HUMCO CAMPHOR SPIRIT- camphor 10% liquid Humco Holding Group, Inc.

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.

HUMCO Camphor Spirit

Drug Facts

Active Ingredient

Camphor 10%

Purpose

Counterirritant

Use

For the temporary relief of minor aches, muscle and joint pain associated with arthritis, strains, brusies and simple backache.

Warnings

For external use only.

Do not drink. If swallowed, immediately give 3 or 4 glases of water. Do not induce vomiting. If vomiting occurs, give fluids again. Do not give anything by mouth to an unconscious or convulsing person. Get medical attention immediately.

Avoid contact with eyes or mucous membranes.

Do not apply to irritated skin.

When using this product

Do not bandage tightly.

Discontinue use and consult a doctor if

condition worsens, or if excesive irritation develops.

symptoms persit for more than 7 days, or clear up and occur again within a few days.

Keep out of reach of children

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center right away. In case of eye contact, flush eyes with running water for 15 minutes, get medical attention.

Directions

(dilute 3 parts olive oil and mix well)

Adults and children 2 yrs. and older. apply to the affected area not more than 3 or 4 times daily. Children under 2 yrs of age: consult a doctor befre use.

Other Information

Flammable: Keep away from spark, heat and flame.

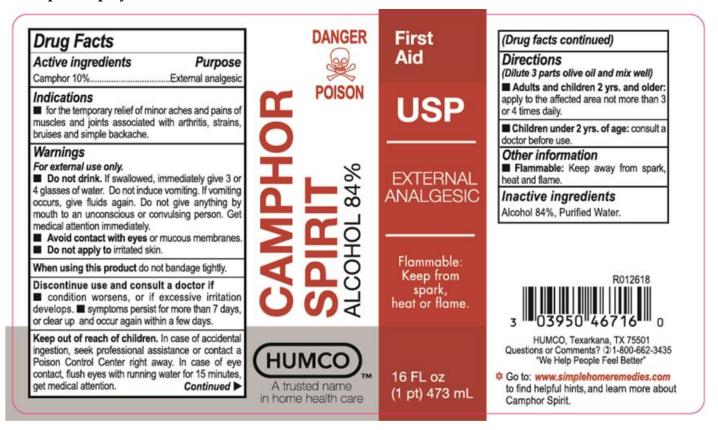
Inactive ingredients

Alcohol 84%, Purified Water

Questions or Comments?

1-800-662-3435

Principal Display Panel



HUMCO CAMPHOR SPIRIT

camphor 10% liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:0395-0467 Route of Administration TOPICAL Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (NATURAL) (UNII: N20 HL7Q941) (CAMPHOR (NATURAL) - UNII:N20 HL7Q941)	CAMPHOR (NATURAL)	100 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

l	Packaging						
	# Item Code	Package Description	Marketing Start Date	Marketing End Date			
	1 NDC:0395-0467-	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 1/0 1/20 13				
	2 NDC:0395-0467- 92	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 1/0 1/20 13				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part348	0 1/0 1/20 0 8			

Labeler - Humco Holding Group, Inc. (825672884)

Establishment			
Name	Address		Business Operations
Humco Holding Group, Inc.		825672884	manufacture(0395-0467), analysis(0395-0467), pack(0395-0467), label(0395-0467)

Revised: 2/2020 Humco Holding Group, Inc.