

CVS SORE MUSCLE RUB VANISHING SCENT- menthol cream

CVS Pharmacy

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

Active Ingredient

Menthol 2.5%

Purpose

Topical Analgesic

Uses

Temporarily relieves minor pain associated with:

- arthritis
- simple backache
- muscle strains
- muscle sprains
- bruises
- cramps

For external use only

When using this product

- Use only as directed
- wash hand thoroughly with soap and water after use
- do not bandage tightly or use with a heating pad
- avoid contact with eyes or mucous membranes
- do not apply to wounds or damaged skin

Stop use and ask doctor if

- Condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days
- redness is present
- irritation develops

If pregnant or breast-feeding

ask a health care professional before use.

If swallowed, get medical help or contact a poison control center immediately.

Directions

Adults and children over 2 years:

- wash hand thoroughly with soap and water after use
- apply generously to affected area
- squeeze desired amount of gel onto affected area
- using the sponge-top applicator, massage dispensed gel into painful area until thoroughly absorbed

- repeat as necessary, but no more than 4 times daily

Children 2 years or younger:

- Ask a doctor

Allantoin, Aloe Barbadosensis Leaf Juice, Carbomer, DMDM Hydantoin, Glycerin, Methylparaben, Phenoxyethanol, Propylparaben, SD Alcohol 40 (15.47%), Steareth-2, steareth-21, Triethanolamine, water.

Questions or comments

1-800-635-3696



CVS SORE MUSCLE RUB VANISHING SCENT

menthol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-712
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	0.025 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ALCOHOL (UNII: 3K9958V90M)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-712-01	1 in 1 CARTON	06/02/2008	
1		70.8 g in 1 TUBE; 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	06/01/2008	

Labeler - CVS Pharmacy (062312574)

Revised: 1/2017

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