# Adaptogenic and Anabolic Botanicals Combined to Restore Metabolic Homeostasis and Support Recovery

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### Allostasis and Cellular Metabolism

Allostasis, the adaptive response of our body's internal systems to seen and unseen factors, is the constant adjusting of our internal functions to maintain homeostasis and health. It is an ongoing process of achieving stability through change. With prolonged stress and when our stress load exceeds the body's ability to adapt (allostatic overload) and maintain homeostasis there is a decline in health as physiological function becomes impaired. When our physiology remains on constant alert due to overactivity of the arousal systems, the body is unable to fully recover or adapt. This erodes areas of constitutional weakness and leads to depletion of vital energy reserves with concomitant fatigue and possibly disease.

Biological aging and disease are associated with functional decline and with deficits at the cellular, tissue, organ and system levels of the body. Aging can be viewed as a process of shifting from a primarily anabolic metabolic balance to increasing levels of catabolic activity. This metabolic shift accounts for wear and tear, enzyme depletion, free radical expression, oxidative stress and other models of aging including expression of genetic errors.

Cellular metabolism, with its ongoing process of anabolic and catabolic activities, forms the dynamic basis of all physiological activity. Anabolic processes involve the synthesis of complex molecules and requires energy. Catabolic processes include the breakdown of nutrients and are associated with the manufacture and release of energy. Catabolic activity includes the breakdown of materials no longer needed by the cell that are subsequently removed from the cell.

Our lifestyle, aging and stress responses affect the balance of anabolic and catabolic activity. This influences our endocrine function, immunity, cell proliferation, bioenergetics and cell communication along with mood and behavior. Aging is associated with reduced anabolic hormone activity. The ratio of serum testosterone to cortisol is often used as an anabolic/catabolic index.

Cortisol and catecholamines mediate stress response and

adaptation in many physiological systems. While beneficial in the short-term, their prolonged activation results in pathological changes including immunosuppression, cardiovascular issues and metabolic imbalance. A prolonged stress response leads to impairment of cellular metabolism, dysfunction in the HPA (hypothalamic-pituitary-adrenal) axis and depletion of the adrenal glands. Allostatic overload also causes excessive activity of inflammatory cytokines and disruption of cellular metabolism.

The stress reaction is based on activation of the organism's energy supply by activating catabolic processes. The key trigger of this process is the stress-induced hyper-production and hyper-secretion of adrenal cortical glucocorticoid hormones. During prolonged stress, energy

## HPA axis dysfunction can lead to:

- dysregulation of catecholamines (fight or flight stress hormones)
- dysregulation of glucocorticoids from the adrenals
- dysregulation of cytokines (disrupted immune response)
- atrophy of nerve cells in the amygdala and hippocampus of the brain
- bone mineral loss
- high lipids
- abdominal obesity
- chronic fatigue
- mild depression and anxiety
- poor sleep patterns
- · altered cognitive performance
- decreased sexual behavior



metabolism at the cellular level is impaired along with the inflow of nutrients and outflow of waste materials through the cell walls. This, in turn, influences the balance of anabolic and catabolic metabolism resulting in a decrease in cellular energy and function. It also results in an accumulation of waste materials that interfere with cellular function. This leads to functional decline of the tissues, organs and systems that the cells comprise. Mitochondria are the energy production centers of the cell. Mitochondrial dysfunction is considered a leading cause of disease and bioenergetic decline.<sup>2</sup>

# Adaptogens Enhance Anabolic Metabolism

Adaptogenic botanicals enhance cellular energy processes, restore mitochondrial function and build adaptive energy reserve. They enhance cellular metabolism and provide an anabolic, anticatabolic effect to reduce the loss of lean muscle mass and optimize anabolic metabolism. They also enhance the regulation of bio-rhythms.<sup>3,4</sup>

By definition, adaptogenic botanicals normalize homeostasis, revitalize exhausted organ systems and improve resilience, vitality and resistance to adverse factors. They enhance appropriate immunological response and act in a nonspecific manner to strengthen physiological adaptation. Adaptogens support the body's natural processes and aid organ systems to work optimally, without side effects. A restorative agent is one that helps restore health, strength or consciousness, one that strengthens and invigorates the body, mind and spirit. Adaptogenic herbs are restorative and, at the same time, support the body's capacity for the ongoing process of adaptation that is referred to as allostasis.

Because they effectively enhance adaptive response and normalize neuroendocrine function, adaptogenic botanicals offer great potential for promoting wellness and longevity.

# Adaptogens:

- non-specific, normalizing action
- support physiological adaptation
- enhance the body's normal processes
- nourish vital life force
- promote optimal function
- · no side effects or toxicity
- normalize hypo- and hyper-conditions
- modulate neuroendocrine function

Adaptogenic herbs are ideally taken to support healthy function, enhance vitality, promote allostasis and help prevent disease. They are highly beneficial during the first stage of the stress response to support adaptation, a quick recovery and return to allostasis. They are also effective to facilitate recovery from depletion and exhaustion.

When multiple herbs are blended together to create harmonious formula, a synergistic effect is created that supports multiple systems and optimizes the healing response. It is often beneficial to use two or more formulas in tandem or in rotation to harmonize physiological systems, provide multiple actions and to address issues unique to the individual. Formulas may change as a person progresses on their healing path. Initially, when depletion is severe, the focus is on nourishing and restorative tonics. Once a person is stronger and can integrate more energy in their system, then dynamically stimulating herbs may be given.

# Adaptogenic and Anabolic Botanicals that Restore Metabolic Homeostasis and Support Recovery

Eleuthero (Eleutherococcus senticosis)

Eleuthero is the perhaps the most widely researched adaptogen. The unique qualities of this herb led the Russian scientist Dr. Nikolai Lazarev to coin the term "adaptogen" in the late 1940s. Later, Israel I. Brekhman, MD used the remarkable properties of Eleuthero to define the parameters of "adaptogenic" herbs.

The most striking biological property of Eleuthero is its ability to prevent or alleviate the stress response by normalizing function. It helps decrease the level of the alarm reaction and delay the onset of adrenal exhaustion (the third phase of Hans Selye's General Adaptation Syndrome). Eleuthero allows a more economical and efficient release of corticosteroids and adrenaline.<sup>5</sup>

Eleuthero is highly effective in improving the body's adaptive capacity to respond in a variety of situations from athletic performance to stress and illness. Eleuthero enhances mental acuity and work along with physical capacity and stamina.<sup>3,6-8</sup>



It enhances capacity for work and exercise with long-term benefits. Eleuthero improves oxygen uptake during exercise – enabling longer workouts, increased stamina and quicker recovery time especially in performance athletes.<sup>7,9</sup>

Eleuthero demonstrates anabolic activity<sup>3,9</sup> and stimulates protein synthesis in the pancreas, liver and adrenal cortex.<sup>3,10</sup> One of Eleuthero's bioactive compounds, Eleutheroside B, is the key compound that plays a significant role in the antistress and anabolic actions of Eleuthero.<sup>10-12</sup>

Eleuthero normalizes function more than any other adaptogenic agent.<sup>3,10</sup> It increases biological resistance, normalizes individual physiological responses and enhances vital energy systems. Eleuthero supports and optimizes endocrine and adrenal gland function when the body is challenged by stress. It is found to normalize blood sugar levels<sup>10,13-15</sup> and helps combat adverse effects of sleep deprivation<sup>16</sup>.

