CVS PAIN RELIEVING- camphor 5.5% menthol 16% lotion 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 Relieving Lotion RollOn

Camphor 5.5 %

Menthol 16 %

Pain Relieving Lotion

For the temporary relief of minor aches and pains of muscles and joints associated with

Arthritis

simple backache

strains

sprains

bruises

Provides penetrating pain relief

For extranal use only

avoid contact with eyes and mucous membranes

do not bandage tightly

do not use with a heating pad, medicated patch or other of local heat

do not apply to wounds or damaged, broken or irritated skin

condition worsens or irritation develops

pain, swelling or blistering develops where product was applied

redness or severe burning develops when product was applied

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

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Adults and children 12 years of age and older:

Apply to affected area not more than 3 to 4 times daily

Wash hands with soap and water after use if product comes with hands

Children under 12 years of age

Ask a doctor

store between 20° - 25°C (68° - 77°F)

Carbomer, Clove Oil, Disodium EDTA, Eucalyptus Oil, Pentylene Glycol, Peppermint Oil, Phenoxyethanol, Polysorbate 80, SD Alcohol 40 B, Sorbitan Oleate, Tocopherol Acetate, Water



camphor 5.5% menthol 16% lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:51316-885

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	16 mg in 1 g	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	5.5 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLYSORBATE 80 (UNII: 6OZP39ZG8H)		
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
SORBITAN MONOOLEATE (UNII: 06XEA2VD56)		
PENTYLENE GLYCOL (UNII: 50C1307PZG)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
PEPPERMINT OIL (UNII: AV092KU4JH)		
CLOVE OIL (UNII: 578389D6D0)		
ALCOHOL (UNII: 3K9958V90M)		
EUCALYPTUS OIL (UNII: 2R040NI662)		

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:51316-885- 16	72 g in 1 TUBE; Type 0: Not a Combination Product	12/19/2022	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	12/19/2022		

Revised: 12/2022 CVS