CVS PAIN RELIEF- menthol 4% gel
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.

CVS Pain Relief Colorless Gel

Menthol 4%

Topical Analgesic

Temporary relief from minor aches and pains of sore muscles and joints associated with:

arthritis

backache

strains

sprains

For external use only

Flammable: Keep away from excessive heat or open flame

Ask a doctor before use if you have: sensitive skin

Stop use and ask a doctor if

condition worsens or symptoms persist for more than 7 days or clear up and occur again within a few days.

When using this product

Use only as directed

Avoid contact with eyes or mucous membranes

Do not apply to wounds or damaged skin

Do not use with other ointments, creams, sprays or liniments

Do not apply to irritated skin or if excessive irritation develops

Do not bandage or use with heating pad or device

Store in a cool dry place away from direct sunlight

Wash hands after use with cool water

If pregnant or breastfeeding, ask a health professional before use

Keep out of reach of children: If accidentally ingested, get medical help or contact a poison Control immediately (1800-222-1222)

Directions

Adults and children 2 years of age and older: Rub a thin film over affected areas not more than daily: massage not necessary

Children under 2 years of age : Consult physician

Aloe Barbadensis Leaf Juice, Arnica Montana Flower Extract, Camellia Sinensis Leaf Extract, Camphor, Carbomer, Dimethyl Sulphone (MSM), Glycerin, Isopropyl Alcohol, Phenoxyethanol, Triethanolamine, Water

CVS PAIN RELIEF

menthol 4% gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51316-997
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	4 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
TROLAMINE (UNII: 9O3K93S3TK)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ARNICA MONTANA (UNII: O80TY208ZW)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-997-04	85 g in 1 TUBE; Type 0: Not a Combination Product	09/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/29/2020	

Labeler - CVS (062312574)

Revised: 9/2022 CVS