

Effects of Adaptogens on the Central Nervous System and the Molecular Mechanisms Associated with Their Stress—Protective Activity

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Abstract

Adaptogens were initially defined as substances that enhance the “state of non-specific resistance” in stress, a physiological condition that is linked with various disorders of the neuroendocrine-immune system. Studies on animals and isolated neuronal cells have revealed that adaptogens exhibit neuroprotective, anti-fatigue, antidepressive, anxiolytic, nootropic and CNS stimulating activity. In addition, a number of clinical trials demonstrate that adaptogens exert an anti-fatigue effect that increases mental work capacity against a background of stress and fatigue, particularly in tolerance to mental exhaustion and enhanced attention. Indeed, recent pharmacological studies of a number of adaptogens have provided a rationale for these effects also at the molecular level. It was discovered that the stress—protective activity of adaptogens was associated with regulation of homeostasis via several mechanisms of action, which was linked with the hypothalamic-pituitary-adrenal axis and the regulation of key mediators of stress response, such as molecular chaperons (e.g., HSP70), stress-activated c-Jun N-terminal protein kinase 1 (JNK1), Forkhead box O (FOXO) transcription factor DAF-16, cortisol and nitric oxide.

1. Introduction

The idea that a pill could improve mental and physical performance in healthy people was devised during World War II with various stimulants given to pilots and members of submarine crews [1,2,3,4,5,6]. For instance, the first studies on the stimulating and tonic effects of *Schisandra chinensis* were published in Soviet Union WWII military journals [5,6]. The Russian interest in *S. chinensis* arises from ethnopharmacological investigations by Komarov (1895) and Arsenyev (1903–1907) in the Far East regions. It was discovered that the berries and seeds were used by Nanai (Goldes or Samagir) hunters as a tonic, to reduce thirst, hunger and exhaustion, and to improve night vision [7,8,9,10]. During the period 1950–60, the idea of using herbal medicinal plants to increase stamina and survival in harmful environment was developed, and a new concept of “adaptogens” was introduced by the toxicologist Lazarev to describe compounds which could increase “the state of non-specific resistance” in stress [11,12]. This concept was based on Hans Selye’s theory of stress and general adaptation syndrome, which have three phases: alarm phase, phase of resistance and phase of exhaustion [13], Figure 1.

Figure 1. Adaptogens increase the state of non-specific resistance in stress and decrease sensitivity to stressors, which results in stress protection, and prolong the phase of resistance (stimulatory effect). Instead of exhaustion, a higher level of equilibrium (the homeostasis) is attained the heterostasis. The higher it is, the better the adaptation to stress. Thus, the stimulating and anti-fatigue effect of adaptogens has been documented in both in animals and in humans. Adapted from [26,78,133].

[Figure image available in the original published article.]

At the end of the 1960s Brekhman and Dardimov proposed that adaptogens are innocuous agents, nonspecifically increasing resistance against physically, chemically, biologically and psychologically noxious factors (“stressors”), normalizing effect independent of the nature of pathologic state [14]. In the early 1960’s, the study of adaptogens developed into a field of biomedical research in its own right in the USSR. This was due to a major targeted project or direction of research, such as mapping or screening of biologically active substances from the plant kingdom. The aim of the stress research was to develop drugs and methods able to stimulate the intrinsic adaptive mechanisms of the organism to help it survive in situations of intense or prolonged stress, whilst preferably maintaining the capability for physical and mental work [15]. The extent of the research carried out was enormous with 1009 studies (primarily pharmacological and clinical) published in Russia up until 1982 and most of them concerned extracts or isolates prepared from *Eleutherococcus senticosus* [16].

Today, research into adaptogens comprises the following four areas: (a) phytochemistry: isolation and structure elucidation of active constituents of adaptogenic plants; (b) biochemistry and molecular biology: mechanisms of stress protective activity of adaptogens on the molecular and cellular levels; (c) experimental and clinical pharmacology: efficacy and safety of adaptogens in stress related disorders on animals and humans; (d) pharmaceutical development of herbal preparations/products that have well established medicinal use in evidence based medicine. Some of the most interesting developments are studies that clearly indicate that certain adaptogenic substances can activate the protective mechanisms of cells, which is linked to an increase in survival rate both *in vitro* and *in vivo* [17,18,19,20]. These studies have so far been directed at the regulation of molecular chaperons (Heat Shock Proteins), such as Hsp70 and other key stress mediators [21,22,23].

As a pharmacotherapeutic group adaptogens were recently defined as herbal preparations that increased attention and endurance in fatigue, and reduced stress-induced impairments and disorders related to the neuro-endocrine and immune systems [23,24]. This definition was based on evidence obtained from clinical trials, which we evaluated in accordance with the European Medicines Agency Assessment Scale [25] and US Natural Standards Evidence–based Validated Grading Rationale [<http://www.naturalstandard.com/grading.html>].

The term adaptogen is often applied to plants, even when the criteria of an adaptogen is not met with, such as the important and significant general adaptive effect on stress involving the whole organism and its main organ and functions. In this review article, we will focus our attention on some of the most extensively studied adaptogens: *Rhodiola rosea*, *Schisandra chinensis* and *Eleutherococcus senticosus* [26,27,28,29,30,31,32,33,34,35,36,37,38] and discuss the results of pharmacological and clinical studies, which are relevant to the main topic of discussion.

The normal paradigm one drug for one disease is not appropriate for adaptogens as they can have numerous pharmacological effects and indications. Table 1 and Table 2 show their pharmacological profiles, which are different but similar in terms of their stress-protective action. Therefore, all these pharmacological effects can be combined into the groups associated with stimulating and stress-protective effects in CNS and vegetative nervous systems, endocrine system and immune system, comprising by definition the parts of neuroendocrine-immune complex - stress-system, Table 3 [26].

