

TITLE

Combination composition, comprising as active ingredients L-carnitine or propionyl L-carnitine, for the prevention or treatment of chronic venous insufficiency.

5 FIELD OF THE INVENTION

The present invention relates to a combination composition comprising as active ingredients L-carnitine or a derivative thereof, or a salt thereof, troxerutine, diosmine and hesperidine, useful for the prevention and/or treatment of chronic venous insufficiency (CVI).

10 BACKGROUND OF THE INVENTION

In humans, arteries bring oxygen-rich blood from the heart to the rest of the body and veins return oxygen-poor blood back to the heart. When human leg veins cannot pump enough blood back to heart, there is the onset of chronic venous insufficiency (CVI). CVI is also sometimes
15 called chronic venous disease, or CVD.

Humans have three kinds of veins: superficial veins, which lie close to the skin, deep veins, which lie in groups of muscles, and perforating veins, which connect the superficial to the deep veins. Deep veins lead to the vena cava, human body's largest vein, which runs directly to the
20 heart.

When humans are in the upright position, the blood in their leg veins must go against gravity to return to our heart. To accomplish this, leg muscles squeeze the deep veins of their legs and feet to help move blood back to our heart. One-way flaps, called valves, in the veins keep
25 blood flowing in the right direction. When the leg muscles relax, the

valves inside the veins close. This prevents blood from flowing in reverse, back down the legs. The entire process of sending blood back to the heart is called the venous pump.

When humans walk and the leg muscles squeeze, the venous pump works well. But when patient sit or stand, especially for a long time, the blood in the leg veins can pool and increase the venous blood pressure. Deep veins and perforating veins are usually able to withstand short periods of increased pressures. However, sitting or standing for a long time can stretch vein walls because they are flexible. Over time, in susceptible individuals, this can weaken the walls of the veins and damage the vein valves, causing CVI.

During CVI ankles may swell and calves may feel tight. The legs may also feel heavy, tired, restless, or achy. Patient may feel pain while walking or shortly after stopping.

CVI may be associated with varicose veins. Varicose veins are swollen veins that we can see through the skin. They often look blue, bulging, and twisted. Large varicose veins can lead to skin changes like rashes, redness, and sores.

CVI can also cause problems with leg swelling because of the pressure of the blood pooling in the veins. The lymphatic system may also produce fluid, called lymph, to compensate for CVI. Human leg tissues may then absorb some of this fluid, which can increase the tendency for the legs to swell. In severe cases, CVI and the leg swelling can cause ulcers to form on the lower parts of the leg.

Over the long-term, blood pressure that is higher than normal inside the leg veins causes CVI. Other causes of CVI include deep vein thrombosis (DVT) and phlebitis, both of which cause elevated pressure in our veins by obstructing the free flow of blood through the veins.

5 DVT occurs when a blood clot (properly called a thrombus) blocks blood from flowing toward the heart, out of a deep or perforating vein. The blood trying to pass through the blocked veins can increase the blood pressure in the vein, which, in turn, overloads our valves. Vein valves that do not work properly are called incompetent because they
10 stretch and no longer work efficiently, and incompetent valves contribute to CVI. DVT is a potentially serious condition that causes leg swelling and requires immediate medical attention because sometimes the blood clots in the veins can break off and travel to the lungs. This condition is called a pulmonary embolus.

15 Phlebitis occurs when a superficial or deep vein becomes swollen and inflamed. This inflammation causes a blood clot to form, which can also lead to DVT.

Factors that can increase the risk for CVI include a family history of varicose veins, being overweight, being pregnant, not exercising
20 enough, smoking, and standing or sitting for long periods of time. Although CVI can affect anyone, age and sex can also be factors that may increase the tendency to develop CVI; women older than 50 most often get CVI.

For mild cases of CVI, physician may recommend compression
25 stockings. Compression stockings are elastic stockings that squeeze the

veins and stop excess blood from flowing backward. In this way, compression stockings can often also help heal skin sores and prevent them from returning. Patient may need to wear compression stockings daily for the rest of the life.

5 More serious cases of CVI may be treated with injections, called sclerotherapy, or with surgical procedures. Fewer than 10 percent of patient with CVI require surgery to correct the problem. Surgical treatments include ablation, vein stripping, bypass surgery, valve repair, and angioplasty or stenting of a vein.

