UP AND UP COLD AND HOT MEDICATED- menthol patch
TARGET CORP

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.

Up & Up Cold & Hot Medicated Patch 5ct Large 678 ZDP

Active ingredient Purpose

Menthol 5%.....Topical analgesic

Uses
- temporarily relieves minor aches and pains of muscles and joints associated with:
- simple backache
- arthritis
- strains
- bruises
- sprains

Warnings

For external use only

Do not use
- on wounds 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 and 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

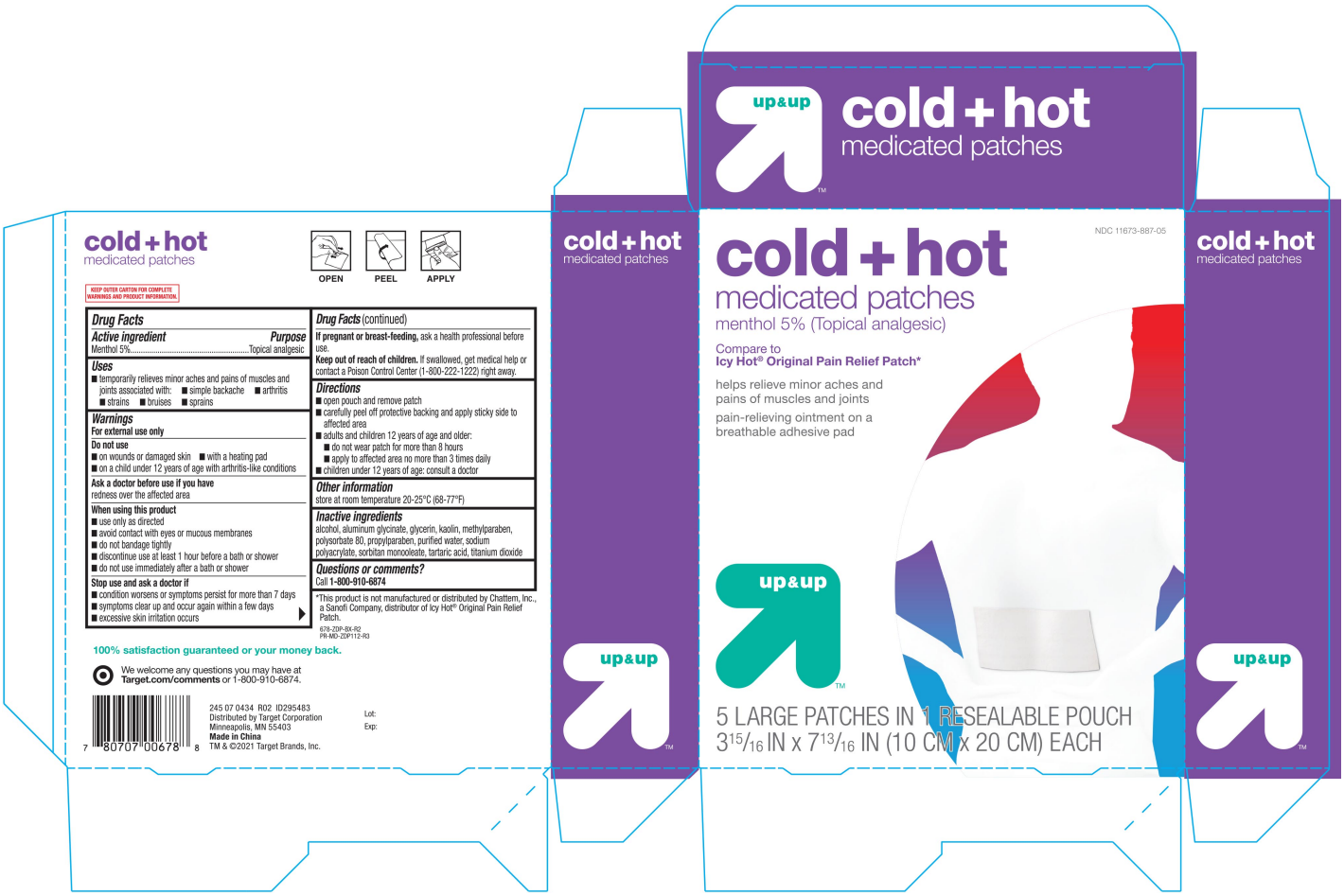
Other information
- store at room temperature 20-25°C (68-77°F)

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

Distributed by:

Target Corporation

Made in China



UP AND UP COLD AND HOT MEDICATED

menthol patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-887
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
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
KAOLIN (UNII: 24H4NWX5CO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
ALCOHOL (UNII: 3K9958V90M)	
TARTARIC ACID (UNII: W4888I119H)	
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JNI2J)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIHYDROXYALUMINUM AMINOACETATE ANHYDROUS (UNII: 1K713C615K)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-887-05	1 in 1 CARTON	11/11/2021	
1		5 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	06/01/2021	

Labeler - TARGET CORP (006961700)

Revised: 12/2022 TARGET CORP