DR GOLDSONS PAIN- menthol 8%, methyl salicylate 30% cream Derma Care Research Labs, LLC

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.

Dr Goldsons Pain Relief Cream

Menthol 8%, Methyl Salicylate 30%

External Analgesic

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

For external use only. Do not use on wounds or damaged skin or with a heating pad. When using this product avoid contact with eyes or mucous membranes, do not bandage tightly. Stop use and ask a doctor if condition worsens, if symptoms persist for more than 7 days or clear up and occur again within a few days, excessive skin irritation occurs.

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

Use only as directed. **Adults and children over 12 years**: apply to affected area not more than 3 to 4 times daily. **Children 12 years oryounger**: ask a doctor. Close cap tightly after use.

Water, lanolin, glyceryl stearate SE, stearic acid, polysorbate 80, curcuma longa (turmeric) root extract, glycerin, carbomer, triethanolamine, potassium hydroxide, disodium EDTA



DR GOLDSONS PAIN

menthol 8%, methyl salicylate 30% cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72839-145
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII: O414PZ4LPZ)	METHYL SALICYLATE	30 g in 100 g	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	8 g in 100 g	

Strength

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:72839-145- 04	113.4 g in 1 JAR; Type 0: Not a Combination Product	09/21/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/21/2021	

Labeler - Derma Care Research Labs, LLC (116817470)

Registrant - Derma Care Research Labs, LLC (116817470)

Establishment				
Name	Address	ID/FEI	Business Operations	
Derma Care Research Labs, LLC		116817470	manufacture(72839-145)	

Revised: 5/2023 Derma Care Research Labs, LLC