QUALITY CHOICE CAMPHOR SPIRIT- camphor 10% liquid
Chain Drug Market Association

Quality Choice Camphor Spirit

Drug Facts

Active Ingredient

Camphor 10%

Purpose

External Anagesic

Indication

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.

Do not use undiluted product.

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 th affected area not more than 3 or 4 times daily.

Children under 2 yrs of age: consult a doctor befre use.

Other Information

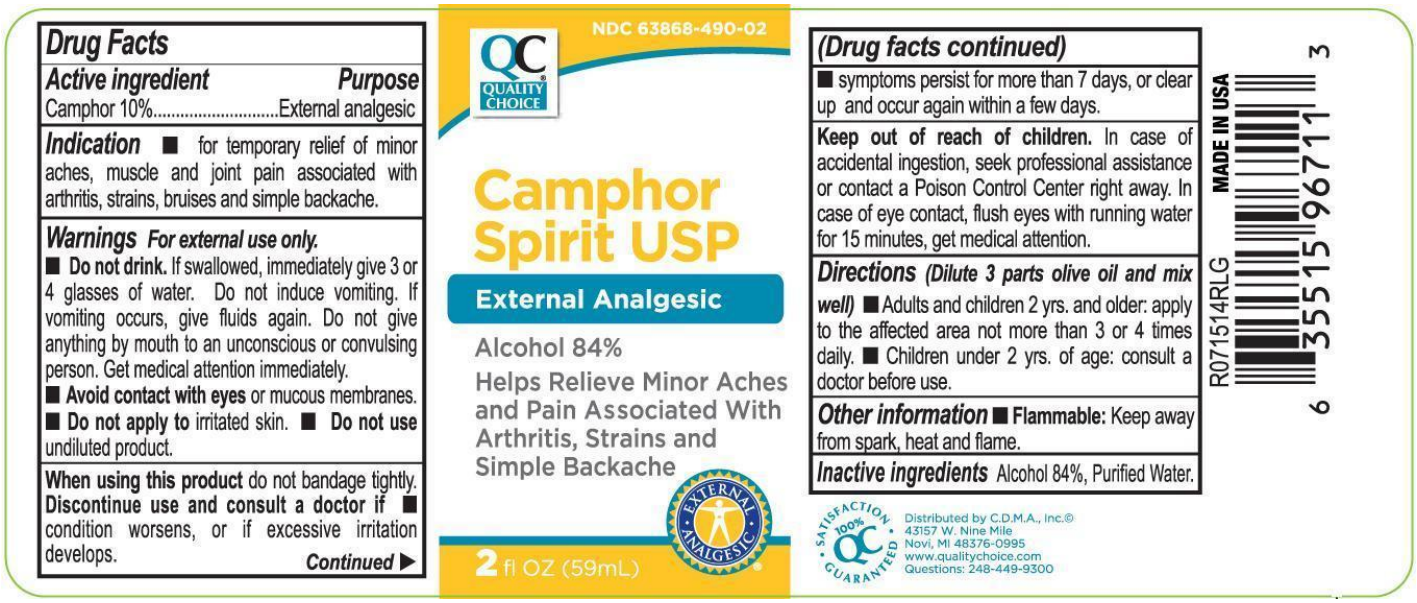
Flammable: Keep away from spark, heat and flame.

Inactive ingredients

Alcohol 84%, Purified Water

Principal Display Panel

Camphor Spirits, USP



QUALITY CHOICE CAMPHOR SPIRIT			
camphor 10% liquid			
Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-490
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	100 mg in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-490-02	473 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 - Chain Drug Market Association (011920774)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(63868-490) , analysis(63868-490) , pack(63868-490) , label(63868-490)

Revised: 12/2023 Chain Drug Market Association