10 The flavonoids, troxerutine, diosmine and hesperidine, also referred to as rutins oxerutins or rutosides, are well known for their antioxidant properties; in addition they also display other properties that impact upon the circulatory system i.e. blood and microvascular endothelial cells (G.I.O.T. 2009;35:23-33).

15 Many epidemiological studies suggest that these flavonoids are associated with improvement of circulation and a general reduced risk of cardiovascular diseases. The mechanisms underlying these protective actions include antithrombotic, anti-ischemic, anti-oxidant, and vasorelaxant properties. In particular these flavonoids have actions that
20 improve circulation by vasodilatation, decreasing platelets clotting and preventing low-density lipoproteins (LDLs) from oxidizing (Journal of Agricultural and Food Chemistry 2008 56 (15), pp 6185-6205).

 These flavonoids also demonstrate strong anti-inflammatory properties that are thought to be the result of the inhibition of the
25 synthesis and biological activities of different pro-inflammatory

mediators, mainly the arachidonic acid derivatives, prostaglandins E₂, F₂, and thromboxane A₂. These processes improve venous tone and lymphatic drainage, and reduce capillary hyperpermeability by protecting the microcirculation from inflammatory processes.

5 This effect has been shown also in clinical studies. The study of Belcaro et al (Angiology, 59; 5S) showed that these flavonoids were effective in the control and treatment of oedema and increased capillary filtration in venous hypertension and diabetic microangiopathy.

10 Another study by Belcaro et al. (Angiology, March 1, 2008; 59 (1) suppl. 7S-13S) also looked at the use of these flavonoids in the treatment of chronic venous disease and signs and symptoms of chronic venous insufficiency (CVI), varicose veins, and deep venous disease.

15 In Rev Fr Gynecol Obstet. 1991 Feb 25; 86 (2 Pt 2) : 209-12, is reported that troxerutine in women (half in the context of premenstrual syndrome and half in pregnant) was useful for the treatment of vulval varicosities and venous insufficiency of the lower limbs.

20 In Br. J. Surg. 2000, 87; 868-872, it is reported that diosmine has anti-inflammatory actions that play a role in protecting the blood vessels and contributes to the maintenance of good circulation of the blood and maintains the venous tone.

 In Farmaco, 40 (11); 709-712, it is reported that hesperidine has an antioxidant action and together with troxerutine and diosmine promotes healthy vessel functions.

In US 4255449 it is reported that L-carnitine is useful for increasing the HDL cholesterol and for treating diseases linked to high cholesterol level.

In WO04091602 9 it is reported that L-carnitine is useful the
5 treatment of cardiovascular diseases.

In US 5811457 it is reported that propionyl L-carnitine is useful the treatment of chronic arteriosclerosis obliterans

In WO 2007045639 it is reported that propionyl L-carnitine is useful the treatment of left ventricular hypertrophy in dialysed patients.

10 In US 4343816 it is reported that propionyl L-carnitine is useful the treatment of peripheral vascular diseases.

Furthermore, while there are other publications available in which is shown that the compounds of the invention are useful for the treatment of diseases of the veins, none of them mention nor suggest
15 about the unexpected synergistic effect shown by the composition of the invention.

DESCRIPTION OF THE INVENTION

It has now been found that a combination composition comprising as active ingredients L-carnitine or a derivative thereof (e.g. propionyl L-carnitine) or a salt thereof, troxerutine, diosmine and hesperidine, is
20 endowed with a surprisingly synergistic effect for the prevention and/or treatment of diseases of the veins selected from Chronic venous insufficiency (CVI) and chronic venous disease (CVD) or their complications.

An example of complications or diseases of the veins (due to CVI or CVD) are reported in the following: swelling and inflammation of veins in the rectum, anus and vulva; venous hypertension; increased permeability; oedema; capillary damage; skin changes; venous leg
5 ulcers; swelling ankles; heavy legs; varicose veins; swelling leg; ulcers; vein thrombosis; phlebitis; thrombo-phlebitis; pulmonary embolus; hemorrhoids.

It is therefore one object of the present invention a combination composition comprising as active ingredients L-carnitine or a salt
10 thereof, troxerutine, diosmine and hesperidine.

It is a further object of the present invention a combination composition comprising as active ingredients propionyl L-carnitine or a salt thereof, troxerutine, diosmine and hesperidine.

The compositions mentioned above may further comprise other
15 active ingredients useful for treating diseases of the vein.

It is a further object of the present invention a composition comprising:

(a) L-carnitine or propionyl L-carnitine, in a dose of from 10 to 3000 mg, preferred doses is of from 50 mg to 400 mg, the most preferred
20 dose is 136 mg.