The adaptogens *Eleutherococcus senticosus*, *Rhodiola rosea* and *Schisandra chinensis* were reported to be safe in acute and subacute toxicity studies. Moreover, adaptogen induced state of non-specific resistance to highly toxic chemicals (e.g., chlorophos, phosphorus, cyclophosphane, strychnine, aniline, sodium nitrite, narcotics like sodium barbital, hexenal, chloralhydrate, benzene, acetone, ether, etc.) and microbes demonstrated in many pharmacological/toxicological studies [28], actually implies that they have an anti-toxic activity. For example, it was demonstrated that repeated administration of *Rhodiola rosea* extract during 10 consecutive days decreased LD50 of 40% ethanol in mice from 24.1 ml/kg to 55.2 ml/kg. It was also shown that salidroside shortened (from 100% to 19%) the duration of benzene induced sleep in rats [28].

Table 1 Pharmacological profile of adaptogens: summary of *in vitro* or in animal studies.

Regulatory System: effect	Pharmacological Effects	<i>Rhodiola</i>	<i>Eleutherococcus</i>	<i>Schisandra</i>
Stress-system (neuro-endocrine-immune complex): Anti-stress/stress-mimetic/stress-protective	CNS-stimulating: enhancing of physical performance, cognitive performance (learning and memory)	+	+	+
	Neuroprotective	+		+
	Hepatoprotective	+	+	+
	Cardioprotective	+		+
	Gastroprotective		+	+
	Oxidative stress/Radioprotective	+	+	+
	Anti-atherosclerosis		+	+
	Vasodilatory/hypotensive			+
	Anti-hyperglycemic		+	
	Anti-inflammatory/allergy	+	+	+
	Immunotropic	+	+	+
	Antidepressive	+		
	Anxiolytic	+	+	

Table 2 A selected pharmacological profile of adaptogens, clinical efficacy in humans relative to CNS.

	Pathophysiological condition	<i>Rhodiola</i>	<i>Eleutherococcus</i>	<i>Schisandra</i>
Neuro-endocrine system	Physical fatigue	+	+	++
	Mental fatigue (declined attention)	++	+	+
	Stress induced chronic fatigue	+	+	
	Depression	+		

Table 3 Summary of the pharmacological activities of *Schisandra chinensis*, adapted from [26].

Regulatory system		Pharmacological effect	
Stress-system	Central and vegetative nervous system	Stimulating effect	Adapto-genic effect
	Endocrine system	Stress-mimetic and stress-protective effect	
	Immune system	Stress protective effect	

2. Active Compounds

Chemically, the adaptogens are typically either complex phenolics or tetracyclic triterpenoids/ steroids (Figure 2). The phenolic compounds include phenylpropanoids and phenylethane derivatives, such as salidoside (rhodioloside), rosavin, syringin, triandrin, tyrosol, [28,35,38,39,40,41,42,43] and lignans, such as eleutherosid E [44,45,46,47,48] and schisandrin B [49,50,51,52,53,54,55,56,57,58,59,60]. They are structurally similar to the catecholamines -the mediators of the sympathoadrenal system (SAS) involved in activation of the stress system in the early stages of stress response. The tetracyclic triterpenoids, such as cucurbitacin R diglucoside [61,62], ginsenosides [63,64,65,66,67,68] and phytosterol-glycosides (e.g. SG, eleutheroside A, sitoindosides, daucosterol) [27,69,70,71] structurally resemble the corticosteroids that act as stress hormones involved in protective inactivation of the stress system [72,73,74]. In addition, the monoterpene glucoside rosiridin, which is isolated

from *Rhodiola rosea*, was found to inhibit monoamine oxidases A and B *in vitro* implying its potential beneficial effect in depression and senile dementia [75].

Figure 2. Active compounds isolated from *Rhodiola rosea* (tyrosol, salidroside, rosiridin), *Eleutherococcus senticosus* (eyringin, eleutheroside E), *Schisandra chinensis* (schizandrin), *Panax ginseng* (ginsenosides), *Withania somiphra* (sitoindosides), *Bryonia alba* (cucurbitacin R glucoside).

[Figure image available in the original published article.]

Salidroside - the active principles of *R. rosea* extracts - was found to have neuro-protective activity, which reduced stress induced impairments and disorders related to the neuro-endocrine and immune system through: Stimulation of CNS system [28,39,76,77,78]. Protection of cultured neuronal cells from sodium azide and glutamate induced injuries [79,80]. Attenuation of glutamate-induced apoptotic cell death in primary cultured hippocampal neurons of rats [81]. Blockage of H₂O₂-induced apoptosis in rat neuronal PC12 cells [82]. The effect of anti-neuronal apoptosis related to its function of decreasing intracellular free calcium concentration [83,84]. Protection of rat neuronal PC12 cells against amyloid beta-peptide (A beta)-induced cytotoxicity reduced accumulation of reactive oxygen species and malondialdehyde (MDA) [85]. Stimulation of glucose uptake in skeletal muscle cells by activating phosphorylation of AMP-activated protein kinase [86]. Protection from oxidative damage during fatigue [87]. Reduction of the degree of cerebral edema in rats with global cerebral ischemia-reperfusion injury, relieves the metabolism abnormality of free radical and improves the function of cognition [88].

A number of these findings might raise the possibility of potential therapeutic applications of salidroside in preventing and treating cerebral ischemic and neurodegenerative diseases [89].

Tyrosol - another active principle of *Rhodiola* extract - increases phosphorylation of eNOS and FOXO3a, which are key molecular targets involved in this mechanism. Furthermore, it has recently been shown that tyrosol induces the expression of the longevity protein SIRT1 [90].

It is noteworthy that administration of the amino acid tyrosine, which is a common precursor of biosynthesis of tyrosol, salidroside and catecholamines (Figure 2), alleviates both stress-induced depletion of brain catecholamines (norepinephrine and dopamine in the alarm phase of stress syndrome) and reduces fatigue, as noted in animal task performances [91]. A number of clinical studies suggest that supplementation of tyrosine might improve stress-induced (e.g., cold, noise, anxiety and fatigue) accuracy of mental performance [92].

