MUSCLE RUB ULTRA STRENGTH CVS- camphor 4.00% menthol 10.00% methyl salicylate 30.00% spray CVS

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.

Drug Facts

Purpose
Topical Analgesic
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Warning For external use only

Flammable.

- · do not use while smoking or near heat or flame
- avoid long term storage above 104oF
- do not puncture or incinerate. Contents under pressure
- do not store at temperature above 120oF

When using this product

- avoid contact with eyes and mucous membranes
- do not apply to wounds or damages skin
- do not bandage tightly or use with a heating pad
- · use only as directed

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days
- needed for longer than 1 week

Keep out of reach of the children

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

Uses

temporarily reliefs minor pain associated with • arthritis • simple backache • muscle strains • bruises • muscle sprains

Directions

- shake well
- adults and children 12 years of age and older: spray on affected area, not more than 3 to 4 times daily
- children under 12 years of age: consult a doctor

Inactive Ingredient



MUSCLE RUB ULTRA STRENGTH CVS

camphor 4.00% menthol 10.00% methyl salicylate 30.00% spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-492
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	4 g in 100 g	
Menthol (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	Menthol	10 g in 100 g	
Methyl Salicylate (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	Methyl Salicylate	30 g in 100 g	

Inactive Ingredients			
Ingredient Name	Strength		
ALCOHOL (UNII: 3K9958V90M)			

Packaging					
l	# Ite	m Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:59	779-492-04	113 g in 1 CAN; Type 0: Not a Combination Product	12/22/2014	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	12/22/2014		

Labeler - CVS (062312574)

Registrant - Product Quest Mfg (927768135)

Establishment					
Name	Address	ID/FEI	Business Operations		
Product Quest Mfg		927768135	manufacture(59779-492)		

Revised: 12/2017 CVS