(b) troxerutine, in a dose of from 900 mg to 50 mg, preferred doses is of from 400 mg to 200 mg, the most preferred dose is 300 mg;

(c) diosmine, in a dose of from 900 mg to 50 mg, preferred doses is of from 400 mg to 200 mg, the most preferred dose is 300 mg; and

(d) hesperidine in a dose of from 10 mg to 500 mg, preferred doses is of from 50 mg to 200 mg, the most preferred dose is 100 mg.

It is a further object of the present invention the composition mentioned above, for use as anti-chronic venous insufficiency and
5 complications thereof.

It is a further object of the present invention the composition mentioned above, for use as anti-chronic venous disease and complications thereof.

It is a further object of the present invention the use of the
10 composition mentioned above, for preparing a medicament for the prevention or treatment of chronic venous insufficiency, chronic venous disease and complications thereof, in which said complications are selected from the group comprising: swelling and inflammation of veins in the rectum, anus and vulva; venous hypertension; increased
15 permeability; oedema; capillary damage; skin changes; venous leg ulcers; swelling ankles; heavy legs; varicose veins; swelling leg; ulcers; vein thrombosis; phlebitis; thrombo-phlebitis; pulmonary embolus or hemorrhoids.

It is a further object of the present invention the use of the
20 composition mentioned above, for preparing a dietary supplement for the prevention or treatment of chronic venous insufficiency, chronic venous disease and complications thereof.

The composition of the invention may further comprise co-enzymes, mineral substances, antioxidants, vitamins, anticlotting agents
25 and agents useful for treating diseases of the veins.

What is meant by salt of L-carnitine is any salt of the latter with an acid that does not give rise to toxic or side effects.

Non-limiting examples of such salts are: chloride, bromide, orotate, aspartate, acid aspartate, acid citrate, magnesium citrate, phosphate, acid phosphate, fumarate and acid fumarate, magnesium fumarate, lactate, maleate and acid maleate, oxalate, acid oxalate, pantoate, acid pantoate, sulphate, acid sulphate, glucose phosphate, tartrate and acid tartrate, glycerophosphate, mucate, magnesium tartrate, 2-amino-ethanesulphonate, magnesium 2-amino-ethanesulphonate, methanesulphonate, choline tartrate, trichloroacetate, and trifluoroacetate.

A list of FDA-approved pharmaceutically acceptable salts is given in the publication Int. J. of Pharm. 33 (1986), 201-217.

L-carnitine, propionyl L-carnitine, troxerutine, diosmine and hesperidine according to the present invention can be administrated in a "co-ordinated manner". What is meant by "co-ordinated manner" of the aforesaid compounds is, indifferently, either the co-administration, i.e. the substantially concomitant or sequential supplementation of L-carnitine or propionyl L-carnitine and at least one troxerutine, diosmine, hesperidine, or the administration of a composition comprising the aforesaid active ingredients in combination and in a mixture optionally further comprising one or more excipients or diluents pharmaceutically acceptable.

The composition of the present invention can be administered orally, parenterally, intravenously, topically and/or transdermally, in any suitable form. The oral administration is preferred.

An example of form of administration is in a liquid, semi-liquid or
5 solid form in sachets, pills, vials, ointment, gel or liposome.

L-carnitine and propionyl L-carnitine are known compounds and their preparation process is described in US 4,254,053.

Troxerutine, diosmine and hesperidine are widely sold on the market and their CAS-No. are: 7085-55-4; 520-27-4 and 520-26-3
10 respectively.

The pharmaceutical composition according to the present invention is composed of active ingredients which are familiar to operators in the medical field and already in use.

Their procurement therefore is very easy, inasmuch as these are
15 products which have been on the market now for a long time and are of a grade suitable for human administration.

For any compound, the therapeutically effective dose can be estimated initially either in cell culture assays or in animal models, usually mice or rats.

20 The animal model may also be used to determine the appropriate concentration range and route of administration. Such information can then be used to determine useful doses and routes for administration in humans.

The precise effective dose for a human subject will depend upon the severity of the disease state, general health of the subject, age, weight, and gender of the subject, diet, time and frequency of administration, drug combination(s), reaction sensitivities, and tolerance/response to therapy. This amount can be determined by routine experimentation and is within the judgement of the clinician.

The following non limiting examples further illustrate the invention.