Indeed, schisandrin B has a similar pharmacological profile associated with stress-protective activity. Apparently the neuroprotective effect of schisandrin B [54,55,59,93,94] is associated with the expression of heat shock proteins Hsp70. It has been demonstrated that schisandrin B stimulate the expression of Hsp70 in normal cells [95, 96,97,98,99,100,101], which is associated with the enhancement of mitochondrial glutathione status [95,97,100, 101,102,103], antioxidant activity [59,96,104,105,106,107,108,109], ATP generation [110], mitigation of age-related impairments in mitochondrial antioxidant status and functional ability in various tissues, enhancement in cognitive functions and an increase in the survival of aging in rodents [56,97,112].

3. Antistress and Stimulating Activity of Adaptogens in Animal Model Systems

A typical pharmacological assessment of adaptogens includes the evaluation of stimulating, tonic and stress-protective activities in model animal systems, which were subjected to various stress conditions [112,113].

3.1. Anti-Stress Effect of Adaptogens

The stress protective effect of adaptogens has been demonstrated on simple organisms and on isolated cells [17,18, 114,115]. Thus, there may be an association with regulation and homeostasis of the neuro-endocrine-immune

complex. In addition, there may also be a connection with more evolutionary older congenial mechanisms of regulation in cellular homeostasis and the adaptive/defense response to external stressors. We hypothesize that this type of defence system is apparently common for all cells and living organisms and probably includes heat shock proteins among the number of key mediators of innate nonspecific resistance to stressors.

It has been shown that the same mechanism can be found in stress tolerance and lifespan extension, which makes them parallel phenomenon. Therefore, it is not surprising that adaptogens prolong the lifespan of the nematode *Caenorhabditis elegans* [18] and *Drosophila melanogaster* in a dose dependent manner [19], e.g., see Figure 3, where effect of ADAPT-232—a fixed combination of *Rhodiola rosea*, *Schisandra chinensis* and *Eleutherococcus senticosus* extracts—on the lifespan of *C.elegans* is demonstrated.

Figure 3. Fixed combination of *Rhodiola rosea*, *Schisandra chinensis* and *Eleutherococcus senticosus* extracts (ADAPT-232/Chisan) causes a concentration-dependant increase in life span of N2 wild-type *C. elegans*. A highly significant change in longevity is observed when Kaplan-Meier survival analysis (log-rank test) is used to compare between groups ($p = 0.001$) (Panosyan and Wiegant, unpublished data, 2005).

[Figure image available in the original published article.]

3.2. Molecular Mechanisms behind Anti-Stress Effect of Adaptogens

It has been demonstrated that the beneficial stress-protective activity of adaptogens was associated with the hypothalamic-pituitary-adrenal axis and the regulation of key mediators of the stress response common to all cells, of such as: Heat shock proteins Hsp70 and Hsp16 [17,22,23,95,96,97,98,99,100,101,116,117,118,119,120], which are molecular chaperons involved in stress-induced cytoprotection and in adaptation of repeated exposure to an initial stressor. Stress-activated c-Jun N-terminal protein kinase 1 (JNK1) [21,121]. Forkhead box O (FOXO) transcription factor DAF-16 [17,90]. HPA- axis, including cortisol and glucocorticoid receptors [21,123,124]. Beta-endorphin [125,126,127]. Nitric oxide [21,128,129,130,131]. The biosynthesis of ATP, thus inducing an alteration in energy source [23,131], Figure 4.

Figure 4. A simplified schematic showing the hypothetical molecular mechanism of ADAPT-232 as it induces stress resistance (adaptation to stress) and enhances cognitive functions and, possibly, longevity. Adapted from [23].

[Figure image available in the original published article.]

Typically, a cell is either in: balance (dynamic equilibrium - homeostasis), or functioning under stressful conditions (threatened homeostasis-imbalance), the state of adaptation (tolerance) to stress (i.e., state of non-specific resistance to stress; heterostasis or homeostasis with a higher level of equilibrium), the state of apoptosis (dying).

Panel (a) shows that mitochondria generate aggressive oxygen-containing radicals that can damage native or repair proteins by distorting their 3-D structure, so that they can no longer fulfil their functions in the cell.

There are many “players” involved in the regulation of homeostasis at both the cellular level and the organism level, such as: the stress hormone cortisol (a molecule that is secreted from glands and regulates the functions of organs and systems of the organism). glucocorticoid receptors that modulate/regulate cortisol secretion (feedback regulation). NO, an intracellular signalling molecule that mediates stress response and modulates stress-induced activation of hormonal, nervous and immune systems. FoxO, a Forkhead protein that controls the synthesis of proteins involved in stress resistance, cell survival and longevity.

Panel (b) shows that under stress (e.g., infection, cold, heat, radiation, physical load, emotional stress) an external stress signal activates a cascade of “signalling” proteins/enzymes including JNK, a stress-activated enzyme that plays important roles in the regulation of a diverse array of cellular functions such as neuronal development, activation of the immune system and programmed cell death (apoptosis). The functions of JNK are as follows: To increase the formation of aggressive radicals and nitric oxide, which in turn suppresses the generation of energy

providing molecules (ATP). As a result of lack of energy, many proteins cannot work, several functions are suppressed, and the first symptoms of fatigue and exhaustion are observed. ATP is also required for the normal functioning of heat shock proteins (e.g., Hsp70), which are produced as a defence response to stress and assist in the repair of misfolded and damaged proteins. To suppress glucocorticoid receptors (GR) such that the feedback inhibition of cortisol secretion ceases to function and levels of circulatory cortisol increase. The cortisol inhibits the immune system, and has anti-inflammatory effects on the body. It is also required to protect the organism from over-reaction/over-activation in response to stress. However, chronically high levels of cortisol are associated with depression, chronic fatigue and impaired cognitive function, such as decreased attention and learning ability. To activate translocation of FoxO to the nucleus and initiates the synthesis of proteins that confer stress-resistance and increased longevity.

