

COMBINATION THERAPY FOR TREATMENT OF PATIENTS WITH NEUROLOGICAL DISORDERS AND CEREBRAL INFARCTION

FIELD OF THE INVENTION

5 The present invention provides methods and compositions for the treatment of stroke and neurological disorders. The methods and compositions of the present invention bring together aspects of Traditional Chinese Medicine (TCM) and Western medicine.

BACKGROUND

10 Stroke is a major cause of death and disability. Primary stroke prevention focuses on lifestyle modifications of risk factors while secondary stroke prevention aims to reduce the overall risk of recurrence in persons who have had a stroke.

There is currently a lack of treatment methods for stroke recovery in Western medicine, these
15 being frequently limited to the following options:

- a) Intra-arterial thrombolysis with intravenous tissue plasminogen activator (rt-PA), which is applicable only to 3 to 5% of stroke patients (as it has to be applied only for acute ischemic forms, only if patients do not present any contra-indication, and only within 3 to 6 hours after the onset of symptoms), can restore blood perfusion and prevent neurological and functional damage to some degree;
- 20 b) administration of aspirin / other antiplatelets / or sometimes anticoagulant to most of the cerebral stroke patients for secondary stroke prevention, which gives an improvement effect of about 1% (The International Stroke Trial (IST) a randomised trial of aspirin, subcutaneous heparin, both or neither among 19435 patients with acute ischaemic stroke. International Stroke Trial Collaborative Group. Lancet. 1997 May 31; 349(9065):1569-81);
- 25 c) medications such as analgesics may be needed to control associated symptoms
- d) rehabilitation includes physical therapy such as physiotherapy, massage, speech therapy, or occupational therapy.

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One known TCM product is NeuroAid®. Clinical studies performed in China on compositions the same as NeuroAid® have shown that this natural product combination increases stroke patients' neurological disability recovery and functional outcomes with extremely few side effects or other adverse effects. The composition of NeuroAid® has been approved by and
35 registered with the State Food and Drug Administration (SFDA) and is administered for the treatment of cerebral infarct patients during their recovery at an early or late stage, and the indications in Traditional Chinese Medicine are: to supplement qi and activate blood circulation. It is applied to treat those patients who are suffering from ischemic or hemorrhagic (for the latter, it is at present indicated in late phase only) cerebral infarction of qi deficiency

and blood stasis with manifestations of hemiplegia, hemianesthesia, wry mouth, aphasia (inarticulateness) and etc during their channel and collateral convalescent period of ischemia apoplexy, and its efficacy is supported by clinical trial data.

- 5 It may be useful for treating other types of stroke than cerebro-vascular stroke such as cardio-vascular disease (heart stroke mainly due to coronary artery stroke) as well as other neurological disorders. Neurological disorders are disorders that affect the central nervous system, the peripheral nervous system and the autonomic nervous system such as neurodegenerative diseases (for example, Alzheimer's disease and Parkinson's disease),
10 epilepsy, seizure, demyelinating diseases (for example, multiple sclerosis), cerebral palsy, traumatic injuries to or tumours in the brain, spinal cord and peripheral nerves.

NeuroAid® is capable of use as a Western medicine or a dietary supplement to provide nutrition to healthy individuals as well as patients afflicted with stroke or neurological disorders.

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NeuroAid® is typically administered orally (per os) as such or by diluting the capsules in water or via a gastric tube, 3 times each day and 4 capsules each time for a 4-week course of treatment. The duration of treatment is typically 3 months/3 courses, adaptable with regard to the patient's condition.

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- The use of TCM is, however, particularly challenging for European clinicians because of the lack of guidelines, clinical data and the small number of studies conducted under Western guidelines. Potential interactions between TCM and Western medicine that may lead to adverse side effects are also a major concern among both practitioners of TCM and Western
25 medicine. Of particular concern are the increase or decrease in the effects of a blood thinner such as Warfarin that may lead to either a bleeding episode or formation of a blood clot, and the decrease in the effect of a blood pressure medication that may lead to high blood pressure and a stroke. For example, the potential interaction of the Chinese herb salvia with the Western drug Warfarin leading to excessive blood thinning with bleeding has been well
30 documented with confirmatory laboratory studies. Non-steroidal anti-inflammatory drugs (NSAIDs), in particular aspirin, also have the potential to interact with Chinese herbs and increase bleeding risks. For these, and other reasons, TCM is generally not used in conjunction with Western medicines.

- 35 As current treatment options do not address the needs of difficult-to-treat patients with important stroke disabilities, such as hemiparalysis or aphasia, the present invention seeks to combine TCM with established agents for the treatment of stroke patients to present a new therapeutic treatment option for stroke patients.

SUMMARY

According to a first aspect, there is provided a composition comprising:

(A) at least one of the following components: DanGui, Honghua, Taoren, and Deer Horn; and

5 (B) at least one of the following components: Radix Astragali root (Membranous Milkvetch root or Huang Qi), Radix et Rhizoma Salviae Miltiorrhizae root (Red Sage root or Dan Shen), Radix Paeoniae Rubra root (Red Peony root or Chi Shao), rhizome of Ligusticum Chuanxiong (Chuan Xiong), Radix et Rhizoma Notoginseng (Sanqi), Cortex moutan (Peony or Mudanpi), Wood of Odoriferous Rosewood (Jiang Xiang), dried body of Scorpion (Quan
10 Xie), Radix Polygalae root (Yuan Zhi), Grassleaf sweetflag rhizome (Shi Changpu), Leeches (Hirudo or Shuizhi), Ground Beetle (or Tu Bie Chong), Natural or Artificial Cow-bezoar (calculus Bovis artifactus or Rengong Niuhuang), Gambirplant stem with hooks (Ramulus uncariae cum uncis or Gou Teng).

In one embodiment, there is provided a composition as defined in the first aspect,
15 wherein for (A), the composition comprises at least 2, 3 or 4 of DanGui, Honghua, Taoren, and Deer Horn. Optionally, for (B), the composition comprises at least 4, 5, 6, 7, 8, 9, 10, 11, 12, 13 or 14 of the components as defined in the first aspect. In yet another embodiment, the composition further comprises an agent used in Western medicine for the treatment of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system
20 trauma or conditions related to neuroplasticity.

Optionally, any any one of (A) and (B) are present as separate formulations. Furthermore, any one of the composition as defined above and Western medicine may be present as separate formulations.

According to a second aspect, there is provided a composition as defined in the first
25 aspect for use as a medicament.

According to a third aspect, there is provided the use of the composition as defined above, in the manufacture of a medicament for treating a patient having a condition selected from the group of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma and conditions related to neuroplasticity.

30 According to a fourth aspect, there is provided a product comprising:

a composition as claimed in any one of claims 1 to 3;

76 an agent used in Western medicine for the treatment of stroke, for simultaneous, separate or sequential use in the treatment of patient with a cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to
35 neuroplasticity; and

optionally instructions for use of the product.

According to a fifth aspect, there is provided a kit treating a patient for a cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity, the kit comprising:

a composition as claimed in any one of claims 1 to 3; and
instructions for administering said composition to a patient.

The kit may further comprise an agent used in Western medicine for the treatment of stroke, for simultaneous, separate or sequential use in the treatment of the patient.

5 According to a sixth aspect, there is provided a method of treating a patient having a condition selected from the group of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma and conditions related to neuroplasticity, the method comprising administering to the patient a composition as define above.

10 The method may further comprise administering an agent used in Western medicine for the treatment of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity.

GLOSSARY OF TERMS

15 This section is intended to provide guidance on the interpretation of the words and phrases set forth below (and where appropriate grammatical variants thereof).

As used herein, the term "about" as used in relation to a numerical value means, for example, $\pm 50\%$ or $\pm 30\%$ of the numerical value, preferably $\pm 20\%$, more preferably $\pm 10\%$, more preferably still $\pm 5\%$, and most preferably $\pm 1\%$. Where necessary, the word "about" may be
20 omitted from the definition of the invention.

The term "comprising" means "including" or "consisting". Thus, for example, a composition "comprising" X may consist exclusively of X or may include one or more additional components.

25 The term "treatment" includes any and all uses which remedy a disease state or symptoms, prevent the establishment of disease, or otherwise prevent, hinder, retard, or reverse the progression of disease or other undesirable symptoms in any way whatsoever. Hence, "treatment" includes prophylactic and therapeutic treatment.

30 By "a cerebral stroke patient" we include a patient who has suffered an ischemic or haemorrhagic cerebral stroke. A cerebral stroke is a sudden and permanent death of brain cells that occurs when the flow of blood is blocked and oxygen cannot be delivered to the brain. Ischaemic stroke most commonly occurs when the flow of blood is prevented by clotting (known as 'thrombosis' of the artery) or by a detached clot that lodges in an artery (referred to
35 as an 'embolic stroke'). Haemorrhagic stroke results from rupture of an artery wall, and from blood leaking into the surrounding brain. Haemorrhagic stroke, like ischemic stroke, causes the death of tissue by depriving the brain of blood and oxygen, and results in a number of neurological disabilities (motor, speech) as well as functional disabilities.