EXAMPLE 1

10 Reduction of the swelling of the recto-anus, in croton oil-induced hemorrhoid model in rats.

Male SD rats (6 weeks old, approximately 140 g) were purchased from Harlan Sprague Dawley and allowed to acclimate for 1 week. The rats were maintained in a pathogen-free facility in accordance with the National Research Council of Laboratory Animal care and use guidelines. Each experiment was performed with age-matched rats 7-8 weeks old. The croton oil-induced hemorrhoid model in rats was performed according to the method published by Nishiki (Nishiki et al. (1988) Folia Pharmacology Japan 92:215-225; Nishiki et al. (1988) Folia Pharmacol. Japan 92:227-240). Briefly, a cotton swab with a diameter of 4 mm soaked with 0.16 mL of inducer (deionized water: pyridine: ethyl ether: 6% croton oil/ethyl ether (1:4:5:10) was applied to the rat's anus for 12 seconds. The final concentration of croton oil was 3%. The edema

developed linearly until 7-8 hours after application and the severity of the edema was sustained for more than 24 hours. Twenty-four hours later, recto-anus tissue (approx. 10 mm long) was isolated after the rats were euthanized. The weights of rat body and recto-anus were measured. The recto-anus coefficient (RAC) was calculated using the formula: weight of recto-anus (mg)/body weight (g).

The compounds of the invention were administered orally suspended in saline (1 mL) once per day for five days. Last treatment with the compounds of the invention was made the day of induction of oedema with croton oil.

The compounds of the invention were administered alone or in combination at the following doses: 20 mg/kg propionyl L-carnitine or L-carnitine; 30 mg/kg troxerutine, diosmine or hesperidine.

Twenty-four hours after hemorrhoid induction with croton oil the RAC of rats was determined.

The results obtained are reported in the following Table 1.

TABLE 1

n=5 rats per group; degrees of freedom= 8.

		RAC	SE	P<
1	Vehicle alone	1.28	0.14	-
2	No treatment	0.97	0.11	- NS vs 1
3	L-carnitine + Troxerutine + diosmine + hesperidine. 20 mg/kg + 30 mg/kg + 30 mg/kg + 30 mg/kg.	1.33	0.12	- NS vs 1
4	Propionyl L-carnitine + Troxerutine + diosmine + hesperidine. 20 mg/kg + 30 mg/kg + 30 mg/kg + 30 mg/kg.	1.31	0.11	- NS vs 1
5	crotonoil (3%)	253	0.18	0.001 vs 1
6	L-carnitine + Troxerutine + diosmine + hesperidine + croton oil.	1.34	0.09	- NS vs 1 - NS vs 3 - NS vs 4 - 0.001 vs 5 - 0.001 vs 8 - 0.001 vs 9 - 0.001 vs 10 - 0.001 vs 11 - 0.001 vs 12 - 0.05 vs 21
7	Propionyl L-carnitine + Troxerutine + diosmine + hesperidine + croton oil.	1.32	0.11	- NS vs 1 - NS vs 3 - NS vs 4 - 0.001 vs 5 - 0.001 vs 8 - 0.001 vs 9 - 0.001 vs 10 - 0.001 vs 11 - 0.001 vs 12 - 0.05 vs 21
8	Troxerutine + croton oil	2.11	0.19	0.05 vs 1
9	Diosmine + croton oil	2.02	0.18	0.05 vs 1
10	Hesperidine + croton oil	2.07	0.20	0.05 vs 1
11	L-carnitine + croton oil	1.99	0.18	0.05 vs 1
12	Propionyl L-carnitine + croton oil	1.96	0.16	0.05 vs 1
13	Troxerutine + diosmine + croton oil	2.00	0.20	0.05 vs 1
14	Troxerutine + hesperidine + croton oil	1.99	0.18	0.05 vs 1
15	troxerutine + L-carnitine + croton oil	1.91	0.17	0.05 vs 1
16	Troxerutine + propionyl l-carnitine + croton oil	1.89	0.17	0.05 vs 1
17	Diosmine + hesperidine + croton oil	2.01	0.21	0.05 vs 1
18	Diosmine + L-carnitine + croton oil	1.87	0.15	0.05 vs 1
19	Diosmine + propionyl L-carnitine	1.83	0.13	0.05 vs 1
20	Hesperidine + L-carnitine + croton oil	1.84	0.14	0.05 vs 1
20	Hesperidine + propionyl L-carnitine + croton oil	1.84	0.13	0.05 vs 1
21	Troxerutine + hesperidine + diosmine + croton oil	1.65	0.14	NS vs 1 - 0.05 vs 5

The results reported in Table 1 show that using the compositions of the invention results statistically significant more active (respect to the use of single component) were obtained.