Panel (c) shows that adaptogens such as ADAPT-232 decrease NO, cortisol and JNK under stress and stimulate/activate the expression of Hsp70 and p-FoxO1. The stimulation of Hsp70 biosynthesis is a key point in the mechanism of action of adaptogens since the heat shock protein: enhances the repair of damaged proteins, inhibits the stress-induced expression of NO genes. ATP is increased to normal levels in the adapted cell. This is due to the inability of reduced levels of NO to suppress the formation of energy providing molecules, inhibits JNK and consequently apoptotic death and suppression of immune system via activation of GR and other mechanisms. Normal GR function and normal ATP levels are associated with the anti-fatigue and anti-depressive effects of adaptogens and with normal cognitive function (e.g., good attention, memory and learning), is probably associated with the effect of adaptogens on the phosphorylation of FoxO and its translocation into the nucleus of isolated cells (i.e., human monocytes) or simple organisms (i.e., DAF-16 in *C. elegans*) and, consequently, with increased resistance to stress and increased life span.

In summary, ADAPT-232 works like a stress vaccine (stress-mimetic) by activating stress-induced self-defence mechanisms in order to adapt the cell and organism to mitigate stress-induced harmful effects.

It seems that activation of Hsp70 expression is a key point in the mode of action of adaptogens [23]. We have demonstrated that adaptogens induce an increase of serum Hsp72 in animal studies [22]. This induction is considered as a defense response to stress, which increases tolerance to stress in a combination of physical and emotional stresses. Our data suggest that increased tolerance to adaptogen-induced stress is associated with its stimulation of expression of circulating serum Hsp72 [22]. In fact, Hsp72 production and release is a known mediator of the stress response involved in repairing proteins during physical load. Our working hypothesis is that adaptogens adapt (or make less sensitive) the organism to stress. Thus, adaptogens act like low molecular weight “vaccines” or stress-mimetics, which induce mild activation of the stress system in order to cope with more severe stress. The adaptogens act as challengers and mild stressors (stress-mimetics) [23]. This gives rise to adaptive and stress-protective effects, which are mainly associated with the HPA axis, a part of the stress system that also contributes to the nervous, cardiovascular, immune, gastrointestinal and endocrine systems [64,132,133].

The anti-depressive effect of *Rhodiola rosea* [134] may be associated with parts of the stress-system (e.g., secretion of cortisol and the JNK mediated effects on the glucocorticoid receptors) [21,122,134] through its effect on mono-amine oxidase A [75].

A recent review [135] focused on the neuroprotective effects of adaptogens. The authors concluded that the adaptogens *Eleutherococcus senticosus*, *Rhodiola rosea* and *Schisandra chinensis* were all involved in the protection of brain neurons from various injuries, which means that they may have an influence on neurodegenerative mechanisms in Parkinson's disease [135].

3.3. CNS Stimulatory Effect

Apparently, stimulating (acute/single dose effect) and tonic (effect of repeated/multiple administration) effects of adaptogens are actually consequences of their stress-protective activity. CNS stimulating and tonic effects of adaptogens are well documented in numerous publications and reviewed in *Phytomedicine* [77]. In contrast to conventional stimulants, such as sympathomimetics (e.g., ephedrine, fenfluramine, phentermine, prolintane) and general tonics, the adaptogens don't possess addiction, tolerance and abuse potentials, they don't impair mental function and lead to psychotic symptoms in long term use, Table 4. Their clinical and pharmacological effects are due to a different mode of action. Their stimulating effect is more pronounced against a background of fatigue and stress.

Table 4 The differences in properties between adaptogens and other stimulants, adapted from [77,78,113,136].

Characteristic		Stimulants	Adaptogens
1.	Recovery process after exhaustive physical load	Low	High
2.	Energy depletion	Yes	No
3.	Performance in stress	–	Increased
4.	Survival in stress	–	Increased
5.	Quality of arousal	Poor	Good
6.	Addiction potential	Yes	No
7.	Side effects	Yes	Rare
8.	DNA/RNA and proteins synthesis	Decreased	Increased

4. Clinical Studies

The most important characteristics of adaptogens, such as stress-protection and a stimulatory effect are common to all adaptogens. However, the effects may differ under various circumstances (Table 1 and Table 2) as has been documented in a number of clinical studies (Table 5) and reviews [14,23,26,28,31,32,36,37,77,78,112,113,137,138,139,140].

The majority of the reviews focused on narrative observations of clinical studies and only a handful conducted a systematic assessment and quality scoring of the level of evidence as recommended by the Natural Standards and the EMEA [23,77].

One such review [31] focused on over 35 clinical trials on *Eleutherococcus senticosus* in healthy human subjects (ca. 6,000 subjects aged from 19 to 72), which were performed in normal and stressful conditions (e.g., high temperature environment, forced work periods, loud noise conditions, motion sickness, varying degrees of deafness, heavy physical burden, hypertension, mountain rescuers under forced conditions, athletes, deep sea divers, intense mental work and physical work, factory workers under extreme working condition). Farnsworth *et al.* observed that there was an improvement of the physical and mental work capacities in all cases. In addition, over 35 studies have focused on the effect of *Eleutherococcus senticosus* on more than 2,200 sick patients [31]. The studies included patients with atherosclerosis, acute pyelonephritis, diabetes, hypertension, trauma, neuroses, rheumatic heart disease, chronic bronchitis, insomnia, cancer and several other ailments. In most cases, a moderate improvement relative to the initial conditions was observed. The extracts were well tolerated and no side effects were observed [31,32].

However, the most convincing evidences of the efficacy of adaptogens were found in studies related to its neuro-protective effects, effects on cognitive functions and mental performance in fatigue, and on its efficacy in asthenia

and depression [23]. The evidence points to adaptogens may be beneficial on neurodegenerative disorders [54,83,84,89,135,141].

4.1. Adaptogens in Fatigue, Effect on Cognitive Functions

In total, more than 30 publications on the clinical efficacy of various *Rhodiola rosea* preparations can be found in the PubMed database. The majority of these studies are of varying methodological rigor and concern cognitive functions and mental performance under fatigue, Table 5.