The term "stroke" refers to the sudden death of tissue cells due to a lack of oxygen when the blood flow is impaired by blockage or rupture of an artery. Stroke is a vascular accident that can occur in the brain or in the cardiac system. The latter condition is medically known as "myocardial infarction" and more commonly known as a "heart attack". Because of the similarity of both stroke mechanisms, it may be useful to use NeuroAid® to help patients with a heart stroke recovering better from their disability.

DETAILED DESCRIPTION

The present invention provides a new combination treatment for patients having one or more of the following conditions: cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity.

The present invention provides methods and compositions for treating patients having a condition selected from the group consisting of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma and conditions related to neuroplasticity. The patients are administered with:

(A) at least one of DanGui, Honghua, Taoren, and Deer Horn; and

(B) at least one of a composition which comprises at least 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13 or 14 of the following ingredients: Radix Astragali root (Membranous Milkvetch root or Huang Qi), Radix et Rhizoma Salviae Miltiorrhizae root (Red Sage root or Dan Shen), Radix Paeoniae Rubra root (Red Peony root or Chi Shao), rhizome of Ligusticum Chuanxiong (Chuan Xiong), Radix et Rhizoma Notoginseng (Sanqi), Cortex moutan (Peony or Mudanpi), Wood of Odoriferous Rosewood (Jiang Xiang), dried body of Scorpion (Quan Xie), Radix Polygalae root (Yuan Zhi), Grassleaf sweetflag rhizome (Shi Changpu), Leeches (Hirudo or Shuizhi), Ground Beetle (or Tu Bie Chong), Natural or Artificial Cow-bezoar (calculus Bovis artifactus or Rengong Niuhuang), Gambirplant stem with hooks (Ramulus uncariae cum uncis or Gou Teng); and

(i) an agent used in Western medicine for the treatment of stroke.

This invention is based on the discovery that NeuroAid® is not only efficacious in treating stroke patients but that it may be safely used in combination with treatments commonly used in Western medicine for treating stroke patients without adverse side effects such as haemostasis disorders or high blood pressure as described above.

As chronic forms of neurodegeneration such as Alzheimer's and Parkinson's disease (associated with cognitive dysfunction) share common mechanisms of neuronal death with acute forms of neurodegeneration which accompanies stroke, head trauma, cardiac arrest and subarachnoid hemorrhage, NeuroAid® may be useful to improve or treat these brain

disorders as it is believed to have potential activity on neuroprotection or on plasticity. Neuroplasticity (also referred to as brain plasticity or cortical plasticity) refers to changes that occur in the organization of the brain and its circuits of neurons, in particular changes that occur to the location of specific information processing functions. This process supports the learning of new functions as the result of experience during development as mature animals and the creation of new information with healthy neurons by-passing damaged neurons affected by trauma or a medical condition.

Hence, in addition to treating cerebral stroke patients it is envisaged that NeuroAid® or a NeuroAid®-like composition (e.g. a composition according to (i) above), optionally in combination with available agents used in Western medicine, may be useful for the treatment of such neurodegenerative diseases. Typically, the Western medicine used in combination with NeuroAid® or a NeuroAid®-like composition is one that targets a different mechanism from NeuroAid® or a NeuroAid®-like composition. For example, the Western medicine may be antiplatelets and anticoagulants typically used in secondary stroke prevention, and neuroprotectants typically used in improving recovery potential in the acute phase of stroke via mechanisms described below.

NeuroAid® and similar compositions

The ingredients set forth in (i) above may be present in the composition in a relatively crude form (e.g. unprocessed or crushed herbs) or in a more refined form (e.g. purified extracts).

In one embodiment, NeuroAid® is used. NeuroAid® is a TCM product in capsule form comprising 9 herbal components and 5 animal components. NeuroAid® comprises Radix Astragali root (Membranous Milkvetch root or Huang Qi), Radix et Rhizoma Salviae Miltiorrhizae root (Red Sage root or Dan Shen), Radix Paeoniae Rubra root (Red Peony root or Chi Shao), rhizome of Ligusticum Chuanxiong (Chuan Xiong), Radix et Rhizoma Notoginseng (Sanqi), Cortex moutan (Peony or Mudanpi), Wood of Odoriferous Rosewood (Jiang Xiang), dried body of Scorpion (Quan Xie), Radix Polygalae root (Yuan Zhi), Grassleaf sweetflag rhizome (Shi Changpu), Leeches (Hirudo or Shuizhi), Ground Beetle (or Tu Bie Chong), Natural or Artificial Cow-bezoar (calculus Bovis artifactus or Rengong Niu Huang), Gambirplant stem with hooks (Ramulus uncariae cum uncis or Gou Teng). NeuroAid® may further comprise any one or more of DanGui, Honghua, Taoren, or Deer Horn

NeuroAid®, which may be registered under different names in different countries (e.g. in South Africa it is marketed as Strocaid® or Danqi Piantan Jiaonang®) is manufactured by and available commercially in the People's Republic of China from Tianjin Shitian Pharmaceutical Group Co., Ltd (located in the Jianxin Industrial area, Wangwenzhuang town, Xiqing district, Tianjin City, China; Postal Code 300381;). It is also available from Moleac Pte Ltd (formerly

Molecular Acupuncture Pte Ltd), the main licensee outside of the People's Republic of China (11 Biopolis Way, Helios #09-08 Singapore 138667; Tel: 65 64789430; Fax: 6564789435).

For the avoidance of doubt, NeuroAid® not only includes NeuroAid® in the form in which it is currently marketed but also includes future formulations of NeuroAid® which may, for example, be marketed by Tianjin Shitian Pharmaceutical Group Co., Ltd or Moleac Pte Ltd. Such future formulations may, for example, vary in dosage amounts or the concentration of its active ingredients etc.

10 **Combination therapy with NeuroAid®**

In the present invention, NeuroAid® can be used in combination with one or more agents used in Western medicine for the treatment of stroke. By "an agent used in Western medicine", we include any form of mainstream medicine or Western medicine, including dietary supplements. For the avoidance of doubt, by "an agent used in Western medicine", we do not include medicaments used in TCM medicaments or the like.

Examples of suitable agents include antiplatelets, anticoagulants, neuroprotectants, compounds for treating stroke and compounds for activating potassium channels TREK-1. One, two, three, four, five or more of such further agents may, for instance, be used in combination with NeuroAid®. Thus, the following are envisaged as suitable examples for use as combination partner (ii): an antiplatelet; an anticoagulant; a neuroprotectant; a compound for treating stroke; a compound for activating potassium channels TREK-1; an antiplatelet in combination with an anticoagulant; an antiplatelet in combination with a neuroprotectant; an antiplatelet in combination with a compound for treating stroke; an antiplatelet in combination with a compound for activating potassium channels TREK-1; an anticoagulant in combination with a neuroprotectant; an anticoagulant in combination with a compound for treating stroke; an anticoagulant in combination with a compound for activating potassium channels TREK-1; a neuroprotectant in combination with a compound for treating stroke; a neuroprotectant in combination with a compound for activating potassium channels TREK-1; a compound for treating stroke in combination with a compound for activating potassium channels TREK-1; an antiplatelet in combination with an anticoagulant and a neuroprotectant; an antiplatelet in combination with an anticoagulant and a compound for treating stroke; an antiplatelet in combination with an anticoagulant and a compound for activating potassium channels TREK-1; an antiplatelet in combination with a neuroprotectant and a compound for treating stroke; an antiplatelet in combination with a neuroprotectant and a compound for activating potassium channels TREK-1; an antiplatelet in combination with a compound for treating stroke and a compound for activating potassium channels TREK-1; an anticoagulant in combination with a neuroprotectant and a compound for treating stroke; an anticoagulant in combination with a neuroprotectant and a compound for activating potassium channels

TREK-1; an anticoagulant in combination with a compound for treating stroke and a compound for activating potassium channels TREK-1; a neuroprotectant in combination with a compound for treating stroke and a compound for activating potassium channels TREK-1; an antiplatelet in combination with an anticoagulant, a neuroprotectant and a compound for
 5 treating stroke; an antiplatelet in combination with an anticoagulant, a neuroprotectant and a compound for activating potassium channels TREK-1; an antiplatelet in combination with an anticoagulant, a compound for treating stroke and a compound for activating potassium channels TREK-1; an antiplatelet in combination with a neuroprotectant, a compound for treating stroke and a compound for activating potassium channels TREK-1; an anticoagulant
 10 in combination with a neuroprotectant, a compound for treating stroke and a compound for activating potassium channels TREK-1; and an antiplatelet in combination with an anticoagulant, a neuroprotectant, a compound for treating stroke and a compound for activating potassium channels TREK-1.