### Schisandra (Schisandra chinensis)

Both Schisandra seed and fruit extract are utilized in adaptogenic formulations, as the seed extract contains powerful adaptogenic compounds including higher amounts of lignin compounds called schisandrins. 6,17

Schisandra seed extract paired with the fruit extract offers the unique property of stimulating the central nervous system without causing over-excitation. This supports improved physical and mental capacity, motor coordination and efficiency. Schisandra seed is widely used for the treatment of stress-induced nervous system exhaustion and fatigue, insomnia, weakness, depression, forgetfulness, vision problems, diarrhea and chemical toxicity. Studies show that Schisandra seed extract enhances physical performance and facilitates recovery time. Section 18,20

The orange-red Schisandra berry has a long history of medicinal and food use in China, Japan, Korea, Tibet and Russia. Throughout time, hunters in the wilds of Siberia have used the dried berries chewed or prepared as a tea to provide energy, stave off exhaustion and improve night vision during long trips. Known as "Five Flavor Fruit" in Chinese medicine, it is considered a tonic that benefits all five energetic/organ systems according to Chinese medical principles, where each flavor relates to a specific energetic system. However, Schisandra was used particularly to nourish the lungs, support liver function and benefit the eyes.

In modern research, it is found to improve visual acuity, increase adaptation to darkness and widen the borders of the visual field.<sup>21</sup> In multiple studies, Schisandra is reported

to increase mitochondrial glutathione redox status<sup>22-24</sup>, which suggests a role in preventing oxidative stress.

# Rhodiola (Rhodiola rosea)

The active consituents unique to *Rhodiola rosea* and thought to be responsible for its adaptogenic qualities are two glycosides, particularly a group called "rosavins."<sup>25</sup> Other compounds in *R. rosea* include flavonoids, volatile essential oils and triterpenes. It also includes 17 amino acids along with vitamins, minerals and trace elements.<sup>26-29</sup>

Known as Golden or Artic Root, this perennial plant has red, pink or yellow flowers and grows prolifically in the colder north of Russia. Chinese emperors sent expeditions to Siberia to bring back the "golden roots" prized for medicinal use. Rhodiola is used in Tibetan medicine to promote circulation and prevent hypertension.

Rhodiola is valued for its ability to enhance mental and physical performance and stamina. It exerts profound protective effects on the nervous and endocrine systems. *R. rosea* extract is found to enhance immune function, improve resistance to stress, and help eliminate fatigue. It benefits concentration, focus, memory and learning capacity. Rhodiola demonstrates the ability to reduce stress-induced depression and decrease levels of anxiety. It harmonizes the neuroendocrine system and improves HPA axis response to stress.

Studies find that extracts of Rhodiola stimulate ATP (adenine triphosphate) and synthesis of creatine phosphate and glycogen in the muscles and liver along with anabolic activity in muscles such as protein synthesis. Thodiola is cardioprotective and helps normalize the heart rate after intense exercise. Some studies indicate it can normalize a system in conditions of stress and arrhythmia. Several human and animal studies found that *R. rosea* increases physical work capacity and dramatically shortens recovery time between rounds of high-intensity exercise.

Another study reported it enhances mental performance and helps reduce fatigue in physicians working night shifts.<sup>43</sup> Rhodiola is also reported to exert antioxidant activity, helping cells to withstand oxidative stress.<sup>44,45</sup>

## Panax ginseng

This highly-revered herb has been used as both food and medicine in Asian countries for over 4,000 years.

The genus name "Panax" is derived from the Greek word "pan", meaning all and "axos" (medicine or remedy) –

reflecting the root's reputation as a panacea. Because the root is shaped like a man it is believed to embody a human's three essences (body, mind and spirit) and to also contain the essence of the earth. It is highly esteemed in Oriental medicine as an herb of longevity that nourishes all the viscera and benefits the mind.

Fresh, unprocessed ginseng has a cooling, calming and quiescent effect and is known as White Panax Ginseng. When fresh ginseng is specially prepared and steamed with herbs, it becomes more warming in nature and is known as Red Panax Ginseng. Red Panax Ginseng is recognized for its ability to restore dynamic vigor and vitality to the body.

Ginseng is high in saponins, the most studied of which are the group of ginsenosides. Red Ginseng is said to be higher in ginsenosides due to the conversion of naturally-occurring compounds during the steaming process to ginsenosides. The outer skin, which is naturally high in ginsenosides, is retained for the making of Red Ginseng. *Panax ginseng* is also high in polysaccharides, peptide glycans and flavones.<sup>46</sup>

Ginseng is well-known for its ability to enhance vital energy and benefit physical performance and stamina. It also benefits mental cognition and concentration. Ginseng enhances recovery from exertion and muscle fatigue. 47-50 It has been used for thousands of years to help resolve serious illness, combat fatigue and promote longevity. It is beneficial for neurasthenia and mild depression.

Ginseng exerts modulating activity on the central nervous system. It facilitates the stress response, acts on the hypothalamus and modulates the activity of the adrenal cortex. It helps normalize glucocorticoid levels to enhance recovery time after stress. It modulates healthy blood glucose levels. 51-54

Ginseng is reported to be neuroprotective and enhance cognitive ability.<sup>55</sup> Ginsenosides from Ginseng have antioxidative properties and enhance the antioxidative defense system, including glutathione. It promotes the activity of the antioxidant enzymes GSH-Px (glutathione peroxidase) and SOD (superoxide dismutase).<sup>56</sup> At the same time, it helps decrease levels of proinflammatory cytokines,<sup>55,57</sup> while it modulates and enhances immune system response, NK (natural killer) cell activity and interferon production.<sup>58</sup>

Rhaponticum carthamoides

R. carthamoides (RC) grows in the pristine alpine and subalpine zones and alpine meadows of Southern Siberia. The root and underground stems of Rhaponticum are valued as powerful medicines in Siberian

folk medicine where RC has been used for centuries to treat fatigue, anemia and impotence. Noted for its ability to support recovery from disease, <sup>59</sup> it was also used to strengthen those suffering from weakness after illness or from overwork. <sup>60</sup>

