CVS PAIN RELIEF- menthol 10.5% 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.

CVS Pain Relief Spray

Menthol 10.5%

Topical analgesic

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

Arthritis

Backache

Strains

Sprains

For external use only

Flammable: Keep away from excessive heat or open flames. Contents Under Pressure.

Do Not Puncture Or Incinerate

Do Not Store At Temperature Above 120°F

Ask a Doctor before using if you have sensitive skin

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

Do not bandage

Do not use with heating pad or device

Store in a cool dry place away from sunlight

Wash hands after use with cool water

Stop use and ask doctor if condition worsens or symptoms persist for more than 7 days or clear up and occur again within a few days

If swallowed, get medical help or contact a Poison Control Center immediately (1-800-222-1222)

If pregnant or breast-feeding, ask a health professional before use.

Adults and children 12 years of age and older: Spray on to the affected areas not more than 4 times daily; message not necessary

Children under 12 years of age: Consult physician

Aloe Barbadensis leaf Juice, Arnica Montana Flower Extract, Camellia Sinensis Leaf Extract, Dimethyl Sulphone (MSM), Isobutane, Propylene Glycol, SD Alcohol 40 B (30%), Water

CVS PAIN RELIEF

menthol 10.5% spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51316-996
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
DEHYDRATED ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOBUTANE (UNII: BXR49TP611)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARNICA MONTANA (UNII: O80TY208ZW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-996-10	85 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/13/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/13/2022	

Labeler - CVS (062312574)

Revised: 9/2022 CVS