15 **Antiplatelet agents**

Antiplatelet agents are medications that block the formation of blood clots by preventing the clumping of platelets. Examples of antiplatelet agents include without limitations: Aspirin, the thienopyridine derivatives such as ticlopidine (Ticlid) and clopidogrel (Plavix), the Phosphodiesterase III inhibitors such as Cilostazol (Pletal), Adenosine re-uptake inhibitors
 20 such as Dipyridamole (Persantine or Aggrenox (in combination with aspirin)), and the glycoprotein IIb/IIIa inhibitors such as Abciximab (ReoPro), Eptifibatide (Integrilin) and Tirofiban (Aggrastat) and orally active RGD mimetic prodrugs such as Orbofiban, Sibrafiban, SR121566, or Roxifiban.

25 These agents differ in the way in which they prevent platelets from clumping: for example, Aspirin (Cyclo-oxygenase inhibitor) blocks thromboxan A-2 by inhibiting the enzyme cyclo-oxygenase-1 (COX-1), the thienopyridine derivatives (ADP inhibitors) block the adenosine diphosphate (ADP) receptor on the surface of platelets membrane, and glycoprotein IIb/IIIa inhibitors prevent platelet aggregation by inhibiting a different receptor at the surface of
 30 platelets (the attachment of glycoprotein IIb/IIIa to its receptor is the final step in all pathways that cause platelets aggregation).

One or more antiplatelet agents can be employed in the present invention, for instance, a combination of, for example, 2 or 3 or more antiplatelet agents may be employed.

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Anticoagulant therapies

Antiplatelet agents are part of anticoagulation therapies. There are two other groups of anticoagulant agents which may also be employed in the present invention:

- Inhibitors of clotting factor synthesis such as without limitation, vitamin K antagonists like coumarins or indanedione derivatives (Warfarin or Coumadin).
- Inhibitors of thrombin that include several products such as, but not limited to Heparin (Standard Unfractionated Heparin (UFH), Low Molecular Weight Heparin (LMWH) such as Enoxaparin, Tinzaparin) or to recombinant forms of hirudin such as desirudin and lepirudin (Refludan).

Neuroprotectants

Using various mechanisms, neuroprotectants are compounds that preserve neuronal tissue at risk of dying during stroke and in the aftermath of stroke. Some neuroprotectant agents are sometimes used to treat human stroke patients and include antioxidants (e.g. selenium, 30 vitamin E, vitamin C, glutathione, cysteine, flavinoids, quinolines, enzymes with reducing activity, etc), N-methyl-D-aspartate Receptor Antagonists (Dextrophan, Selfotel, Magnesium), Narcotic Receptor antagonist (Nalmefene (Cervene), Ca-channel blockers, Na-channel modulators (Lubeluzole), Alpha-aminobutyric acid agonist (Clomethiazole), glutamate receptor modulators, serotonin receptor agonists (repinotan), phospholipids, free-radical scavenger (Tirilazad, and NXY-059), astrocyte activation inhibitor (ONO 2506), monoclonal antibodies such as anti-ICAM-1 (Enlimomab), Human anti-leukocytic antibody, Hu23F2G, membrane stabilization agent CDP-choline (Citicholine), Fibroblast growth factor (Fiblast), 20 unsaturated- and polyunsaturated fatty acids, estrogens and selective estrogen receptor modulators (SEAMS), progestins, thyroid hormone and thyroid hormone- mimicking compounds, cyclosporin A and derivatives, thalidomide and derivatives, methylxanthines, Mono-Amine-Oxydase inhibitors (IMAO), serotonin-, noradrenaline and dopamine uptake blockers, dopamine I agonists, L-DOPA, nicotine and derivatives, and NO synthase 25 modulators.

Compounds for treating stroke

Compounds used to treat human stroke patients include Calcium Channel Blockers (D-600, Diltiazem, Nitrendipine, Nimodipine, Nifedipine, Flunarizine, Fluspirilene, Isradipine, 30 Nicardipine, PY 108-068, Verapamil and Triapamil), Calcium chelator (DP-b99), free radical scavengers (Ebselen, Tirilazad, NXY-059), GABA receptor agonists (Diazepam, Baclofen), AMPA agonists (ZK 200775/ MPQX), competitive NMDA antagonists (aptiganel / cerestat, CP 101,606, dextrophan, MK 801/ dizocilpine, remacemide), glycine site antagonists (GV 150526, ACEA 1021), polyamine site antagonists (eliprotil), growth factors (bFGF), Sodium 35 channel blockers (fosphenytoin, 619C89), potassium channel opener (BMS 204352, cromakalim, levcromakalim, aprikalim, pinacidil, diazoxide, nicorandil, minoxidil), piracetam, adenosine transport inhibitor (propentofylline), gangliosides GM (non NMDA antagonist), presynaptic glutamate release inhibitors, clazosentan, desmoteplase, viprinex (ancrod), tenecteplase (TNKase; Metalyse), alteplase, cyclic nitrones, TWEAK (TNF-like weak inducer

of apoptosis) receptor, thrombolytica treatments (urokinase, streptokinase, t-PA / tissue plasminogen activator or recombinant urokinase), anistreplase, riluzole, and disufenton sodium (NXY 059), candesartan, AX-200 (G-CSF, Fligrastim), caffeinol (caffeine+ethanol), enecadin, microplasmin, sonolysis + tPA, V-10153, HTUPA, solulin, piclozotan, S-0139, S-18986, AEOL-10150, AL-208, KN-38-7271, phridoxal 5-phosphate, Neu-2000KL, ONO-2231, PGX-100, RVX-208, SUN-N4057, SUN-N8075, TAT-NR2B9c, GLP-1-expressing stem cell therapy, Msc-1 (SA-4503, AGY-94806)), NH-02D, S-0139 259, tissue protective cytokines (Lu-AA24493), V10153 270 (BB-10153, TAPgen), combined use of statins and other cholesterol lowering drugs, erythropoietin, cerebrolysin and CDP-choline (cytidine-5'-diphosphocholine).

Compounds for activating potassium channels TREK-1

Other suitable agents used in Western medicine include compounds capable of activating the potassium channels TREK-1. The activation of TREK-1 has been found to have a neuroprotective effect. In addition, TREK-1 has been shown to play a major role in neuroprotection against epilepsy and brain and spinal chord ischemia. Since NeuroAid® does not activate potassium channels TREK 1, compounds that are capable of activating the potassium channels TREK-1 may be used in combination with NeuroAid® to enhance the neuroprotective effect of NeuroAid®.

One example of a compound that is capable of activating the potassium channels TREK-1 is Polyunsaturated fatty acids (PUFAs). Polyunsaturated fatty acids are fatty acids that contain more than one double bond. PUFAs can be categorized as methylene-Interrupted Polyenes or Conjugated fatty acids.

Methylene-Interrupted Polyenes are fatty acids that have two or more cis double bonds that are separated from each other by a single methylene group. The essential fatty acids are all omega-3 and -6 methylene-interrupted fatty acids. Examples of Omega-3 fatty acids include without limitation: Alpha-linolenic acid (ALA), Stearidonic acid (STD), Eicosatrienoic acid (ETE), Eicosatetraenoic acid (ETA), Eicosapentaenoic acid (EPA), Docosapentaenoic acid (DPA), Docosahexaenoic acid (DHA), Clupanodonic acid, Tetracosapentaenoic acid, Tetracosahexaenoic acid or Nisinic acid. Examples of Omega-6 fatty acids include without limitation: Linoleic acid (LIN), Gamma-linolenic acid (GLA), Eicosadienoic acid, Dihomo-gamma-linolenic acid (DGLA), Arachidonic acid (AA), Docosadienoic acid, Adrenic acid, Docosapentaenoic acid or Osbond acid. Omega-9 fatty acids are also methylene-Interrupted polyenes, and may be monosaturated or polysaturated. Examples of Omega-9 fatty acids include without limitation: Oleic acid, Eicosenoic acid, Mead acid, Erucic acid or Nervonic acid.

Conjugated fatty acids are fatty acids that have two or more conjugated double bonds. Examples of Conjugated fatty acids include without limitation: Remenic acid, α -Calendic acid, β -Calendic acid, Jacaric acid, α -Eleostearic acid, β -Eleostearic acid, Catalpic acid, Punicic acid, Rumelenic acid, α -Parinaric acid, β -Parinaric acid, Bosseopentaenoic acid.

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Some other PUFAs which are not catergorized as methylene-Interrupted Polyenes or Conjugated fatty include without limitation: Pinolenic acid and podocarpic acid.