This plant has been researched for over 30 years and acknowledged for its powerful adaptogenic benefits. It is noted for its ability to increase protein biosynthesis and to enhance physical and mental work capacity along with physical performance and endurance. It modulates immune function, exerts antioxidant activity and enhances cardiovascular functions. <sup>60,61</sup>

Rhaponticum is high in flavonoids and contains lignans, carotenoids, tannins, resins, vitamin C and glycosides. <sup>62-64</sup> It is especially studied due to its naturally-occurring steroids, particularly ecdysteroids which are anabolic. <sup>60,65</sup> The biological activity of RC is attributed to the phytoecdisone content, which influences metabolic processes. <sup>66</sup>

Rhaponticum extract (RE) increases protein anabolism to build lean muscle and reduces body fat.<sup>60,67</sup> This has led to its use by athletes to increase muscle mass and support strength and endurance.<sup>60</sup> RE improves the capacity for physical work, motor coordination and enhances athletic performance.<sup>67</sup>

Researchers find that RE helps prevent fatigue by increasing non-specific resistance and supporting mental and physical performance.<sup>68,69</sup> It also facilitates recovery after intensive activity.<sup>67,70</sup>

Rhaponticum normalizes the central nervous system and the cardiovascular system and improves vascular perfusion of the muscles and brain.<sup>71</sup> The flavonoids in RE are thought to enhance their ability to enhance vascular wall strength and to inhibit platelet aggregation.<sup>60</sup>

Rhaponticum powerfully improves adaptive capacity and normalizes function in many physiological systems. It is found to enhance sleep, appetite, mood and both physical and mental work performance under stressful conditions.<sup>71</sup>

Ashwagandha (Withania somnifera)

Ashwagandha, or Winter Cherry, is a woody shrub in the Solanaceae family that grows in diverse areas including Africa, India and the Mediterranean.

This powerful herb has been revered in Ayurvedic medicine for over five thousand years. Often called Indian Ginseng, it belongs to an elite class of Ayurvedic restorative, tonic herbs, known as Rasayana. The name Ashwagandha means "the smell of a horse," referring to the strong smell of the root. It also refers to the traditional belief that Ashwagandha root



confers the vigor, virility and strength of a stallion.

Active compounds in Ashwagandha root include alkaloids, steroidal lactones, saponins and iron.<sup>72</sup> Ashwagandha extract can prevent depletion of vitamin C and cortisol in subjects under stress. Its anti-stress and anabolic activity is considered similar to that of *Panax ginseng*.<sup>73</sup> It normalizes biological markers induced by stress including blood sugar, cortisol levels and adrenal function.<sup>74</sup> Ashwagandha is found to exert antioxidant and anti-inflammatory activity. One study found that an extract of Ashwagandha significantly suppresses production of pro-inflammatory cytokines in both normal individuals and in rheumatoid arthritis patients.<sup>72</sup>

Ashwagandha is known for its neurocognitive benefits including nervous system restoration. The Studies demonstrate that Ashwagandha benefits cognition and offers significant brain- and neuro-protective qualities. One study found 80% reduction in cell degeneration in the brain of stressed animals. Another discussed the ability of an isolate of Ashwagandha to positively influence regeneration of neurons and synapses in damaged neurons and neural circuits – vital components of the nervous system and brain. Ashwagandha may be protective of brain cells and support healthy brain function in degenerative brain conditions. It was found to increase acetylcholine receptor activity which may partially explain its ability to enhance cognition and memory.

Ashwagandha supports healthy anabolic activity and nourishes those in a weakened physical or mental condition. Chinese medicine reveres it as a Kidney Yang (metabolicallywarming) tonic to promote longevity and health. Modern studies find that Ashwagandha helps to normalize glucose levels, supports healthy aging and enhances male sexual function.<sup>83-85</sup>

Deer antler is the only mammalian organ that

# Pantocrine (Cervus elaphus)

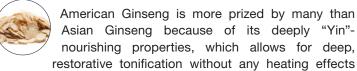
regenerates itself in an annual rhythm. Each spring, male deer cast off the hard antler from the previous year and the growth of a new antler begins. It starts as a soft, velvet material that regenerates at an amazing pace. It's not surprising that ancient cultures in the Far East were fascinated by this yearly miracle and intuited that deer antler may offer rejuvenative qualities to help slow the aging process. Deer antler velvet has an amazing nutrient profile including collagen, amino acids, essential fatty acids, enzymes, vitamins, minerals and trace minerals. It also contains high concentrations of somatodein-C, an important

A vast array of literature on deer antler exists in Asia. Natura uses a humanely-harvested extract of deer antlers known

as Pantocrine. Studies find Pantocrine improves athletic performance of healthy athletes and increases work capacity, strength and stamina. It helps enhance sexual health and supports healthy heart function.<sup>88</sup>

Pantocrine decreases the rate of muscle fatigue, enhances muscular strength and possesses anti-catabolic activity. It also increases red and white blood cell production and accelerates healing and recovery. Pantocrine has an anabolic/anti-catabolic effect in elderly animals, stimulating lean muscle, bone, cartilage and nerve growth. Pantocrine is anabolic – it promotes protein synthesis, building lean muscle and tissue.

# American Ginseng (Panax quinquefolius)



such as happens with "Yang" tonics (including Red and Korean Ginseng). American Ginseng is particularly known as a lung tonic.

The Eclectic physicians were a branch of American medicine in the latter half of the 19th and first half of the 20th centuries. They used botanical remedies extensively and valued American Ginseng as a tonic to support energy, stamina and endurance. It was considered a soothing nervine energy tonic beneficial for cerebral anemia, nervous dyspepsia and in mental exhaustion from overwork.<sup>95</sup>

Found to modulate the HPA axis, it increases endurance and reduces fatigue after exercise. Fa A large amount of research supports American Ginseng's traditional use as a nervine and tonic as it shows multiple beneficial neurocognitive effects. Extracts are found to be neurotrophic, neuro-regenerative and neuro-protective.