In fact, the results reported in Table 1 show that the compositions of the invention completely inhibited the swelling of the recto-anus induced by croton oil.

EXAMPLE 2

Relaxant effect on aortic rings of the compounds tested.

The objective of this study was to investigate the vasoarelation effects of the compositions of the invention using isolated aorta from spontaneously hypertensive rats.

Materials and Methods

The compounds tested were used at a concentration expressed as final concentration in the organ chamber.

Animals

Male spontaneously hypertensive rats (SHR) 10-12 weeks old, weighing 250-300 g, were housed at 24 ± 2 °C with $60 \pm 20\%$ relative humidity, on a 12-h light-dark cycle. Rats were given free access to a diet of standard food and water. All experiments were performed according to the guidelines for the ethical treatment of animals of the European Union. The rats were killed by cervical dislocation and the aortae were rapidly dissected.

Aortic ring preparation

The descending thoracic aorta was placed in a modified Krebs-Henseleit solution (PSS) containing (mM): NaCl 118, KCl 4.75, NaHCO₃ 25, MgSO₄ 1.2, CaCl₂ 1.8, KH₂PO₄ 1.2 and glucose 11.

After excess fat and connective tissue were removed, the aortae were cut into 2-3-mm rings. Aortic rings were mounted under a basal tension of 2 g in 20 mL organ baths containing PSS and attached to an isometric transducer (Harvard UF-1); the signal was recorded by a Powerlab data acquisition system (AD-Instruments). The tissue bath was maintained at 37°C and bubbled with a 95% O₂ and 5% CO₂ gas mixture.

Relaxant effect of the compounds tested.

The relaxant effect of compounds tested was assessed by adding the compound to aortic rings precontracted by phenylephrine (1 micromolar).

After the aortae were contracted with phenylephrine and when the contractile response was reached, the test compound was added to the bath. All the results were expressed as a percentage of the maximal contraction of phenylephrine-induced responses.

Statistical analysis

Results are expressed as percentages from the initial precontraction level.

The significance was calculated on the actual contraction values means $\pm$ SE (n = 8 aorta preparations per group). Eight preparations

were studied from a single aorta. The Student t-test was used as the comparison test for statistical analysis.

$p < 0.05$ Values were considered to represent a significant difference.

5 The results obtained are reported in the following Table 2.

TABLE 2

	TREATMENT	% RELAXATION	P<
1	Saline	0	-
2	L-carnitine + troxerutine + hesperidine + diosmine (100 micro-molar + 100 micro- molar + 100 micro-molar + 100 micro-molar).	52	- 0.001 vs 1 - 0.01 vs 4 - 0.01 vs 5 - 0.01 vs 6 - 0.05 vs 7 - 0.05 vs 8 - 0.05 vs 18
3	Propionyl L-carnitine + troxerutine + hesperidine + diosmine. (100 micro-molar + 100 micro- molar + 100 micro-molar + 100 micro-molar).	53	- 0.001 vs 1 - 0.01 vs 4 - 0.01 vs 5 - 0.01 vs 6 - 0.05 vs 7 - 0.05 vs 8 - 0.05 vs 18
4	Troxerutine (100 micro-molar).	10	0.05 vs 1
5	Diosmine (100 micro-molar).	10	0.05 vs 1
6	Hesperidine (100 micro-molar).	9	0.05 vs 1
7	L-carnitine (100 micro-molar).	25	0.01 vs 1
8	Propionyl L-carnitine (100 micro-molar).	28	0.01 vs 1
9	Troxerutine + diosmine	12	0.05 vs 1
10	Troxerutine + hesperidine	13	0.05 vs 1
11	Troxerutine + L-carnitine	27	0.01 vs 1
12	Troxerutine + propionyl L- carnitine	29	0.01 vs 1
13	Diosmine + hesperidine	14	0.05 vs 1
14	Diosmine + L-carnitine	26	0.01 vs 1
15	Diosmine + propionyl L- carnitine	28	0.01 vs 1
16	Hesperidine + L-carnitine	26	0.01 vs 1
17	Hesperidine + propionyl L- carnitine	29	0.01 vs 1
18	Troxerutine + hesperidine + diosmine	35	0.001 vs 1

The results reported in Table 2 show that the addition of compositions of the invention to aortic rings precontracted with phenylephrine elicited endothelium-dependent relaxation which was statistically significant more active respect to the single components.