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Other compounds that may be capable of activating the potassium channels TREK-1 include the drug Riluzole (Rilutek®), Lysophospholids (LPLs), Caffeic Acid esters and Xenon. These compounds may also be used in combination with NeuroAid ® to enhance the neuroprotective effect of NeuroAid®.

Modes of Administration

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The combination partners (i) and (ii) may be present in a single formulation or may be present as separate formulations. In one embodiment there may be a synergistic effect. As mentioned above, combination partner (ii) may comprise more than one agent, for example, two antiplatelet agents, or an antiplatelet agent and a neuroprotectant may be used.

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The combination partners (i) and (ii) may be administered to the patient at the same time (e.g. simultaneously) or at different times (e.g. sequentially) and over different periods of time, which may be separate from one another or overlapping. The combination partners (i) and (ii) may be administered in any order.

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The combination partner (ii) utilized and the appropriate administration route and dose level will be known to those in the art or could be readily determined by one skilled in the art. Typically, as is well known in the medical art, dosage regimens may depend on various factors including the patient's size, body surface area, age, the particular compound to be administered, sex, time and route of administration, general health, and other drugs being administered concurrently. While individual needs vary, determination of optimal ranges of effective amounts of each component is within the skill of the art. The dosage would be similar to that administered when the agent is used without NeuroAid®.

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Dosage amounts for ticlopidine and for dipyridamole are described in the Physicians' Desk Reference, as are dosage amounts for other antiplatelet and neuroprotectant agents. Dosage amounts of aspirin for the indicated effects are known to those skilled in the medical arts, and generally range from about 20mg to about 325 mg per day. For example, a formulation may contain about 20mg, 30mg, 80 mg, 160 mg, 250 mg, 300 mg, 325 mg or 350 mg of aspirin.

NeuroAid® may be administered orally as such, typically with four 0.4g capsules being taken 3 times a day. For patients with swallowing difficulties, capsules may be opened and powder diluted in water that can be drunk as such or injected via a gastric tube. Hence, a daily dose of about 4.8g is envisaged. In one embodiment, the patient's daily dose of NeuroAid® (or other composition according to (i) above) is about 2g to 8g; 3g to 7g; 4g to 6g; 4.25g to 5.75g; 4.5g to 5.25g; 4.5g to 5g; 4.6g to 4.10g; or 4.7g to 4.9g. A "daily dose" can be a single tablet or capsule etc. or multiple tablets or capsules etc. to be taken on a given day. Suitably, the composition according to (i) is taken orally.

- 10 In one embodiment, each course of NeuroAid® treatment lasts about 4 weeks. Typically 3 courses are administered, most commonly back to back. No therapeutic window is required but additional courses can be added even after a few days of treatment cessation. Hence, in one embodiment, each NeuroAid® treatment lasts about 12 weeks. In another embodiment, the treatment course of NeuroAid® (or other composition according to (i) above) is about 4 to 15 24 weeks; 7 to 16 weeks; 9 to 15 weeks; 10 to 14 weeks; or 11 to 13 weeks.

- In instances of ischemic stroke, treatment with anti-platelet drugs usually commences as soon as possible after onset of stroke symptoms while in instances of haemorrhagic stroke, anticoagulation treatments such as Coumadin or Heparin, are highly contra-indicated and discontinued immediately if they were part of patient's usual treatment. Protamine and vitamin K may be given to reduce bleeding in patients with anticoagulant-induced bleeding.

- In addition to combination partners (i) and (ii), other compounds may be administered, for example, proton pump inhibitors such as Nexium, Protonix and Aciphex. Daily doses of proton pump inhibitors are typically administered to reduce the risk of ulcer development and bleeding in patients under long-term low-dose aspirin or antiplatelet therapy.

BRIEF DESCRIPTION OF DRAWINGS

- 30 * **Figure 1** Phase II clinical trial results on use of NeuroAid® versus BNJ (a) Neurological deficit improvement (DTER scores) before and after treatment with NeuroAid® or BNJ (b) Functional outcomes before and after treatment with NeuroAid® or BNJ.

- 35 * **Figure 2** Phase III clinical trial results on use of NeuroAid® versus BNJ (a) Neurological deficit improvement (DTER scores) before and after treatment with NeuroAid® or BNJ (b) Functional outcomes before and after treatment with NeuroAid® or BNJ.

Figure 3 Clinical study results on the interaction between NeuroAid® and aspirin in healthy volunteers and its effect on blood pressure. (a) systolic blood pressure (b) diastolic blood pressure.

5 * For Figures 1 and 2:

Level of severity of stroke (DTER scores) = LOW, MILD, COMMON, SEVERE

Functional outcomes: 0 pts = able to take care of oneself and speak freely; 2 pts = able to live independently and to do some simple work with some incomplete functions; 4 pts = able to walk and take care of oneself but must be helped partially; 6 pts = able to stand and take a step but must be taken care of at all times; 8 pts = confined to bed.

EXAMPLES

In the following examples, tables and drawings, the term "NeuroAid®" refers to a composition that is the same as NeuroAid®.

EXAMPLE 1

Phase II trial: NeuroAid® versus Buchang Naoxintong Jioanang (BNJ)

A randomized, double-blinded, stratified, control design was adopted for the clinical trial on the efficacy of NeuroAid® in treating patients suffering from apoplexy compared to BNJ which is known for its effectiveness in treating patients suffering from apoplexy (see Example 3). A total of 200 subjects were involved; 100 cases were treated with NeuroAid® while 100 cases were treated with BNJ (control). BNJ is produced and provided by Xianyang Buchang Pharmaceutical Co., Ltd. Four capsules of each drug were administered 3 times daily, with each course of treatment lasting 4 weeks.

The evaluation criteria for neurological and functional recovery from apoplexy (DTER scoring diagnostic standard) and TCM symptom therapeutic effects (TCM diagnostic symptom scoring standard) were assessed in accordance with the Clinical Guiding Principles for the Treatment of Apoplexy with New Chinese Herbs promulgated by the Ministry of Health of the PRC in 1993.

Severity of symptoms in the DTER scoring standard was classified according to 4 levels (SEVERE, COMMON, MILD and LOW) while functional outcomes were classified in points from 0 to 8.

The data (Figure 1) demonstrated NeuroAid®'s clinical efficacy and superiority in improving patients' neurological deficit and functional (autonomy/dependency post-stroke) outcomes versus that of the control treatment BNJ.

EXAMPLE 2**Phase III trial: NeuroAid® versus BNJ**

A randomized, double-blinded, stratified, control design was adopted. A total of 405 subjects were involved, where 300 cases were treated NeuroAid® while 105 cases were treated with the control drug BNJ produced and provided by Xianyang Buchang Pharmaceutical Co., Ltd. Four capsules of each drug were administered 3 times daily, with each course of treatment lasting 4 weeks.

The evaluation criteria for neurological and functional recovery from apoplexy (DTER scoring diagnostic standard) and TCM symptom therapeutic effects (TCM diagnostic symptom scoring standard) were assessed as in the Phase II trial.

The data (Figure 2) demonstrated that NeuroAid® was superior to BNJ in improving the patients' neurological deficit particularly in helping patients recover from their hemi-paralysis. With regard to functional outcomes, even if NeuroAid® failed to demonstrate any superiority to BNJ, both treatments had comparable effect and about 50% of the stroke patients included in the study returned to a functionality dependent state after 4 weeks of treatment (comprehensive function score similar to Rankin score used in the West equal to 0 or 2).

No adverse effects of treatment with NeuroAid® were observed in the Phase II or Phase III trials. The finding of a broad therapeutic window (>14 days after stroke onset) in both trials also gives a huge advantage over fibrinolytic agents, which have to be given within the first 3 to 6 hours.

EXAMPLE 3**Comparative study on efficacy of BNJ versus Citicoline and aspirin**

A randomized, double-blinded, stratified, control design was adopted for the comparative study on the efficacy of BNJ and Citicoline in treating patients with apoplexy. Citicoline is the only putative neuro-protectant that has shown results in Western randomized, double-blinded trials given within 24 hours after symptom onset. Davalos et al. (Oral Citicoline in Acute Ischemic Stroke. An Individual Patient Data Pooling Analysis of Clinical Trials (2002) *Stroke* 33:2850) documents the ability of Citicoline to improve complete recovery at 3 months.

A first comparative study performed in the PRC involved 150 subjects treated with Citicoline (0.5g IV) for 15 days in combination with a TCM (Xueshuantong) and 160 subjects treated with Citicoline (0.5g IV) for 15 days in combination with Xueshuantong and BNJ. The latter group treated with BNJ showed improvements in scores on a neuro-functional defects scale, plasma viscosity level and cholesterol level.