Constituents include ginsenosides (which are triterpene saponin glycosides) in ratios different than those found in Asian Ginsengs. American Ginseng also contains polysaccharides, proteins and proteoglycans.<sup>96</sup>

Studies demonstrate a positive influence on the cardiovascular system with antioxidant effects. Extracts are found to exert both antioxidant and anti-inflammatory activity. American Ginseng benefits the immune system and promotes immune cytokine production. <sup>97-101</sup> It demonstrates cytoprotective and hepatoprotective capacity. <sup>97</sup> Extracts of American Ginseng are found to normalize blood sugar <sup>96</sup> and to protect against renal damage in diabetics. <sup>102</sup> Studies find it reduces inflammation in the colon and may exert a beneficial influence on the colon microbiome. <sup>103</sup>

growth hormone.86,87

### Cordyceps (Cordyceps sinensis)

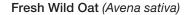
Known in China as "winter worm summer grass" or the "caterpillar mushroom," Cordyceps is a parasite fungus that traditionally grows on the larvae

of caterpillars and other hosts that provide a growth medium for the mycelia. Today, most Cordyceps is made from cultured mycelia. The Cordyceps CS-4 strain cultured on organic brown rice offers superior quality and therapeutic activity. The CS-4 strain is the most widely-researched and highly-regarded.

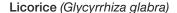
Li Chih Shen, a renowned sixteenth century herbalist, praised Cordyceps for its ability to invigorate and tone the entire body. Chinese herbalists revered it as a respiratory tonic. At the same time it deeply nourishes the deep reserve (Kidney) energy that supports healthy lung function and overall vitality and stamina. Cordyceps contains polysaccharides that contribute to its immunomodulating influence.<sup>104</sup>

Cordyceps first gained international attention when it was discovered that Chinese Olympic athletes included Cordyceps as part of their daily herbal training formula. Studies confirm that Cordyceps increases endurance, vigor and athletic training and performance. 105,106 Recognized for its liver- and kidney-protective qualities, Cordyceps is reported to be especially beneficial for those with chronic kidney disease. 107-110

Cordyceps extract is found to inhibit brain aging, modulate endocrine function,<sup>111</sup> and to restore sexual function with the ability to replenish sperm and support healthy testosterone levels.<sup>112</sup>



Avena is a classic nervine tonic that builds energy and reduces stress. It is traditionally recognized as a nourishing, restorative tonic for the nervous system.<sup>113-114</sup> The milky seed of wild oats nourishes and harmonizes the sympathetic and parasympathetic nervous systems. It is one of the best herbs to restore vital energy especially in cases of adrenal exhaustion. The Eclectic physician Dr. Finley Ellingwood considered it a great remedy to alleviate nervous exhaustion due to stress, overwork and nervous anxiety.<sup>113</sup> Modern research reports that oats exert a wide spectrum of activity including antioxidant, anti-inflammatory and immunemodulating.<sup>114</sup> *Avena sativa* is a rich source of protein, minerals (including calcium and magnesium), flavones, saponins, sterols and tocols.<sup>113-115</sup>



Licorice acts as a synergist because it moderates and harmonizes the characteristics of other botanicals in a formula. It is known as a special herb in Chinese medicine that carries the herbs in a formula throughout the body. Licorice is known as a demulcent (soothing to the mucus membranes of the body) and is noted for its anti-inflammatory, immune-modulating influence. 116-118

Licorice extract is shown to increase immune function including production of interferon and NK (natural killer) cells. Multiple flavonoids have been isolated from licorice, many of which show broad-spectrum antibacterial effects. Licorice is hepatoprotective, enhances adrenal function and supports the stress response through multiple pathways. Licorice is hepatoprotective.

For more information on any of the ingredients listed here, including extensive research or individual monographs compiled by Donnie Yance, please email info@naturaedu. com.



### References

- McEwen, BS PhD. Allostasis and Allostatic Load: Implications for Neuropsychopharmacology. Neuropsychopharmacology 2000 Vol 22, No 2. Elsevier Science Inc.
- 2. Selye H. Nature. 1936:138: 32.
- Blokhin BN The Influence of Eleutherococcus Root and Leaf Extract on Human Work Capacity under Static and Dynamic Workloads. ed. Eleutherococcus and Other Adaptogens from Far East Plants. 1966. Vladivostok: Siberian Department of the Academy of Sciences of the USSR.
- Wahlstrom M. Adaptogens: Nature's Key to Well-Being. 1987. Goteborg, Skandinavisk Bok.
- Brekhman II. Eleutherococcus: 20 Years of research and clinical application. Hamburg 1980. 1st International Symposium on Eleutherococcus.
- Wagner H, Norr H, Winterhoff H. Plant adaptogens. Phytomedicine 1994:163-76.
- Asano K, Takakhasi T, Kugo K, Kuboyama M. The Influence of Eleutherococcus on Muscle Work Capacity in Humans. eds. New Data on Eleutherococcus. Moscow. 1986, 1984. 2nd International Symposium on Eleutherococcus.
- Asano K, Takahashi T, Miyashita M, et al. Effect of Eleutherococcus senticosus extract on human working capacity. Planta Medica: 3(1986): 175-177.
- Kelly GS. Sports Nutrition: A review of selected nutritional supplements for endurance athletes. Alternative Medicine Review. 1997: 2(4): 282-295.
- Todorov IN, Zaikov GE, Degterev IA. Bioactive compounds: biotransformation of biological action, mechanism of antistress and anabolic action of Eleutherococcus. 1993. Commack, New York. Nova Science Publishers, Inc.
- 11. Wagner H, et al. *Die DC und HPLC analyze der Eleutherococcus*. Planta Medica. Vol 44 (1982):193-9.
- 12. Farnsworth N, Walter D and Sterkoff L. Use of Eleutherococcus in the USA: problems, prospects and literature update. New data on Eleutherococcus. Proceedings of the II International Symposium on Eleutherococcus. (Moscow 1984) Vladivostok. 1986:47-52.
- Brekhman II, Kirillov OL. Effect of Eleutherococcus on alarm-phase of stress. Life Sciences: 1969: 8(3)113-121.
- Kupin VJ. Eleutherococcus and other Biological Active Modifiers in Oncology. 1984. Moscow. Medexport.
- Hikino H, Takahashi M, Otake K, Konno C. Isolation and hypoglycemic activity of Eleutherans A, B, C, D, E, F and G: glycans of Eleutherococcus senticosus roots. Journal of Natural Products. 1986: 49(2):293-7.
- Provalova NV, Skurikhin EG, Pershina OV, et al. Mechanisms underling the effects of adaptogens on erythropoiesis during paradoxical sleep deprivation. Bulletin of Experimental Biology and Medicine. 2002: 133(5): 428-432.
- Lupandidn AV, Lapaev II. Schizandra. Khabarovsk book publisher. Moscow. USSR. 1981:125.
- 18. Weiner M. Schizandra. In book: Herbs that heal: Prescription for Herbal Healing. 1996:292-293.
- 19. Lebedev AA. Appraisal of the stimulative action of Schizandra