The clinical trials using *Schisandra chinensis* (13 studies) and *Eleutherococcus senticosus* (11 studies) on mental performance in humans have been the subject of a recent review [23]. A systematic review showed that adaptogens have a significant, beneficial and specific effect on stress-induced symptoms under fatigue [23]. It was seen that *Rhodiola rosea*, in particular, significantly reduced symptoms of fatigue and improved attention after four weeks of repeated administration [122]. Furthermore, it was suggested that the inhibitory effect of *Rhodiola rosea* on the increased basal level of salivary cortisol results in an improvement of cognitive function. This is in line with other studies demonstrating that optimal corticosteroid levels are a requirement for efficient cognitive function. It has been shown that significant changes (upregulation or downregulation) of circulating levels of corticosteroids are associated with cognitive impairment [168]. Moreover, studies on healthy volunteers receiving single and repeated doses of SHR-5 extract (*Rhodiola rosea*) have demonstrated an anti-fatigue effect and improvement in cognitive functions during fatigue and in stressful conditions [143,144,145]. Thus, one may conclude that repeated administration of *R. rosea* extract (SHR-5) exerts an anti-fatigue effect on healthy subjects and burnout patients expressing fatigue syndrome. This in turn, increases the patient's mental performance and ability to concentrate.

Table 5 Results of clinical studies on humans involving effects of plant adaptogens on physical and mental performance related to fatigue.

Plant name Intervention/ control/ Daily dose/ Duration	Study design ^a / Total subjects	Primary endpoint and main results ^b	Quality level of evidence*	Jadad score [183]*	Ref.
<i>Rhodiola rosea</i> Extract SHR-5 (288 mg twice daily)/ placebo for 4 weeks	R, PC, DB 2 parallel groups 60 volunteers with stress- induced fatigue, (30/30) [20–55 years]	Symptoms of fatigue, attention, depression, QOL, salivary cortisol. Symptoms of fatigue, attention and salivary cortisol significantly improved compared with control	Ib	5	[122]
<i>Rhodiola rosea</i> Extract SHR-5 (170 mg once daily)/ placebo for 2 weeks	R, PC, CO, DB 2 parallel groups 56 healthy subjects (??) ^c [24–35 years]	Mental fatigue, perceptive and cognitive functions such as associative thinking, short-term memory, calculation and ability of concentration, and speed of audio-visual perception Statistically significant improvement in the treatment group (SHR-5) during the first 2 week period	Ib	4	[143]
<i>Rhodiola rosea</i> Extract SHR-5 (50 mg twice daily)/ placebo for 20 days	R, PC, DB 2 parallel groups 40 healthy subjects (20/20) [17–19 years]	Mental fatigue, physical performance, general well-being Significant improvement in physical fitness, mental fatigue and neuromotor tests compared with control ($p < 0.01$). General well-being was also significantly ($p < 0.05$) better in the verum group. No significance was seen in the correction of text tests or a neuromuscular tapping test	Ib	3	[144]
<i>Rhodiola rosea</i> Extract SHR-5 (single dose of 370 mg or 555 mg) / placebo	R, PC, DB 3 parallel treatments groups 161 healthy subjects, (41/20/40 treated + 20 untreated) [19–21 years]	Capacity for mental work Significant difference in anti-fatigue effects in SHR-5 groups compared with control ($p < 0.001$), whilst no significant difference between the two dosage groups was observed.	Ib	3	[145]
<i>Rhodiola rosea</i> Extract SHR-5 (170 mg or 340 mg twice daily)/placebo for 6 weeks	R, PC, DB 3 parallel treatment groups 91 patients with mild and moderate depression (31/30/30) [18–70 years]	Depression in total HAMD and BDI scores Significant differences in HAMD and BDI scores and scores reflecting levels of insomnia, emotional instability, somatisation and self-esteem in SHR-5 groups compared to placebo ($p < 0.001$)	Ib	5	[134]
<i>Eleutherococcus senticosus</i> Extract (2mg eleutherosides B and E equivalent to 2–4 g of herbal substance daily)/placebo for 2 months and 2 months of follow-up	R, PC, DB 2 parallel treatment groups 96 patients with chronic fatigue syndrome (49/47) [21–65 years]	RVI measurement of fatigue reduced Significant ($p < 0.05$) improvement in RVI compared with control after 2 months treatment in the subgroups of patients with moderate fatigue at baseline (RVI value 8–12) and a history of fatigue <5 years. However, no significant difference was observed after 4 months treatment	Ib	5	[142]
Chisan-fixed combination of	R, PC, DB 2 parallel groups	Duration of antibiotic therapy associated with the clinical manifestations of the acute phase of the disease, mental	Ib	4	[146]