A second comparative study compared 30 subjects treated with aspirin (150mg), Citicoline (0.75g IV) and salvia miltiorrhizae injection (60mL), with 30 subjects treated with aspirin (150mg), Citicoline (0.75g IV), salvia miltiorrhizae injection (60mL) and BNJ for 1 month. The BNJ-treated group showed significantly better results in neurological and functional outcomes.

5

Citicoline (CDP-choline) is a key intermediary in the biosynthesis of phosphatidylcholine, an important component of the neural cell membrane that stabilizes cells membranes and inhibits the formation of cytotoxic free fatty acids. It has been shown to produce beneficial effects in both animal models and clinical stroke trials. A significant difference between the groups, favoring citicoline treatment, was seen in terms of functional outcome as measured by the Barthel Index and Rankin scale, neurologic evaluation as measured by the National Institutes of Health (NIH) stroke scale, and cognitive function as measured by the Mini Mental Status Examination. ¹As efficacy trials on Citicoline have been demonstrated to be superior to the placebo, it can be inferred from the above two comparative studies that treatment with BNJ would likewise be superior to a placebo.

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EXAMPLE 4

Safety Study: Phase IV open-label study on effect of NeuroAid® on hemostasis

Blood samples of 30 healthy individuals were collected before they received NeuroAid® and 2 and 8 hours after NeuroAid® initiation (4 capsules). Five coagulation tests were performed on the blood samples.

20

Table 1.

Quick Prothrombine Time			
<i>Blood Sampling</i>	<i>First (%)</i>	<i>Second (%)</i>	<i>Third (%)</i>
Average	12.75	12.71	12.92
Std Deviation	0.68	0.72	0.67
Activated Partial Prothrombine Time			
<i>Blood Sampling</i>	<i>First (%)</i>	<i>Second (%)</i>	<i>Third (%)</i>
Average	37.64	37.15	38.66
Std Deviation	4.28	4.44	4.26
Fibrinogen			
<i>Blood Sampling</i>	<i>First (%)</i>	<i>Second (%)</i>	<i>Third (%)</i>
Average	2.98	3.05	3.06
Std Deviation	0.61	0.59	0.57
Platelet Aggregation			
<i>Blood Sampling</i>	<i>First (%)</i>	<i>Second (%)</i>	<i>Third (%)</i>
Average	63.01	62.19	61.52

Std Deviation	15.50	12.08	13.37
D-Dimer			
<i>Blood Sampling</i>	<i>First (%)</i>	<i>Second (%)</i>	<i>Third (%)</i>
Control Device	2.7	2.4	1.9

The results confirmed that NeuroAid® has no effect on hemostasis blood factors and thus does not increase bleeding risks.

5 EXAMPLE 5

Safety Study: Phase IV open-label study on potential interaction between NeuroAid® and acetylsalicylic acid (aspirin) in healthy volunteers and its effect on hemostasis and blood pressure

An open randomized one-day study was conducted on 11 healthy volunteers receiving NeuroAid® treatment (12 capsules per day) and aspirin (ASA 300mg per day) from day 1 to day 5, and 11 healthy volunteers receiving aspirin alone (300mg per day) from day 1 to day 5. Blood samples of the subjects were collected before they received treatment and 2 and 8 hours after treatment initiation. The blood samples were analyzed with the 5 coagulation tests, namely quick prothrombine time, activated partial prothrombine time, fibrinogen dosage, platelet aggregation and D-dimer, to identify the coagulation mechanisms triggered by potential interaction between NeuroAid® and aspirin. Blood pressure of the 22 subjects was also measured before treatment, and 2, 4, 6 and 8 hours after treatment initiation.

Results of this trial show that NeuroAid® can be safely associated with aspirin and does not have any interaction with aspirin regarding blood coagulation (Table 2) and blood pressure (Figure 3).

Table 2.

Quick Prothrombine Time (Seconds)	
Aspirin alone	Mean
T = 0	12.9
T + 2 hours	12.6
T + 8 hours	12.4
Aspirin + NeuroAid®	Mean
T = 0	12.7
T + 2 hours	12.6
T + 8 hours	12.4
Activated Partial Prothrombine Time (Seconds)	
Aspirin alone	Mean
T = 0	38.8

T + 2 hours	39.0
T + 8 hours	38.2
Aspirin + NeuroAid®	Mean
T = 0	37.1
T + 2 hours	38.2
T + 8 hours	37.2
Fibrinogen (G/L)	
Aspirin alone	Mean
T = 0	3.2
T + 2 hours	3.2
T + 8 hours	3.0
Aspirin + NeuroAid®	Mean
T = 0	3.2
T + 2 hours	3.0
T + 8 hours	2.9
Platelet Aggregation (Percentage)	
Aspirin alone	Mean
T = 0	62.82
T + 2 hours	62.87
T + 8 hours	48.01
Aspirin + NeuroAid®	Mean
T = 0	61.82
T + 2 hours	58.28
T + 8 hours	46.98
D-Dimer (μG/L)	
Aspirin alone	Mean
T = 0	0.16
T + 2 hours	0.16
T + 8 hours	0.16
Aspirin + NeuroAid®	Mean
T = 0	0.13
T + 2 hours	0.09
T + 8 hours	0.15

EXAMPLE 6

Safety Study: Pilot open-label study on potential interaction between NeuroAid® and Western drugs in ischemic stroke patients and its effect on hemostasis and blood pressure

A pilot open-label study was conducted on 10 ischemic stroke patients within the first week of the onset of stroke (early phase). The test patients were selected based on the following criteria:

- a) aged above 18 years old;
- b) have had a cerebral infarction with compatible imaging on Computed Tomography (CT) scan
- 5 or Magnetic Resonance Imaging (MRI); and
- c) the time window between stroke onset and the open-label study was less than one week,

Conversely, patients that were excluded from the study include:

- a) female patients who were pregnant, lactating or nursing;
- 10 b) patients showing signs of intra-cerebral hemorrhage on brain Computed Tomography (CT) scan or Magnetic Resonance Imaging (MRI);
- c) patients having a history of easy bruising or blood coagulation disorders;
- d) patients receiving other Traditional Chinese Medicine than NeuroAid®;
- e) patients who received thrombolysis;
- 15 f) patients who have used NeuroAid® within a 3-month period prior to screening and enrolment in the open-label study.

Each patient received 4 capsules of NeuroAid® 3 times a day for one month in addition to the other Western medicine the patient was receiving. The Western medicine includes platelet

20 aggregation inhibitors, nitrates, oral anti-hypertensive drugs, lipid regulating drugs, oral anti-diabetic, or anti-convulsant drugs.

Safety assessment tests were performed on blood samples collected from the test patients at three intervals: a) before NeuroAid® intake (to form the baseline); b) 1 week after NeuroAid®

25 initiation; and c) 4 weeks after NeuroAid® initiation. The blood samples were analyzed using the following hematology and biochemical laboratory analytical tests:

- a) Prothrombine Time (PT);
- b) Activated Partial Prothrombine Time (APPT);
- c) fibrinogene dosage;
- 30 d) platelet aggregation;
- e) D-dimer test;
- f) blood cell count;
- g) creatinine;
- h) SGOT SGPT;
- 35 i) glycemia; and
- j) CRP-C-reactive protein.

The results of the open-label study (Table 3) demonstrated that NeuroAid® does not have any effect on renal and hepatic functions, glycemia, and C-Reactive-protein when used in combination with Western medicine for managing risk factors of secondary stroke. For example, in the five coagulation tests (Prothrombine Time (PT), Activated Partial Prothrombine Time (APPT), fibrinogene dosage, platelet aggregation, D-dimer test) performed on Patients 4 and 5 (both receiving NeuroAid® and four different types of Western medicine) (Table 3, columns 5 and 6), the test readings taken 1 week and 4 weeks after NeuroAid® initiation do not differ significantly from the baseline readings taken before the start of NeuroAid® intake.

- 10 The statistical analysis of the readings for the open-label study (Table 4) also show that the readings from all 10 patients do not differ significantly from the mean values. For example, in the Prothrombine Time (PT) test, the Standard Deviation (SD) values after 1 week and 4 weeks of NeuroAid® initiation are 1.09 and 0.77, respectively, which do not deviate significantly from the baseline value of 0.88. Similarly, in the fibrinogen dosage test, the
- 15 Standard Deviation (SD) values after 1 week and 4 weeks of NeuroAid® initiation are 1.22 and 1.11, respectively, which do not deviate significantly from the baseline value of 1.13.

Hence, the results of the open-label study further demonstrated that NeuroAid® can be safely used in combination with Western medicine such as platelet aggregation inhibitors, nitrates, oral

20 anti-hypertensive drugs, lipid regulating drugs, oral anti-diabetic, or anti-convulsant drugs.

Table 3.