- Chinensis. Materials for the study of the stimulative and tonic actions of Ginseng and Schizandra Chinensis. Moscow, Academy of Science of USSR. 1955:182-185.
- Ahumada F, et al. Studies on the effect of Schizandra Chinensis extract on horses submitted to Exercise and Maximum Effort. Phytotheraphy. 1989: 3(5):175-179.
- Sinovich VA, Akhmerova, ZB. The influence of Schizandra on normal and pathological visual functions- Materials for study of Ginseng and Schizandra. Issue 3. Leningrad. 1958:177-180.
- 22. Chiu PY, Mak DH, Poon MK, Ko KM. In vivo antioxidant action of a lignan-enriched extract of Schisandra fruit and an anthraquinone-containing extract of Polygonum root in comparison with schisandrin B and emodin. Planta Med. 2002 Nov 68(11):951-6. Department of Biochemistry. Hong Kong University of Science and Technology. Clear Water Bay, Hong Kong, China.
- 23. Ip SP, Yiu HY, Ko KM. Differential effect of schisandrin B and dimethyl diphenyl bicarboxylate (DDB) on hepatic mitochondrial glutathione redox status in carbon tetrachloride intoxicated mice. Mol Cell Biochem 2000 Feb:205(1-2):111-4. Department of Biochemistry. The Hong Kong University of Science and Technology. Clear Water Bay, PR China.
- Ram VJ. Herbal preparations as a source of hepatoprotective agents.
  Drug News Perspect. 2001 Aug:14(6):353-63. Medicinal Chemistry Division, Central Drug Research Institute. Lucknow, India.
- 25. Kelly, G, ND. Nutritional and Botanical Interventions of Assist with the Adaptation to Stress. Altern Med Rev 1999:Vol 4, Number 4:255.
- Brown RP, Gerbarg PL, Ramazanov Z. Rhodiola rosea. A phytomedical overview. Journal of the American Botanical Council 2002. 56:40-52.
- 27. Rohloff J. Volatiles from rhizomes of Rhodiola rosea L. Phytochemistry 2002: 59(6) 655-661.
- Ganzera M, Yayla Y, Khan IA. Analysis of the marker compounds of Rhodiola rosea L. (golden root) by reversed phase high performance liquid chromatography. Chemical & Pharmaceutical Bulletin. 2001: 49(4) 465-467.
- Yin G, Guo J, Wang G, et al. The nutritional value and longevity effects of Rhodiola on animals. Acta Nutr Sin 1992: 14(1) 98-102.
- Fintelmann V, Gruenwald J. Efficacy and tolerability of a Rhodiola rosea extract in adults with physical and cognitive deficiencies. Advances in Therapy 2007: 24(4) 929-939.
- Saratikov AS, Krasnov EA. Chapter III: Stimulative properties of Rhodiola rosea. eds. Rhodiola rosea is a valuable medicinal plant. Tomsk State University. 1987:69-90.
- Kelly GS. Nutritional and botanical interventions to assist with the adaptation to stress. Alternative Medicine Review. 1999: 4(4) 249-265.
- Darbinyan V, Kteyan A, Panossian A, et al. Rhodiola rosea in stress induced fatigue – a double blind cross-over study of a standardized extract SHR-5 with a repeated low-dose regimen on the mental performance of healthy physicians during night duty. Phytomedicine 2000: 7(5) 365-371.
- 34. Brichenko VS, Kupriyanova IE, Skorokhova TF. The use of herbal adaptogens with tricyclic antidepressants in patients with psychogenic depression. In: Saratikov AS, ed. Modern problems of pharmacology and search for new medicines. Tomsk State University Press. 1986:



- 58-60.
- Novikov MG, Adamchuk VD. Experience in the organization of medical-preventive work at an industrial enterprise. Sovetskoe zdravookhranenie / Ministerstvo zdravookhraneniia SSSR 1960: 19(3) 35-41.
- Maslova LV, Kondrat'ev BI, Maslov LN, Lishmanov I. The cardioprotective and antiadrenergic activity of an extract of Rhodiola rosea in stress. 1994: 57(6) 61-63.
- Maimeskulova LA, Maslov LN. Anti-arrhythmic effect of phytoadaptogens. Eksperimental'naia i klinicheskaia farmakologiia 2000: 63(4): 29-31.
- Maslov LN, Lishmanov I, Maimeskulova LA, Krasnov EA. Mechanism of the anti-arrhythmic effect of Rhodiola rosea extract. Biulleten' eksperimental'noi` biologii i meditsiny 1998: 125(4) 424-426.
- Maslov LN, Lishmanov I. Cardioprotective and antiarrhythmic properties of Rhodiolae roseae. Eksperimental'naia i klinicheskaia farmakologiia 2007: 70(5) 59-67.
- 40. Brown, R MD, Gerbarg, P MD, Ramazanov, Z, PhD DS. *Rhodiola rosea, A Phytomedical Overview*. HerbalGram 2002. 56:40-52.
- Saratikov, AS, Krasnov, EA. Clinical Studies of Rhodiola. Rhodiola Rosea is a Valuable Medicinal Plant. Tomsk: Medical Institute, Russia 1987:216
- Saratikov, AS, Krasnov, EA. The Adaptogenic Attributes of Rhodiola. Rhodiola Rosea is a Valuable Medicinal Plant. Tomsk: Medical Institute, 1987:194.
- 43. Darbinyan V, Kteyan A, Panossian A, et al. *Rhodiola rosea in stress induced fatigue--a double blind cross-over study of a standardized extract SHR-5 with a repeated low-dose regimen on the mental performance of healthy physicians during night duty.* Phytomedicine. 2000 Oct 7(5):365-71 (ISSN: 0944-7113) Department of Neurology, Armenian State Medical University, Yerevan.
- 44. Ip SP, Che CT, Leung PS. Association of free radicals and the tissue renin-angiotensin system: prospective effects of Rhodiola, a genus of Chinese herb, on hypoxia-induced pancreatic injury. JOP 2001 Jan 2(1):16-25 (ISSN: 1590-8577 School of Chinese Medicine, Department of Physiology, The Chinese University of Hong Kong, Shatin, NT. Hong Kong.
- Zhang Z, Liu J, Shang X, et al. The effect of Rhodiola capsules on oxygen consumption of myocardium and coronary artery blood flow in dogs. Zhongguo Zhong Yao Za Zhi 1998 Feb:23(2):104-6. (ISSN: 1001-5302) Institute of Information on TCM, China Academy of Traditional Chinese Medicine, Beijing 100700.
- 46. Dharmananda, S. PhD. *The Nature of Ginseng: From Traditional Use to Modern Research*. This article first appeared in Herbalgram 54: The Journal of the American Botanical Council. Web Posting Date: Sept 2002. Institute for Traditional Medicine. Portland, Oregon. http://www.itmonline.org/arts/ginsengnature.htm
- Weider, B. Selected herbals and human exercise performance. Am J Clin Nutr 2000 Aug 72(2 Suppl):624S-36S (ISSN: 0002-9165). Weider Nutritional International. Salt Lake City, UT.
- Forgo I, Kirchdorfer AM. The Effects of Different Ginsenoside Concentrations on Physical Work Capacity. Notabene Medici 1982. 12(9):727.
- Jung HL, Kwak HE, Kim SS, et al. Effects of Panax ginseng supplementation on muscle damage and inflammation after uphill treadmill running in humans. Am J Chin Med. 2011:39(3):441-50.
- 50. Lee HM, Lee OH, Kim KJ, Lee BY. Ginsenoside Rg1 Promotes