5 In the following are reported some non limiting examples of the compositions of the invention.

Composition 1

	- L-Carnitine tartrate	200 mg
	Equivalent to L-carnitine	136 mg
10	- Troxerutine NEC®	700 mg
	Comprising Troxerutina	300 mg
	- Diosmine	300 mg
	- Hesperidine 98%	100 mg
	Equivalent to Hesperidine	98 mg.

15 **Composition 2**

	- Propionyl L-Carnitine tartrate	250 mg
	Equivalent to propionyl L-carnitine	136 mg
	- Troxerutine NEC®	700 mg
	Comprising Troxerutina	300 mg
20	- Diosmine	300 mg
	- Hesperidine 98%	100 mg
	Equivalent to Hesperidine	98 mg.

Composition 3

	- L-Carnitine fumarate	200 mg
	Equivalent to L-carnitine	136 mg
	- Troxerutine NEC®	700 mg
5	Comprising Troxerutina	300 mg
	- Diosmine	300 mg
	- Hesperidine 98%	100 mg
	Equivalent to Hesperidine	98 mg.

Composition 4

10	- Propionyl L-Carnitine fumarate	250 mg
	Equivalent to propionyl L-carnitine	136 mg
	- Troxerutine NEC®	700 mg
	Comprising Troxerutina	300 mg
	- Diosmine	300 mg
15	- Hesperidine 98%	100 mg
	Equivalent to Hesperidine	98 mg.

CLAIMS

1. Combination composition comprising as active ingredients L-carnitine or propionyl L-carnitine or a salt thereof, troxerutine,
5 diosmine, hesperidine, and optionally one or more excipient pharmaceutically acceptable.
2. The combination composition of claim 1, which comprises:
 - L-carnitine or propionyl L-carnitine, in a dose of from 10 to 3000 mg, preferred doses is of from 50 mg to 400 mg, the most
10 preferred dose is 136 mg;
 - troxerutine, in a dose of from 900 mg to 50 mg, preferred doses is of from 400 mg to 200 mg, the most preferred dose is 300 mg;
 - diosmine, in a dose of from 900 mg to 50 mg, preferred doses is of from 400 mg to 200 mg, the most preferred dose is 300 mg; and
15 - hesperidine in a dose of from 10 mg to 500 mg, preferred doses is of from 50 mg to 200 mg, the most preferred dose is 100 mg.
3. The combination composition of claim 1, as a dietary supplement.
4. The combination composition of claim 1, as a medicament.
5. The combination composition of claim 1, further comprising
20 coenzymes, mineral substances, antioxidants, vitamins, anticlotting agents, and agents useful for treating diseases of the veins.
6. The combination composition of claim 1 or 2, for use in the prevention or treatment of diseases of the veins.
7. Use of a combination composition comprising as active ingredients
25 L-carnitine or propionyl L-carnitine or a salt thereof, troxerutine, diosmine and hesperidine, for preparing a medicament for the prevention or treatment of diseases of the vein.
8. The use according to claim 7, in which the disease of the veins is selected from the group consisting of: chronic venous
30 insufficiency, chronic venous disease and complication thereof.

9. The use according to claim 8, in which said complication is selected from the group consisting of: swelling and inflammation of veins in the rectum, anus and vulva; venous hypertension; increased permeability; oedema; capillary damage; skin changes; venous leg ulcers; swelling ankles; heavy legs; varicose veins; swelling leg; ulcers; vein thrombosis; phlebitis; thrombo-phlebitis; pulmonary embolus or hemorrhoids.
10. The use according to claim 7, in which the salt of L-carnitine or propionyl L-carnitine is selected from the group consisting of: chloride, bromide, orotate, aspartate, acid aspartate, acid citrate, magnesium citrate, phosphate, acid phosphate, fumarate and acid fumarate, magnesium fumarate, lactate, maleate and acid maleate, oxalate, acid oxalate, pamoate, acid pamoate, sulphate, acid sulphate, glucose phosphate, tartrate and acid tartrate, glycerophosphate, mucate, magnesium tartrate, 2-amino-ethanesulphonate, magnesium 2-amino-ethanesulphonate, methanesulphonate, choline tartrate, trichloroacetate, or trifluoroacetate.
11. The use according to claim 7, for oral, parenteral, intravenous, topical and/or transdermal administration.
12. The use according to claim 7, for oral administration.