	PATIENT 1	PATIENT 2	PATIENT 3	PATIENT 4	PATIENT 5	PATIENT 6	PATIENT 7	PATIENT 8	PATIENT 9	PATIENT 10
Gender	F	F	F	M	F	F	F	M	M	M
Age	79	69	61	58	53	60	64	85	48	69
Days from stroke onset	3	4	1	2	7	7	1	1	4	7
On Aspirin?	N	Y	N	Y	N	N	N	Y	Y	Y
Other Treatment administered	NeuroAid®, ISMN, nimodipine	NeuroAid®, aspirin, nimodipine, gemfibrozil	NeuroAid®, ISMN, metformin, glipizide	NeuroAid®, aspirin, nimodipine, glicazide, indapamide	NeuroAid®, ISMN, nimodipine, amlodipine, KCl	NeuroAid®, digoxine, isosorbide mononitride, nimodipine, phenytoin	NeuroAid®, phenytoin sodium, nimodipine	NeuroAid®, aspirin, ISMN	NeuroAid®, aspirin, ISMN, amlodipine	NeuroAid®, aspirin, ISMN, amlodipine
NIHSS										
Baseline	19	1	16	7	6	4	6	4	3	2
1 Week	17	0	8	4	4	12	4	3	3	1
1 Month	15	1	7	4	3	10	2	2	2	1
Rankin score										
Baseline	5	1	5	4	3	4	4	2	3	1
1 Week	5	1	4	2	1	5	2	2	2	1
1 Month	5	1	4	1	1	4	1	1	1	1
PT (secs)										
Baseline	14.3	13.0	11.6	12.8	13.8	11.3	13.3	13.9	12.1	12.6
1 Week	12.7	11.6	11.1	12.4	14.4	11.5	12.8	13.9	12.0	11.6
1 Month	13.7	12.0	11.5	11.8	13.6	11.2	11.9	12.2	12.4	12.6
APTT (secs)										
Baseline	27.1	32.1	29.0	33.5	33.1	27.9	29.3	33.5	30.3	30.7
1 Week	31.8	29.5	28.6	31.5	38.2	30.8	34.3	33.2	32.1	29.7
1 Month	29.6	29.9	30.5	32.2	28.5	24.9	29.8	30.2	34.7	30.7

[illegible]

Basophil	0.0	0.1	0.1	0.1	0.2	0.1	0.4	0.2	0.1	0.1	0.1
Baseline	0.0	0.1	0.1	0.1	0.2	0.1	0.1	0.2	0.1	0.1	0.1
1 Week	0.1	0.1	0.2	0.2	0.1	0.1	0.1	0.2	0.1	0.1	0.1
1 Month	0.1	0.1	0.1	0.3	0.1	0.1	0.1	0.2	0.1	0.1	0.1
Basophil											
Baseline	0.0	0.0	0.0	0.0	0.0	0.1	0.1	0.0	0.1	0.1	0.1
1 Week	0.0	0.0	0.0	0.0	0.0	0.0	0.1	0.0	0.1	0.1	0.1
1 Month	0.0	0.0	0.1	0.1	0.0	0.0	0.1	0.0	0.1	0.1	0.1
Hemoglobin (Hb)											
High											
Baseline	138	140	134	163	109	141	130	137	162	132	132
1 Week	131	139	138	167	120	147	132	137	161	146	146
1 Month	119	148	138	142	116	153	133	162	157	158	158
Hematocrit (Hct)											
High											
Baseline	40.1	40.1	38.2	44.9	31.0	39.5	37	41.0	46.1	38.7	38.7
1 Week	45.8	39.8	38.8	47.1	33.5	42.4	38.7	39.2	47.3	41.5	41.5
1 Month	34.3	43.4	39.4	41.1	33.5	43.0	38.9	44.6	44.6	45.0	45.0
CRP (mg/dL)											
Baseline	6.34	0.99	6.42	0.49	0.96	0.14	3.53	2.17	0.69	0.17	0.17
1 Week	11.10	0.48	0.64	0.44	0.16	3.73	2.1	0.35	0.13	0.10	0.10
1 Month	0.64	0.40	0.65	5.44	0.74	1.10	1.88	0.10	0.38	0.10	0.10

Table 4.

SUMMARY	MEAN	SD -	SD +	SD
Gender	F=5	M=4		
Age	65.11	53.24	76.98	11.87
Day(s) from stroke onset	3.33	0.94	5.73	2.40
On Aspirin?	Y=5	N=4		
Other treatment administered				
NIHSS				
Baseline	7.11	0.86	13.36	6.25
1 Week	4.89	-0.17	9.95	5.06
1 Month	4.11	-0.37	8.60	4.48
Rankin score				
Baseline	3.11	1.57	4.65	1.54
1 Week	2.22	0.83	3.62	1.39
1 Month	1.78	0.21	3.34	1.56
PT (secs)				
Baseline	13.04	12.17	13.92	0.88
1 Week	12.50	11.41	13.59	1.09
1 Month	12.41	11.64	13.18	0.77
APTT (secs)				
Baseline	30.96	28.70	33.21	2.26
1 Week	32.10	29.17	35.03	2.93
1 Month	30.68	28.87	32.48	1.80
Fibrinogen/dosage (g/L)				
Baseline	4.10	2.97	5.23	1.13
1 Week	4.03	2.81	5.25	1.22
1 Month	4.06	2.95	5.18	1.11
Platelet aggregation				
Baseline	63.61	54.67	72.55	8.94
1 Week	52.31	40.68	63.94	11.63
1 Month	48.44	36.31	60.58	12.14
D-Dimer (ng/mL)				
Baseline	0.25	0.08	0.42	0.17
1 Week	0.45	-0.12	1.02	0.57
1 Month	0.49	-0.23	1.22	0.73
SGPT - ALT (IU/L)				
Baseline	13.87	9.75	17.98	4.11
1 Week	20.80	11.13	30.47	9.67
1 Month	17.77	10.48	25.06	7.29
SGOT - AST (IU/L)				
Baseline	19.14	9.03	29.26	10.11
1 Week	18.09	12.00	24.18	6.09
1 Month	16.12	12.26	19.99	3.87
Glycemia (mmol/L)				
Baseline	5.88	3.60	8.16	2.28
1 Week	5.21	3.44	6.98	1.77
1 Month	4.91	3.81	6.02	1.10
Creatinine (μmol/L)				

Baseline	70.56	48.97	92.14	21.58
1 Week	75.56	55.81	95.30	19.75
1 Month	67.89	48.94	86.84	18.95
Red blood cells				
Baseline	4.42	3.87	4.96	0.55
1 Week	4.51	4.00	5.02	0.51
1 Month	4.56	4.01	5.11	0.55
Mean RBC volume				
Baseline	90.71	85.73	95.70	4.99
1 Week	90.93	86.00	95.86	4.93
1 Month	90.09	84.36	95.82	5.73
white blood cells				
Baseline	6.63	4.33	8.94	2.30
1 Week	5.88	4.10	7.65	1.77
1 Month	6.68	3.79	9.57	2.89
granulocytes				
Baseline	4.67	2.83	6.51	1.84
1 Week	4.10	2.43	5.77	1.67
1 Month	4.43	2.43	6.44	2.01
lymphocytes				
Baseline	1.41	0.65	2.17	0.76
1 Week	1.68	0.77	2.59	0.91
1 Month	1.74	1.01	2.48	0.74
monocytes				
Baseline	0.31	0.15	0.47	0.16
1 Week	0.29	0.15	0.43	0.14
1 Month	0.34	0.19	0.50	0.15
eosinophiles				
Baseline	0.11	0.05	0.17	0.06
1 Week	0.13	0.08	0.18	0.05
1 Month	0.13	0.06	0.20	0.07
basophiles				
Baseline	0.03	-0.02	0.08	0.05
1 Week	0.02	-0.02	0.07	0.04
1 Month	0.04	-0.01	0.10	0.05
hemoglobin (Hb) in g/L				
Baseline	139.56	123.41	155.70	16.15
1 Week	142.89	128.42	157.36	14.47
1 Month	143.67	126.96	160.38	16.71
hematocrit (Hct) in %				
Baseline	39.96	35.65	44.26	4.31
1 Week	41.71	37.19	46.23	4.52
1 Month	40.99	36.58	45.40	4.41
CRP (mg/dL)				
Baseline	2.04	-0.49	4.57	2.53
1 Week	1.90	-1.73	5.54	3.63
1 Month	1.06	-0.61	2.73	1.67

EXAMPLE 7**Effect of NeuroAid® on specific potassium channels**

NeuroAid® powder was dissolved in distilled water at a concentration of 1 µg/ml. After 30
5 minutes infusion, the liquid was filtered on a 0.22µm filter.