- Glucose Uptake Through Activated AMPK Pathway in Insulin-resistant Muscle Cells. Phytother Res. 2011 Dec 15. doi: 10.1002/ptr.3686.
- Reay JL1, Kennedy DO, Scholey AB. Effects of Panax ginseng, consumed with and without glucose, on blood glucose levels and cognitive performance during sustained 'mentally demanding' tasks. J Psychopharmacol. 2006 Nov:20(6):771-81. Epub 2006 Jan 9.
- Wagner H, Bladt S. Plant drug analysis: a thin layer chromatography atlas, 2nd ed. Spinger-Verlag, Berlin, 1996:307.
- Chang HM, But PP. Pharmacology and applications of Chinese materia medica, Vol. 1. World Scientific, Singapore. 1986:17-31.
- Kee CH. The Pharmacology of Chinese Herbs. Second Ed. CRC Press, Boca Raton, FL. 1999:19-44.
- Zhu J, Mu X, Zeng J, Xu C, Liu J, et al. Ginsenoside Rg1 Prevents Cognitive Impairment and Hippocampus Senescence in a Rat Model of DGalactose-Induced Aging. 2014. PLoS ONE 9(6): e101291. doi:10.1371/journal.pone.0101291
- Shen LH, Zhang JT. Ginsenoside Rg1 promotes proliferation of hippocampal progenitor cells. Neurological Research 2004. 26(4):422-428
- Wu SF, Sui DY, Yu XF, et al. Antimyocardial ischemic effects of Panax quinquefolium 20 s-protopanaxdiol saponins(PQDS) and its mechanism. Chinese Pharmaceutical Journal 2002. 37(2): 100-103.
- 58. Mills S, Bone K. *Principles & Practice of Phytotherapy*. Churchill Livingston. Part Three: Ginseng.
- Saratikov AS. On the stimulative action of Siberian Leuzea carthamoides. New Medicinal Plants of Siberia, Their Medicinal Preparations and Use. 1949: 3: 167-190.
- Kokoskaa L, Janovskab D. Chemistry and pharmacology of Rhaponticum carthamoides: A review. Phytochemistry. Volume 70. Issue 7. May 2009:842-855.
- 61. Yakunina GD, Krasnov EA. *Leuzea carthamoides as a prospective resource of new remedies*. Herbal remedies in traditional and folk medicine. Ulan Ude 1987; 155-156.
- 62. Koleckar V, Brojerova E, Rehakova Z, et al. *In vitro antiplatelet activity of flavonoids from Leuzea carthamoides*. Drug and Chemical Toxicology. 2008. 31(1): 27-35.
- 63. Lamer-Zarawska E, Serafinowicz Gasiorowwski K, Brocos B. Immunomodulatory activity of polysaccharide-rich fraction from Rhaponicum carthamoides leaves. Fitoterapia 1996; 4: 371-372.
- 64. Szendrei K, Varga E. *Thiophene acetylenes from Leuzea Roots*. Phytochemistry 1984: 23(4): 901-902.
- 65. Syrov VN, Kurmukov AG. *Anabolic activity of phytoecdisterone, isolated from Rhaponticum carthamoides*. Farmakol Toksikol. 1976: 39(6):690-3.
- Krasnov EA. Saratikov AS. Yakunina GD. Incosterone and Ecdisterone from Rhaponticum Carthamoides. Chemistry of Natural Compounds. 1976. Issue 4:550.
- 67. Lupandin, AV. Adaptation and Rehabilitation in Sports. Khabarovsk: Institute of Physical Culture, 1991.
- 68. Gorovits MB, Zatsny IL, Abubakirov NK. Ecdysones in the world of plants. Plant's Resources 1974: 10(2): 261-274.
- Gorovits MB. Research of plant steroids and synthesis of physiological active substances from it. Abstract of Dissertation. 1977 Doctor's of Science degree in pharmacology.
- 70. Sololov S, Ya VD, Monogaroc VL. Sobolev, et al. The Influence of the Saponins of the Aralia Mandshurica on the Processes of Recovery



- on Athletes after Intensive Physical Workloads. Biologically Active Substances of Flora and Fauna of the Far East and Pacific Ocean Vladivostok. 1971. 113-114.
- Gerasyuta, MA, Koval TN. The Experience of Prolonged Use of Leuzea Carhamoides Extract for the Purposes of Preservation and Increase of Mental And Physical Work Capacity. In: Proceedings of the First International Symposium on Eleuthercoccus (Hamburg, 1980) Vladivostok: Far East Scientific Center of the Academy of Science of the Suur. 1981: 135.
- 72. Atal, CK, Gupta OP, Raghunathan K, Dhar KL. *Pharmacognosy and Phytochemistry of Withania Somnifera*. Central Council for Research in Indian Medicine and Homeopathy, New Delhi. 1975.
- 73. Singh D, Aggarwal A, Maurya R, Naik S. Withania somnifera inhibits NF-kappaB and AP-1 transcription factors in human peripheral blood and synovial fluid mononuclear cells. Phytother Res. 2007 Jun 11.
- 74. Grandhi A, Mujumdar AM, Patwardlhan B. *A comparative* pharmacological investigation of Ashwagandha and Ginseng. J Ethnopharmacol 1994. 44:131-135.
- 75. Archana R, Namasivayam A. *Antistressor effect of Withania* somnifera. J Ethnopharmacol. 1999. 64:91-93.
- Jain S, Shukla SD, Sharma K, Bhatnagar M. Neuroprotective effects of Withania somnifera Dunn. in hippocampal sub-regions of female albino rat. Phytother Res 2001:15(6):544-8.
- 77. Bhattacharya SK, et al. *Anti-Stress activity of Sitoindosides VII and VIII, New Acylsteryglucosides from Withania somnifera*. Phytotherapy Research 1987:1(1):32-37.
- 78. Archana R, Namasivayam A. *Antistressor effect of Withania somnifera*. J Ethnopharmacol 1999: Jan 64(1):91-3.
- 79. Jain S, Shukla SD, Sharma K, Bhatnagar M. Neuroprotective effects of Withania somnifera Dunn. in hippocampal sub-regions of female albino rat. Phytother Res 2001: 15(6):544-8.
- Kuboyama T, Tohda C, Komatsu K. Neuritic regeneration and synaptic reconstruction induced by withanolide A. Br J Pharmacol 2005 Feb 14.
- Tohda C, Kuboyama T, Komatsu K, Vanella A. *Indian medicinal plants as antiradicals and DNA cleavage protectors*. Phytomedicine 2001 Mar 8 (2):125-32.
- 82. Jayaprakasam B, Padmanabhan K, Nair MG. Withanamides in Withania somnifera fruit protect PC-12 cells from beta-amyloid responsible for Alzheimer's disease. Phytother Res. 2009 Dec 2.
- 83. Bhattacharya SK, Bhattacharya A, Sairam K, Ghosal S. *Anxiolytic-antidepressant activity of Withania somnifera glycowithanolides: an experimental study*. Phytomedicine Dec 2000: 7(6):463-9.
- 84. Bhattacharya SK, Muruganandam AV. Adaptogenic activity of Withania somnifera: an experimental study using a rat model of chronic stress. Pharmacol Biochem Behav. 2003 Jun:75(3):547-55. Department of Pharmacology, Postgraduate Institute of Basic Medical Sciences, Calcutta University. 244 B Acharya JC Bose Road, 700 020, Calcutta, India
- 85. Mishra LC, Singh BB, Dagenais S. Scientific basis for the therapeutic use of Withania somnifera (Ashwagandha): a review. Altern Med Rev 2000 Aug;5(4):334-46.
- 86. Academy of Sciences of the USSR Far East Branch. *The Soviet Far East: The USSR's Medicinal Treasury*. 1990. Address of the Presidium: 50 Leninskaya, Vladivostok 690600, USSR.
- 87. Duarte A. (2000). Velvet Deer Antler: The Ultimate Anti-Aging