Rhodiola-Schisandra- Eleutherococcus	60 patients suffering from acute non- specific pneumonia (30/30) [18–65 years]	performance and self-evaluation by WHO-QOL brief questionnaires Adjuvant therapy with ADAPT-232 had a positive effect on the recovery of patients by decreasing the duration of the acute phase of the illness for 2 days, by increasing mental performance of patients in the rehabilitation period, and by improving their QOL			
ADAPT-232 capsules [fixed combination of standardised extracts of <i>Rhodiola rosea</i> (salidroside 3 mg), <i>E. senticosus</i> (syringin 3 mg), <i>S. chinensis</i> (schizandrin 4 mg)] 3 capsules <i>acute</i>	PC, CO, DB 60 PC, CO, DB 5 cosmonauts	Compared with placebo, ADAPT-232 significantly increased short-term memory, speed and reliability in the understanding of information, and precision and accuracy in the ability to reproduce the information in repeated highly sophisticated computer-based tests (Monotonic 2). ADAPT-232 was most effective against a background of pronounced fatigue induced by monotonous night work. The effect was most marked in complicated tests and under extreme conditions. ADAPT-232 significantly decreased the number of mistakes (<i>cf.</i> with placebo) in complicated psychometric tests 4 h after administration and increased working capacity 1.5 and 4 h after administration to Russian cosmonauts during their training in prolonged isolation (90 days) under conditions of long-term, monotonous work. No significant effects were observed in non-complicated tests.	Ila Ila	1	[147] [148]
<i>Eleutherococcus</i> <i>senticosus</i> extract 0.25, 0.5, 1.0, 2.0, 4.0 and 8.0 mL <i>acute</i>	PC, CO, OL 357 Sailors	<i>E. senticosus</i> improved mental performance in correction test; increased activity of the adrenal cortex, the activity of the sympathetic adrenomedullary system, the intensity of metabolic processes, and the intensity of red-ox processes. In stress conditions <i>E. senticosus</i> decreased adrenal cortex activity and sympathetic nervous system; increased the tonus of the parasympathetic nervous system; moderately intensified excitation of the CNS and of energy metabolism; improved endurance to hypoxia.	Ila	1	[149]
<i>Schisandra chinensis</i> extract 0.25, 0.5, 1.0, 2.0, 4.0 and 8.0 mL <i>acute</i>		<i>S. chinensis</i> stimulated the activity of the CNS at night; increased tonus of the sympathetic part of the autonomic nervous system (having no effect on the parasympathetic part) after night duty; activated the adrenal cortex activity; increased the activity of the cardiovascular and respiratory systems; intensified oxidation-reduction and metabolic processes; improved working ability parameters; reduced parameters of non-specific resistance the organism.			
<i>Rhodiola rosea</i> (extract) 10 drops or 3 x 10 drops/day <i>Rhaponticum</i> <i>cartamoides</i> (extract) 40 drops or 3 x 10 drops/day. Acute and 10 days	PC, SB 80 healthy students (control group) and 70 patients with neurosis	Single and repeated administration of adaptogens improved functional state of the CNS in patients with neurosis as characterised by normalisation of the speed and power of neural processes in Ivanov-Smolenski's verbal test with speech-supported locomotor-conditioned reflex measurement. The memory improved and attention became more stable.	Ila	1	[150]
Rhodosin (<i>R. rosea</i> extract, 100 mg/ 20 days	PC, DB 60	Significant improvement in physical fitness, mental fatigue and neuromotor tests compared with control ($p < 0.01$); general well-being was also significantly ($p < 0.05$) better in the verum group. No significance was seen in the correction of text tests or a neuromuscular tapping test.	Ila	1	[151]
Rodelim tablets (fixed combination of the extracts of <i>R.</i>	PC, DB 60	Rodelim improved mental working capacity in computer and correction tests against a background of fatigue.	Ila	1	[152]

<i>rosea</i> , <i>E. senticosus</i> and <i>S. chinensis</i>) 100 mg/acute					
<i>R. rosea</i> (tincture 40% ethanol) 5-10 drops/acute	PC/85	Improved mental performance, reduced the number of errors in Anfimov's correction test: the stimulating effect lasted 4 h or more	I Ib	1	[153]
<i>Rhodiola rosea</i> (40% ethanol tincture, 20 drops) <i>E. senticosus</i> (40% ethanol tincture, 20 drops) Extract of <i>R. rosea</i> rhizome, 0.3 g/ acute	PC/254	Improved mental performance; reduced the number of errors in Anfimov's correction test; increased the accuracy, working capacity and speed of information perception. Stimulating effect lasted 4 h or more	I Ia	1	[154]
Tyrosol (1, 5, 10 and 20 mg), <i>R. rosea</i> extract, 5 drops	? / 82 patients	Improved mental performance, reduced the number of errors in Anfimov's correction test.	III	0	[155]
Salidroside, 2.5 mg/ acute	PC, SB/ 46	Improved mental performance; reduced the number of errors in Anfimov's correction test; stimulating effect lasting 4 h or more.	I Ia	1	[39]
<i>E. senticosus</i> (dry extract), 120 mg/ Acute and 3 weeks	PC, DB/40	Improved mental performance determined by a letter correction test	I Ia	1	[156]
<i>E. senticosus</i> (tincture) <i>P. ginseng</i> tincture/ 2 ml/ acute	PC, SB,CO/ 13	Decreased errors in data sent by radio operators 1 h after drug uptake. Stimulating effect of <i>E. senticosus</i> was stronger and more stable than the effect of Ginseng, which was insignificant.	I Ia	1	[157]
Eleutherococcus (extract 1250 mg) Ginkgo (extract, 28 mg flavonoids, 7 mg ginkgolides) 3 months	PC, CO /24	Selective memory significantly improved compared with placebo ($p<0.02$).	I Ia	1	[158]
<i>S. chinensis</i> (tincture) <i>E. senticosus</i> (tincture) <i>Rhaponticum cartamoides</i> (tincture) <i>Oplopanax elatus</i> (tincture), 0.5, 1.0 and 2.0 mL, acute	PC, CO, OL 1327 healthy subjects	Eleutherococcus and Rhaponticum treatment significantly increased precision in tremometric test compared with placebo. Significant differences compared with placebo were observed with various doses of all tested preparations in psychometric tests including assessment of attention and memory functions.	I Ia	1	[159]
<i>S. chinensis</i> (tincture) <i>E. senticosus</i> (tincture) <i>Rhaponticum cartamoides</i> (tincture) <i>Oplopanax elatus</i>	PC, CO 665 healthy subjects	Pilots were tested before a flight and again 5 –15 min later, and finally 1 and 3 h after landing Significant differences compared with placebo were observed for all tested preparations in tests including assessment of precision, dynamic tremometry, sensomotor response, and attention and memory functions	I Ia	1	[160]