Cos-7 cells, cultured in Dulbecco's Modified Eagle Medium (DMEM; Gibco, Invitrogen, San
Diego, CA) supplemented with 10% fetal bovine serum (Perbio, Thermo Fisher Scientific Inc. ,
Waltham, MA), 100 U/ml penicillin and 100 µg/ml streptomycin (Gibco, Invitrogen, San Diego,
10 CA) in a humidified incubator with 5% CO₂ at 37°C, were used for electrophysiological
procedures. Cells were transfected using DEAE-Dextran (Sigma-Aldrich Corporation, St-Louis,
MO) with 0.5 µg of TREK 1 or TREK 2 cloned in pIRES-EGFP plasmid (Clontech Palo Alto, CA)
expressing the green fluorescent protein.

15 Recordings were realised 48 hours after transfection. For electrophysiological recordings, the
internal solution contained 155mM KCl, 3mM MgCl₂ , 5mM EGTA and 10mM HEPES (Sigma-
Aldrich Corporation, St-Louis, MO) at pH 7.2 with KOH and the external solution contained
150mM NaCl, 5mM KCl, 3 mM MgCl₂, 1mM CaCl₂ and 10mM HEPES at pH 7.4 with NaOH.
Effects of NeuroAid® on TREK currents were measured on whole-cell currents, elicited by
20 voltage ramps (from -100mV to +100mV, 1.5 seconds in duration) before and during application
of NeuroAid® (5 minutes). TREK 1 and TREK 2 currents were not different before and after 5
minutes of NeuroAid® application, comparing with control cells transfected with EGFP only.

The results show that NeuroAid® does not have any effect on potassium channels TREK-1.
25 Hence, the results suggest that agents that activate potassium channels TREK-1 and NeuroAid®
have different mechanisms and different targets. Accordingly, it may be beneficial to combine
agents that activate potassium channels TREK-1 with NeuroAid® to enhance the neuroprotective
effect of NeuroAid®.

30 It will be understood that the invention has been described by way of example only and
modifications may be made whilst remaining within the scope and spirit of the invention.

CLAIMS

1. A composition comprising:
- 5 (A) at least one of the following components: DanGui, Honghua, Taoren, and Deer Horn; and
- 10 (B) at least one of the following components: Radix Astragali root (Membranous Milkvetch root or Huang Qi), Radix et Rhizoma Salviae Miltiorrhizae root (Red Sage root or Dan Shen), Radix Paeoniae Rubra root (Red Peony root or Chi Shao), rhizome of Ligusticum Chuanxiong (Chuan Xiong), Radix et Rhizoma Notoginseng (Sanqi), Cortex moutan (Peony or Mudanpi), Wood of Odoriferous Rosewood (Jiang Xiang), dried body of Scorpion (Quan Xie), Radix Polygalae root (Yuan Zhi), Grassleaf sweetflag rhizome (Shi Changpu), Leeches (Hirudo or Shuizhi), Ground Beetle (or Tu Bie Chong), Natural or Artificial Cow-bezoar (calculus Bovis artifactus or Rengong Niuhuang), Gambirplant stem with hooks
- 15 (Ramulus uncariae cum uncis or Gou Teng).
2. A composition as claimed in claim 1, wherein for (A), the composition comprises at least 2, 3 or 4 of DanGui, Honghua, Taoren, and Deer Horn.
- 20 3. A composition as claimed in claim 1 or claim 2, wherein for (B), the composition comprises at least 4, 5, 6, 7, 8, 9, 10, 11, 12, 13 or 14 of the components.
- 25 4. A composition as claimed in claim 1 or claim 2 or claim 3, further comprising an agent used in Western medicine for the treatment of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity.
- 30 5. A composition as claimed in any one of the preceding claims, wherein any one of (A) and (B) are present as separate formulations.
- 36 6. A composition as claimed in claim 4, wherein any one of the composition and Western medicine are present as separate formulations.
- 35 7. A composition as claimed in any of claims 4 to 6, where the Western medicine is selected from the group consisting of Calcium Channel Blockers (D-600, Diltiazem, Nitrendipine, Nimodipine, Nifedipine, Flunarizine, Fluspirilene, Isradipine, Nicardipine, PY 108-068, Verapamil and Triapamil), Calcium chelator (DP-b99), free radical scavengers (Ebselen, Tirilazad, NXY-059), GABA receptor

agonists (Diazepam, Baclofen), AMPA agonists (ZK 200775/ MPQX), competitive NMDA antagonists (aptiganel / cerestat, CP 101,606, dextrophan, MK 801/ dizocilpine, remacemide), glycine site antagonists (GV 150526, ACEA 1021), polyamine site antagonists (eliprodil), growth factors (bFGF), Sodium channel blockers (fosphenytoin, 619C89), potassium channel opener (BMS 204352, cromakalim, levromakalim, aprikalim, pinacidil, diazoxide, nicorandil, minoxidil), piracetam, adenosine transport inhibitor (propentofylline), gangliosides GM (non NMDA antagonist), presynaptic glutamate release inhibitors, clazosentan, desmoteplase, viprinex (ancrod), tenecteplase (TNKase; Metalyse), alteplase, cyclic nitrones, TWEAK (TNF-like weak inducer of apoptosis) receptor, thrombolytica treatments (urokinase, streptokinase, t-PA / tissue plasminogen activator or recombinant urokinase), anistreplase, riluzole, and disufenton sodium (NXY 059), candesartan, AX-200 (G-CSF, Fligrastim), caffeineol (caffeine+ethanol), enecadin, microplasmin, sonolysis + tPA, V-10153, HTUPA, solulin, piclozotan, S-0139, S-18986, AEOL-10150, AL-208, KN-38-7271, phridoxal 5-phosphate, Neu-2000KL, ONO-2231, PGX-100, RVX-208, SUN-N4057, SUN-N8075, TAT-NR2B9c, GLP-1-expressing stem cell therapy, Msc-1 (SA-4503, AGY-94806)), NH-02D, S-0139 259, tissue protective cytokines (Lu-AA24493), V10153 270 (BB-10153, TAPgen), combined use of statins and other cholesterol lowering drugs, erythropoietin, cerebrolysin and CDP-choline (cytidine-5'-diphosphocholine).

8. A composition as claimed in any of claims 4 to 6, where the Western medicine is capable of activating potassium channels TREK-1.

9. A composition as claimed in any one of the preceding claims for use as a medicament.

10. Use of the composition as claimed in any one of claims 1 to 4, in the manufacture of a medicament for treating a patient having a condition selected from the group of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma and conditions related to neuroplasticity.

11. Use of the composition as claimed in any one of claims 1 to 3, in the manufacture of a medicament for treating a patient having a condition selected from the group of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma and conditions related to neuroplasticity, where the patient also receives an agent used in Western medicine for the treatment of

patients with cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity.

12. The use of claim 11, wherein said Western medicine is selected from the group consisting of Calcium Channel Blockers (D-600, Diltiazem, Nitrendipine, Nimodipine, Nifedipine, Flunarizine, Fluspirilene, Isradipine, Nicardipine, PY 108-068, Verapamil and Triapamil), Calcium chelator (DP-b99), free radical scavengers (Ebselen, Tirilazad, NXY-059), GABA receptor agonists (Diazepam, Baclofen), AMPA agonists (ZK 200775/ MPQX), competitive NMDA antagonists (aptiganel / cerestat, CP 101,606, dextrophan, MK 801/ dizocilpine, remacemide), glycine site antagonists (GV 150526, ACEA 1021), polyamine site antagonists (eliprodil), growth factors (bFGF), Sodium channel blockers (fosphenytoin, 619C89), potassium channel opener (BMS 204352, cromakalim, levromakalim, aprikalim, pinacidil, diazoxide, nicorandil, minoxidil), piracetam, adenosine transport inhibitor (propentofylline), gangliosides GM (non NMDA antagonist), presynaptic glutamate release inhibitors, clazosentan, desmoteplase, viprinex (ancrod), tenecteplase (TNKase; Metalyse), alteplase, cyclic nitrones, TWEAK (TNF-like weak inducer of apoptosis) receptor, thrombolytica treatments (urokinase, streptokinase, t-PA / tissue plasminogen activator or recombinant urokinase), anistreplase, riluzole, and disufenton sodium (NXY 059), candesartan, AX-200 (G-CSF, Fligrastim), caffeinol (caffeine+ethanol), enecadin, microplasmin, sonolysis + tPA, V-10153, HTUPA, solulin, piclozotan, S-0139, S-18986, AEOL-10150, AL-208, KN-38-7271, phridoxal 5-phosphate, Neu-2000KL, ONO-2231, PGX-100, RVX-208, SUN-N4057, SUN-N8075, TAT-NR2B9c, GLP-1-expressing stem cell therapy, Msc-1 (SA-4503, AGY-94806)), NH-02D, S-0139 259, tissue protective cytokines (Lu-AA24493), V10153 270 (BB-10153, TAPgen), combined use of statins and other cholesterol lowering drugs, erythropoietin, cerebrolysin and CDP-choline (cytidine-5'-diphosphocholine).