- Supplement. Grass Valley, CA: Nutri Tapes and Publications, Inc.
- Pavlenko, SM. Pantocrin: A Publication of Articles on Studies of Curative Properties of Pantocrin. 1988. Moscow, USSR: V/O Medexport.
- Chinese Herbal Medicine Materia Medica. translated by D Bensky & A Gamble. revised edition 1993. Eastland Press: 336-7.
- Dobriyakov U. Antistress action of the remedy from Antlers. In Adaptation and Adaptogens. Vladivostok, 1977:132-134.
- Pantokrin, Pharmacological Article. 1995. Ministy of Healthcare of Russia. Moscow. Official Document.
- Wang B. Advances in the research of chemistry, pharmacology and clinical application of pilose antler. Proceedings of the '96 International Symposium on Deer Science and Deer Products. 1996. Changchun, China. 14-32.
- 93. Wang BX, Zhao XH, Qi SB et al. Stimulating effect of deer antler extract on protein synthesis in senescence-accelerated mice in vivo. Chemistry and Pharmacology Bulletin. 1988: 36(7): 2593-2598.
- Ryashchenko (Editor) Russian book on Pantocrine, translation arranged by R. Archer, Properties of New Zealand deer velvet, 1983.
- 95. Cook, W MD. The Physiomedical Dispensatory. Professor of Botany, Therapeutics, and Materia Medica in the Physio-Medical Institute. 1869: 399
- Felter, HW MD, Lloyd, JU PhD. King's American Dispensatory 1898
  Eighteenth Edition. Third Revision. In Two Volumes.
- Upton, R editor. American Herbal Pharmacopeia and Therapeutic Compendium: American Ginseng Root (Panax quinquefolius L.) Standards of Analysis, Quality Control, and Therapeutics. 2012. American Herbal Pharmacopoeia. www.herbal-ahp.org
- Assinewe VA, Amason JT, Aubry A, et al. Extractable polysaccharides of Panax quinquefolius L. (North American ginseng) root stimulate TNFalpha production by alveolar macrophages. Phytomedicine. 2002: 9(5) 398-404.
- 99. Fu Y, Ji LL. Chronic ginseng consumption attenuates age-associated oxidative stress in rats. The Journal of Nutrition 2003: 133(11) 3603-
- 100. Wang M, Guilbert LJ, Ling L, et al. Immuno-modulating activity of CVT-E002, a proprietary extract from North American ginseng (Panax quinquefolium). The Journal of Pharmacy and Pharmacology 2001: 53(11) 1515-1523.
- 101. Wang M, Guilbert LJ, Li J, et al. A proprietary extract from North American ginseng (Panax quinquefolium) enhances IL-2 and IFNgamma productions in murine spleen cells induced by Con-A. International Journal of Immunopharmacology 2004: 4(2) 311-315.
- 102. Kim HY, Kang KS, Yamabe N, Nagai R, Yokozawa T. Protective effect of heat-processed American ginseng against diabetic renal damage in rats. Journal of Agricultural and Food Chemistry 2007: 55(21) 8491-8497.
- 103. Dougherty et al. American ginseng suppresses Western diet-promoted tumorigenesis in model of inflammation-associated colon cancer: role of EGFR. BMC Complementary and Alternative Medicine 2011.11:111 http://www.biomedcentral.com/1472-6882/11/111
- 104. Zhu JS, Halpern GM, Jones K. The scientific rediscovery of an ancient Chinese herbal medicine: Cordyceps sinensis: part I. J Altern Complement Med 1998 Fall;4(3):289-303 (ISSN: 1075-5535) Department of Pediatrics, Stanford University School of Medicine, California, USA.



