HOT/COLD MENTHOL 16% ROLL-ON- hot/cold menthol 16% roll-on liquid Humco Holding Group, Inc

Private Label Hot/Cold Menthol 16% Roll-On

Adults and children over 12 yrs of age: apply to the affected area no more than 3 to 4 times daily

IF MEDICINE COMES IN CONTACT WITH HANDS, WASH WITH SOAP AND WATER.

When using this product: use only as directed. Do not bandage tightly or use with a heating pad. Avoid contact with eyes and mucous membranes. Do not apply to wounds or damaged, broken or irritated skin. If you experience pain, swelling, or blistering of the skin where an OTC topical muscle and joint pain reliever was applied, stop using the product and seek professional attention immediately. These products produce local warmth or coolness.

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Capsaicin, Glycerin, Isopropyl Myristate, Propylene Glycol, SD Alcohol 40 (30%Z), Triethanolamine, Purified Water.

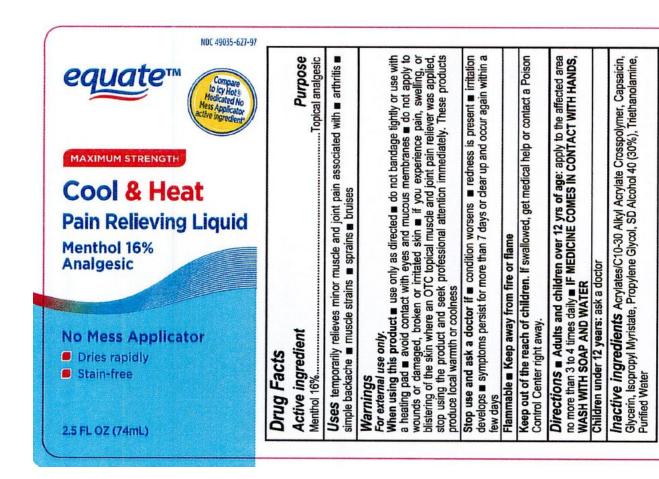
Temporarily relieves minor muscle and joint pain associated with arthritis, simple backache, muscle strains, sprains, bruises.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Topical analgesic

Menthol 16%

Equate Label



Not For Individual Retail Sale,

Satisfaction guaranteed - Or we'll replace it or give you your money back. For questions or comments please call 1-888-287-1915.

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716

"This product is not manufactured or distributed by Chattern, Inc.,
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HOT/COLD MENTHOL 16% ROLL-ON

hot/cold menthol 16% roll-on liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0395-9131
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
M	IENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	16 mg in 100 mL

Inactive Ingredients			
Strength			
TROLAMINE (UNII: 903K93S3TK)			
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)			

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:0395- 9131-97	74 mL in 1 CONTAINER; Type 0: Not a Combination Product	12/21/2018	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	12/21/2018		

Labeler - Humco Holding Group, Inc (825672884)

Registrant - Pharma Nobis, LLC (118564114)

Establishment			
Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	label(0395-9131) , pack(0395-9131) , analysis(0395-9131) , manufacture(0395-9131)

Revised: 12/2023 Humco Holding Group, Inc