(tincture, 0.5, 1.0 and 2.0 mL), 10 days					
<i>S. chinensis</i> (tincture, 2 ml) <i>E. senticosus</i> (tincture, 2 ml) Coffee, 200 ml Black tea, 200 ml (control) Acute and repeated	OL, PC 200 sailors	<i>S. chinensis</i> showed a tonic effect for 4–7 h in those on night duties. <i>E. senticosus</i> was inactive Repeated uptake (for more than 2 weeks) of coffee and <i>S. chinensis</i> produced similar negative effects (insomnia, excitability, etc.).	I Ib	0	[161]
<i>S. chinensis</i> (seed powder, 2g) <i>Panax ginseng</i> (tincture, 2 ml), acute	CO, PC, SB, 122 students	Improved attention in text correction 2 h after drug administration. Both extracts increased quality and quantity of mental work performed.	IIa	1	[162]
<i>S. chinensis</i> (seed tincture, 30 ml) <i>P. ginseng</i> (3% tincture, 30 ml) Phenamine(0.02 g) Glucose (0.5 g), acute	CO, PC /20	<i>P. ginseng</i> and <i>S. chinensis</i> improved accuracy in the work of telegraph operators at exhaustion compared with control group (glucose) and those given phenamine.	IIa	1	[51]
<i>S. chinensis</i> (seed – 3 g, extract and fractions, tablets and capsules, 0.036–0.168 g), acute	OL,CO/20	Improved accuracy in error correction test. Most active stimulating effect was revealed by a crystalline substance identified as the lignan schizandrin.	I Ib	0	[50, 51]
Schizandrin (0.02, 0.01, 0.005 g), Phenamine (0.2 g) Glucose (0.5g) acute	PC/23	Schizandrin improved accuracy in the work of telegraph operators at exhaustion compared with control group (glucose) and those given phenamine.	I Ib	1	[29,51]
<i>S. chinensis</i> (seed powder, 0.25 and 0.5 g), 10 days	OL / 36	Beneficial effects on symptoms of astheno-depressive syndrome	IIIa	0	[163]
<i>S. chinensis</i> (fruit and seed tincture, 1:5, 90% ethanol, 1 ml), 16–40 days	OL/ 40	Stimulating effect-improve mood, increases physical and psychological vivacity, relief of tiredness and fatigue in asthenia and depressions.	III	0	[164]
<i>S. chinensis</i> (tincture, decoction and tablets), 2–10 weeks	OL/ 2000	Stimulating and tonic effect.	III	0	[165]
<i>S. chinensis</i> (tincture, decoction and tablets), 2–10 weeks	OL/ 250	Effective in the treatment of general asthenia, exhaustion and reduced physical and mental performance in group of patients with nervous disorders where an increase in general well-being and working capacity, as well as a decrease in sleepiness and exhaustion, were observed.	III	0	[166]
<i>S. chinensis</i> (tincture 95%) 45–120 drops <i>S. chinensis</i> (tablets, 0.25 g, 0.5 g), 10 days	OL, observational study/ 30	Stimulating effect in astheno-depressive syndrome with relief of somnolence, limpness, tiredness, fatigue	III	0	[167]

^a R – Randomized, PC – placebo– controlled; DB – double– blind; SB – single blind, NC – not controlled; CO – crossover, UC – uncontrolled, OL – open label trial; ^b According to WHO, FDA and EMEA: Ia – meta– analyses of randomised and controlled studies; Ib – evidence from at least one randomised

study with control ; IIa – evidence from at least one well– performed study with control group; IIb – evidence from at least one well– performed quasi– experimental study; III – evidence from well– performed non– experimental descriptive studies as well as comparative studies, correlation studies and case– studies; and IV – evidence from expert committee reports or appraisals and/or clinical experiences by prominent authorities; ^c ? – data not listed or unavailable.* –max 5.

^a R – Randomized, PC – placebo– controlled; DB – double– blind; SB – single blind, NC – not controlled; CO – crossover, UC – uncontrolled, OL – open label trial; ^b According to WHO, FDA and EMEA: Ia – meta– analyses of randomised and controlled studies; Ib – evidence from at least one randomised study with control ; IIa – evidence from at least one well– performed study with control group; IIb – evidence from at least one well– performed quasi– experimental study; III – evidence from well– performed non– experimental descriptive studies as well as comparative studies, correlation studies and case– studies; and IV – evidence from expert committee reports or appraisals and/or clinical experiences by prominent authorities; ^c ? – data not listed or unavailable.* –max 5.

4.2. Adaptogens in Asthenia and Psychiatric Disorders

It should be mentioned, that the majority of clinical studies cited here are the most questionable and poorly documented as standardized psychological measures were not used in the earlier studies. Indeed, some of them did not use randomization, or blinding of subjects. However, the main problem in assessment of these studies is that the Soviet diagnostic criteria were different from commonly used criteria in the rest of the world [169,170,171]. The diagnostic criteria used in the USSR prior to 1990 for schizophrenia was particularly idiosyncratic, overused, and misapplied to other conditions [169,170,171]. For example, some patients who would be diagnosed as having psychotic depression, schizotypal disorder, schizoaffective disorder, or bipolar disorder by a non-Soviet psychiatrist would have been diagnosed as schizophrenic by a Soviet psychiatrists [166,167,168]. Indeed, the term “sluggish-schizophrenia” was particularly misused [172]. The diagnoses of asthenia and neuroasthenia include a very heterogeneous group of patients with mixed psychological and physical disorders, making the studies more difficult to interpret.

Nevertheless, in spite of numerous shortcomings, which reduced the quality of evidences obtained in the early clinical studies in Russia, these scientific evidences provide important information about efficacy and safety of adaptogens in the treatment of psychiatric disorders, table 6. Some of these publications are discussed below.

Table 6 Summary of clinical trials of *Rhodiola rosea* and *Schisandra* preparations in psychiatric disorders.

