side to affected area
- adults and children 12 years of age or older
- do not wear patch for more than 8 hours

- apply to affected area no more than 3 times daily
- children under 12 years of age: consult a doctor

Other information

- Store at room temperature 20-25°C (68-77°F)

Inactive ingredients

alcohol, aluminum glycinate, glycerin, kaolin, methylparaben, polysorbate 80, propylparaben, purified water, sodium polyacrylate, sorbitan monooleate, tartaric acid, titanium dioxide

Distributed By:

Perrigo Direct, Inc.

Peachtree City, Ga 30269

www.PerrigoDirect.com

1-866-696-0957

Made in China

GOODSENSE EXTRA STRENGTH COLD AND HOT MEDICATED

menthol patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-126
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	50 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
KAOLIN (UNII: 24H4NWX5CO)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
TARTARIC ACID (UNII: W4888I119H)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50804-126-05	1 in 1 CARTON	12/26/2018	
1		5 in 1 POUCH; 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	12/26/2018	

Labeler - Geiss, Destin and Dunn, Inc. (076059836)

Revised: 10/2025 Geiss, Destin and Dunn, Inc.