- 105. Koh JH, Kim KM, Kim JM, et al. Antifatigue and Antistress Effect of the Hot-Water Fraction from Mycelia of Cordyceps sinensis. Biol Pharm Bull. 2003 May:26(5):691-4.
- 106. Dai G, Bao T, Xu C, et al. CordyMax Cs-4 improves steady-state bioenergy status in mouse liver. J Altern Complement Med 2001 Jun 7(3):231-40. [medline]
- 107. Zhang HW, Lin ZX, Tung YS, et al. Cordyceps sinensis (a traditional Chinese medicine) for treating chronic kidney disease. Cochrane Database Syst Rev. 2014 Dec 18:12:CD008353. 36, 37.
- Shen LM, Chen YP. Treatment of 18 cases of chronic nephritis mainly with cultivated cordyceps. Liaoning J Trad Chin Med. 1985: 9:32-33.
- Chen YP, Liu WZ, Shen LM, Xu SN. Clinical effects of natural cordyceps and cultured mycelia of Cordyceps sinensis in kidney failure. Chin Trad Herbal Drugs. 1986: 17:256-258.
- 110. Kuo YC, Chen CF, Chen WP, et al. Inhibition of activated human mesangial cell proliferation by the natural product of Cordyceps sinensis (H1-A): an implication for treatment of IgA mesangial nephropathy. J Lab Clin Med 1999 Jan:133(1):55-63 (ISSN: 0022-2143) Lin CY, Ku FM. Department of Pediatrics and Medical Research. Veterans General Hospital-Taipei, Taiwan, Republic of China.
- 111. Yamaguchi Y, Kagota S, Nakamura K. Antioxidant activity of the extracts from fruiting bodies of cultured Cordyceps sinensis. Phytother Res 2000 Dec:14(8):647-9 (ISSN: 0951-418X). Department of Pharmacology, Faculty of Pharmaceutical Sciences, Mukogawa Women's University, Nishinomiya 663-8179, Japan.
- 112. Huang BM, Hsu CC, Tsai SJ. Effects of Cordyceps sinensis on testosterone production in normal mouse Leydig cells. Life Sci 2001 Oct 19: 69(22):2593-602 (ISSN: 0024-3205); Sheu CC; Leu SF Department of Cell Biology and Anatomy, College of Medicine, National Cheng Kung University, Tainan, Taiwan, Republic of China.
- 113. Ellingwood, F MD. *American Materia Medica, Therapeutics*. Eclectic Medical Publications 1919:204-6.
- 114. Singh R1, De S, Belkheir A. Avena sativa (Oat), a potential neutraceutical and therapeutic agent: an overview. Crit Rev Food Sci Nutr. 2013 53(2):126-44. doi: 10.1080/10408398.2010.526725.
- 115. Chu, YF1, Wise, ML, Gulvady AA, et al. In vitro antioxidant capacity and anti-inflammatory activity of seven common oats. Food Chem. 2013 Aug 15:139(1-4):426-31. doi: 10.1016/j.foodchem.2013.01.104. Epub 2013 Feb 10.
- 116. Herold A, Cremer L, Calugaru A, et al. Hydroalcoholic plant extracts with anti-inflammatory activity. Roum Arch Microbiol Immunol. 2003 Jan-Jun 62(1-2):117-29. National Institute of Research-Development for Microbiology and Immunology Cantacuzino, Bucharest, Romania.
- 117. Kwon HM, Choi YJ, Choi JS, et al. Blockade of Cytokine-Induced Endothelial Cell Adhesion Molecule Expression by Licorice Isoliquiritigenin Through NF-{kappa}B Signal Disruption. Exp Biol Med (Maywood). 2007 Feb 232(2):235-45.
- 118. Khan R1, Khan AQ, Lateef A. Glycyrrhizic acid suppresses the development of precancerous lesions via regulating the hyperproliferation, inflammation, angiogenesis and apoptosis in the colon of Wistar rats, PLoS One. 2013. 8(2):e56020. doi: 10.1371/ journal.pone.0056020. Epub 2013 Feb 14.
- 119. Shibata S. A drug over the millennia: pharmacognosy, chemistry, and pharmacology of licorice. Yakugaku Zasshi 2000 Oct;120(10):849-62 (ISSN: 0031-6903) Shibata Laboratory of Natural Medicinal Materials, C/o Minophagen Pharmaceutical Co., Ltd., Tokyo, Japan. Actual study- http://www.ncbi.nlm.nih.gov/pubmed/11082698

- 120. Xing GX, Li N, Wang T, Yao MY. Advances in studies on flavonoids of licorice, Zhongguo Zhong Yao Za Zhi. 2003 Jul 28(7):593-7. Review. Chinese.
- 121. Hatano T, Aga Y, Shintani Y, Ito H, Okuda T. Minor flavonoids from licorice. Phytochemistry 2000 Dec:55(8):959-63 (ISSN: 0031-9422) Yoshida T Faulty of Pharmaceutical Sciences, Okayama University, Tsushima, Japan.
- 122. Fukai T, Marumo A, Kaitou K; et al. Antimicrobial activity of licorice flavonoids against methicillin-resistant Staphylococcus aureus. Fitoterapia 2002 Oct 73(6):536 (ISSN: 0367-326X) Nomura T School of Pharmaceutical Sciences, Toho University, 2-2-1 Miyama, Funabashi, 274-8510, Chiba, Japan.
- 123. Song NR, Kim JE, Park JS, et al. Licochalcone A, a Polyphenol Present in Licorice, Suppresses UV-Induced COX-2 Expression by Targeting PI3K, MEK1, and B-Raf. Int J Mol Sci. 2015 Feb 20:16(3):4453-70. doi: 10.3390/ijms16034453.
- 124. Tanaka Y, Kikuzaki H, Fukuda S. *Antibacterial compounds of licorice against upper airway respiratory tract pathogens*. J Nutr Sci Vitaminol (Tokyo) 2001 Jun 47(3):270-3. (ISSN: 0301-4800) Nakatani N R & D Laboratory, Taiyo Corporation, Higashiyodogawa, Osaka, Japan.
- 125. Jeong HG, You HJ, Park SJ, et al. Hepatoprotective effects of 18betaglycyrrhetinic acid on carbon tetrachloride-induced liver injury: inhibition of cytochrome P450 2E1 expression. Pharmacol Res. 2002 Sep 46(3):221-7.
- 126. Bean P. The use of alternative medicine in the treatment of hepatitis C. Am Clin Lab. 2002 May 21(4):19-21. Review.
- 127. Lin G, Nnane IP, Cheng TY. The effects of pretreatment with glycyrrhizin and glycyrrhetinic acid on the retrorsine-induced hepatotoxicity in rats. Toxicon. 1999 Sep 37(9):1259-70.
- 128. Heilmann P, Heide J, Hundertmark S, Schoneshofer M. Administration of glycyrrhetinic acid: significant correlation between serum levels and the cortisol/cortisone-ratio in serum and urine. Exp Clin Endocrinol Diabetes. 1999. 107(6):370-8.
- 129. Al-Qarawi AA, Abdel-Rahman HA, Ali BH, El Mougy SA. *Liquorice* (*Glycyrrhiza glabra*) and the adrenal-kidney-pituitary axis in rats. Food Chem Toxicol. 2002 Oct 40(10):1525-7.
- Armanini D, Fiore C, Mattarello MJ, et al. History of the endocrine effects of licorice. Exp Clin Endocrinol Diabetes. 2002 Sep 110(6):257-61.
- 131. Stewart PM. Cortisol as a mineralocorticoid in human disease. J Steroid Biochem Mol Biol. 1999 Apr-Jun 69(1-6):403-8. Department of Medicine, University of Birmingham, Queen Elizabeth Hospital, Edgbaston, England, UK.