Pathophysiological condition: Pharmacological activity or effect recorded	Plant name	Reference	Study design	Quality in Jadad's score	Number of patients
Fatigue syndrome:	<i>R.rosea</i>	Olsson, 2009	R,PC,DB	5	60
Adults physical and cognitive deficiency	<i>R.rosea</i>	Fintelman, 2007	OL	0	120
Depression:	<i>R.rosea</i>	Darbinyan, 2007	R,PC,DB	5	91
	<i>R.rosea</i>	Brychenko, 1987**	OL, C**	0	78/56**
	<i>S.chinensis</i>	Staritsina, 1946	OL, UC	0	37
Astheno-depressive syndrom (stress-induced mild depression):	<i>R.rosea</i>	Krasik, 1970*	OL, UC	0	128
	<i>R.rosea</i>	Krasik, 1970*	OL, UC	0	135/27
	<i>R.rosea</i>	Mikhailova, 1983	OL	0	58
	<i>R.rosea</i>	Mesheryakova, 1975	OL	0	25
	<i>S.chinensis</i>	Zakharov, 1956	OL, UC	0	13
	<i>S.chinensis</i>	Leman, 1952	OL, UC	0	40
	<i>S.chinensis</i>	Rossiiskij, 1952	OL, Uc	0	260
Neurosis (stress-induced depression)	<i>R.rosea</i>	Saratikov, 1965*	OL	0	65
	<i>R.rosea</i>	Kaliko, 1966	PC, SB	1	70/80 ^{\$}
	<i>S.chinensis</i>	Sudakov, 1986	OL, UC	0	386*
Anxiety:	<i>R.rosea</i>	Bystritsky, 2008	OL	0	10
Schizophrenia	<i>S.chinensis</i>	Romas, 1958, 1962	OL	0	79/41*
	<i>S.chinensis</i>	Zakharova, 1948*	OL, C	0	30/20*
	<i>S.chinensis</i>	Lastovetskiy,	OL	0	48

^a R – Randomized, PC – placebo– controlled; DB – double– blind; SB – single blind, NC – not controlled; CO – crossover, UC – uncontrolled, OL – open label trial; ^c ? – data not listed or unavailable.* – mixed patients population, sick/heatly subjects. ** – adjuvant therapy with antidepressants, control group – tricyclic antidepressants. \$ – sick/heatly subjects.

^a R – Randomized, PC – placebo– controlled; DB – double– blind; SB – single blind, NC – not controlled; CO – crossover, UC – uncontrolled, OL – open label trial; ^c ? – data not listed or unavailable.* – mixed patients population, sick/heatly subjects. ** – adjuvant therapy with antidepressants, control group – tricyclic antidepressants. \$ – sick/heatly subjects.

4.2.1. *Rhodiola Rosea*

In 1969, the Pharmacological Committee of the Ministry of Health of the USSR recommended the medicinal use of *Rhodiola rosea* extract in patients suffering from illnesses, such as asthenia syndrome, neuroses, vascular dystonia, hypotension, and schizophrenia (remissions of asthenic type). The advised dosage of the extract was 5–25 drops (in a quarter of a glass of water) three times daily and should be taken 15–30 minutes before meals. Although, the duration of the therapy was individually determined and varied from 10 days to 4 months [173,174].

The symptomatic characteristics of asthenic syndrome is general weakness, reduced working capacity, low memory, irritability, headaches, insomnia, and poor appetite. These symptoms are frequently observed after intensive work, which requires high levels of mental exertion. Indeed, the *Rhodiola rosea* extract was reported to be a valuable medicine in essentially healthy people placed under heavy physical or mental workloads [175,176]. Furthermore, to take advantage of the prophylactic measures of *Rhodiola rosea* extract it has been advised that

doses should be taken in the morning, or in the morning and the afternoon, for at least a few days prior to a period of intensive mental work for a during 2–3 weeks. In cases of hard mental work, it has been recommended that duration of *Rhodiola rosea* therapy last for the duration of the period. It has been reported that therapy with *Rhodiola rosea* extract prevented exhaustion and fatigue during work requiring long intensive mental activity [175, 176].

The positive therapeutic effect of *Rhodiola rosea* extract was also observed in an open label, uncontrolled study in 128 patients with pronounced states of fatigue of different genesis [175]. For example, 128 patients between 17 and 55 years of age (male/female: 75/53) were treated with an *R. rosea* extract and as a result the clinical symptoms of fatigue were considerably improved or disappeared completely. Their diagnosis of fatigue was confirmed by psychological examinations and increased mental work capacity. Indeed, there are also therapeutic effects of *Rhodiola rosea* extract in exhaustion due to psychogenic and somatogenic origin (82%) or in patients convalescing from somatic and infectious diseases (80%) [175]. For example, patients suffering from post-influenza fatigue showed improved mental and physical working capacity already on the third day after intake of *Rhodiola rosea* extract. Thus, these patients improved in focusing their attention, and headaches vanished. Furthermore, *Rhodiola rosea* extract reduced and eliminated general weakness and fatigue in cases of traumatic cerebrastronia injury and also significantly facilitated the normalisation of the autonomic functions in this group of patients [175].

That *Rhodiola rosea* extract has a pronounced therapeutic effect on neuroses (exhausting depression) was reported in an open label study on 65 poorly characterized mixed patient population [177] suffering from insomnia, high irritability and various somatic disorders. The state of the higher nervous activity was tested in these patients with the help of a verbal test and the conditional motor reflexes were tested with verbal reinforcement, in conjunction with routine clinical examinations. The results of the verbal tests showed that the latent period of verbal reactions in most of the subjects was long-up to 1.8–6.0 s (a normal period is 1.5 s). Indeed, results of the motor-verbal test prior to treatment showed that two thirds of the patients performed poorly in mental work situations. However, after a course of therapy with *Rhodiola rosea* extract (10 drops; 3 times daily for 10 days) the excitation and inhibitory processes intensified and mobility normalized. In fact, the conditional motor reflexes also changed. It was seen that their value and stability increased and that the latent period reduced, whilst concentration improved and the interaction of both signal systems were normalized. Furthermore, the extract therapy reduced the latent time of verbal reactions and improved attention and memory.

Indeed, the authors observed a clinical improvement, such as disappearance of irritability and unpleasant sensations in the heart region and improvement of sleep and appetite, in these patients. Similar clinical results were obtained in another study with 177 patients suffering from vascular hypotension. The authors observed that a course of therapy with *Rhodiola rosea* extract stabilized, completed or partially normalized 92% of the hypotensive patients. