CAMPHOR SPIRIT- camphor 10% liquid Pharma Nobis, LLC

Private Label Camphor Spirit USP

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 thaaffected 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

Sunmark Label

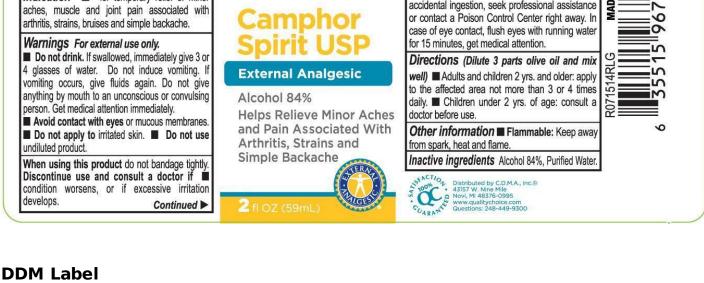


Leader Label



Quality Choice Label





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**Continued ▶*



CAMPHOR SPIRIT USP ALCOHOL 84%

COUNTERIRRITANT

DANGER POISON

DISTRIBUTED BY: **DRUG MART-FOOD FAIR** 211 Commererce Dr. MEDINA, OH 44256

www.discount-drugmart.com

2 FL OZ (59mL)

(Drug facts continued)

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B053013.RLG



CAMPHOR SPIRIT

camphor 10% liquid

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:82645-920

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NII: N20HL7Q941) (CAMPHOR (NATURAL) -	CAMPHOR (NATURAL)	100 mg

CAMPHOR (NATURAL) (UN

UNII:N20HL7Q941)

' in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82645- 920-92	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/12/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	01/01/2008		

Labeler - Pharma Nobis, LLC (118564114)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	$manufacture (82645-920) \ , \ analysis (82645-920) \ , \ pack (82645-920) \ , \\ label (82645-920)$

Revised: 12/2023 Pharma Nobis, LLC