13. The use of claim 11, wherein said Western medicine is capable of activating potassium channel TREK-1.

14. A product comprising:

- (i) a composition as claimed in any one of claims 1 to 3;
- (ii) an agent used in Western medicine for the treatment of stroke, for simultaneous, separate or sequential use in the treatment of patient with a cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity; and

(iii) optionally instructions for use of the product.

15. The product as claimed in claim 14, wherein said Western medicine is selected from the group consisting of Calcium Channel Blockers (D-600, Diltiazem, Nitrendipine, Nimodipine, Nifedipine, Flunarizine, Fluspirilene, Isradipine, Nicardipine, PY 108-068, Verapamil and Triapamil), Calcium chelator (DP-b99), free radical scavengers (Ebselen, Tirilazad, NXY-059), GABA receptor agonists (Diazepam, Baclofen), AMPA agonists (ZK 200775/ MPQX), competitive NMDA antagonists (aptiganel / cerestat, CP 101,606, dextrophan, MK 801/ dizocilpine, remacemide), glycine site antagonists (GV 150526, ACEA 1021), polyamine site antagonists (eliprodil), growth factors (bFGF), Sodium channel blockers (fosphenytoin, 619C89), potassium channel opener (BMS 204352, cromakalim, levcromakalim, aprikalim, pinacidil, diazoxide, nicorandil, minoxidil), piracetam, adenosine transport inhibitor (propentofylline), gangliosides GM (non NMDA antagonist), presynaptic glutamate release inhibitors, clazosentan, desmoteplase, viprinex (ancrod), tenecteplase (TNKase; Metalyse), alteplase, cyclic nitrones, TWEAK (TNF-like weak inducer of apoptosis) receptor, thrombolytica treatments (urokinase, streptokinase, t-PA / tissue plasminogen activator or recombinant urokinase), anistreplase, riluzole, and disufenton sodium (NXY 059), candesartan, AX-200 (G-CSF, Fligrastim), caffeineol (caffeine+ethanol), enecadin, microplasmin, sonolysis + tPA, V-10153, HTUPA, solulin, piclozotan, S-0139, S-18986, AEOL-10150, AL-208, KN-38-7271, phridoxal 5-phosphate, Neu-2000KL, ONO-2231, PGX-100, RVX-208, SUN-N4057, SUN-N8075, TAT-NR2B9c, GLP-1-expressing stem cell therapy, Msc-1 (SA-4503, AGY-94806)), NH-02D, S-0139 259, tissue protective cytokines (Lu-AA24493), V10153 270 (BB-10153, TAPgen), combined use of statins and other cholesterol lowering drugs, erythropoietin, cerebrolysin and CDP-choline (cytidine-5'-diphosphocholine).
16. The product as claimed in claim 14, wherein said Western medicine is capable of activating potassium channel TREK-1.
17. A kit for treating a patient for a cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity, the kit comprising:

 - (i) a composition as claimed in any one of claims 1 to 3; and
 - (ii) instructions for administering said composition to a patient.
18. A kit as claimed in claim 17, wherein the kit further comprises:

- (ii) an agent used in Western medicine for the treatment of stroke, for simultaneous, separate or sequential use in the treatment of the patient.

5 19. The kit as claimed in claim 18, wherein said Western medicine is selected from the group consisting of Calcium Channel Blockers (D-600, Diltiazem, Nitrendipine, Nimodipine, Nifedipine, Flunarizine, Fluspirilene, Isradipine, Nicardipine, PY 108-068, Verapamil and Triapamil), Calcium chelator (DP-b99), free radical scavengers (Ebselen, Tirilazad, NXY-059), GABA receptor agonists (Diazepam, Baclofen), AMPA agonists (ZK 200775/ MPQX), competitive NMDA
10 antagonists (aptiganel / cerestat, CP 101,606, dextrophan, MK 801/ dizocilpine, remacemide), glycine site antagonists (GV 150526, ACEA 1021), polyamine site antagonists (eliprodil), growth factors (bFGF), Sodium channel blockers (fosphenytoin, 619C89), potassium channel opener (BMS 204352, cromakalim, levromakalim, aprikalim, pinacidil, diazoxide, nicorandil, minoxidil), piracetam,
15 adenosine transport inhibitor (propentofylline), gangliosides GM (non NMDA antagonist), presynaptic glutamate release inhibitors, clazosentan, desmoteplase, viprinex (ancrod), tenecteplase (TNKase; Metalyse), alteplase, cyclic nitrones, TWEAK (TNF-like weak inducer of apoptosis) receptor, thrombolytica treatments (urokinase, streptokinase, t-PA / tissue plasminogen activator or recombinant
20 urokinase), anistreplase, riluzole, and disufenton sodium (NXY 059), candesartan, AX-200 (G-CSF, Fligrastim), caffeinol (caffeine+ethanol), enecadin, microplasmin, sonolysis + tPA, V-10153, HTUPA, solulin, piclozotan, S-0139, S-18986, AEOL-10150, AL-208, KN-38-7271, phridoxal 5-phosphate, Neu-2000KL, ONO-2231, PGX-100, RVX-208, SUN-N4057, SUN-N8075, TAT-NR2B9c, GLP-
25 1-expressing stem cell therapy, Msc-1 (SA-4503, AGY-94806)), NH-02D, S-0139 259, tissue protective cytokines (Lu-AA24493), V10153 270 (BB-10153, TAPgen), combined use of statins and other cholesterol lowering drugs, erythropoietin, cerebrolysin and CDP-choline (cytidine-5'-diphosphocholine).

30 20. The kit as claimed in claim 18, wherein said Western medicine is capable of activating potassium channel TREK-1.

26 21. A method of treating a patient having a condition selected from the group of cerebral stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous
35 system trauma and conditions related to neuroplasticity, the method comprising administering to the patient a composition as claimed in any one of claim 1 to 3.

22. A method as claimed in claim 21, wherein the method further comprises administering an agent used in Western medicine for the treatment of cerebral

stroke, heart stroke, neurodegenerative diseases, brain trauma, nervous system trauma or conditions related to neuroplasticity.

23. The method as claimed in claim 22, wherein said Western medicine is selected from the group consisting of Calcium Channel Blockers (D-600, Diltiazem, Nitrendipine, Nimodipine, Nifedipine, Flunarizine, Fluspirilene, Isradipine, Nicardipine, PY 108-068, Verapamil and Triapamil), Calcium chelator (DP-b99), free radical scavengers (Ebselen, Tirilazad, NXY-059), GABA receptor agonists (Diazepam, Baclofen), AMPA agonists (ZK 200775/ MPQX), competitive NMDA antagonists (aptiganel / cerestat, CP 101,606, dextrophan, MK 801/ dizocilpine, remacemide), glycine site antagonists (GV 150526, ACEA 1021), polyamine site antagonists (eliprodil), growth factors (bFGF), Sodium channel blockers (fosphenytoin, 619C89), potassium channel opener (BMS 204352, cromakalim, levcromakalim, aprikalim, pinacidil, diazoxide, nicorandil, minoxidil), piracetam, adenosine transport inhibitor (propentofylline), gangliosides GM (non NMDA antagonist), presynaptic glutamate release inhibitors, clazosentan, desmoteplase, viprinex (ancrod), tenecteplase (TNKase; Metalyse), alteplase, cyclic nitrones, TWEAK (TNF-like weak inducer of apoptosis) receptor, thrombolytica treatments (urokinase, streptokinase, t-PA / tissue plasminogen activator or recombinant urokinase), anistreplase, riluzole, and disufenton sodium (NXY 059), candesartan, AX-200 (G-CSF, Fligrastim), caffeinol (caffeine+ethanol), enecadin, microplasmin, sonolysis + tPA, V-10153, HTUPA, solulin, piclozotan, S-0139, S-18986, AEOL-10150, AL-208, KN-38-7271, phridoxal 5-phosphate, Neu-2000KL, ONO-2231, PGX-100, RVX-208, SUN-N4057, SUN-N8075, TAT-NR2B9c, GLP-1-expressing stem cell therapy, Msc-1 (SA-4503, AGY-94806)), NH-02D, S-0139 259, tissue protective cytokines (Lu-AA24493), V10153 270 (BB-10153, TAPgen), combined use of statins and other cholesterol lowering drugs, erythropoietin, cerebrolysin and CDP-choline (cytidine-5'-diphosphocholine).

24. The method as claimed in claim 22, wherein said Western medicine is capable of activating potassium channel TREK-1.

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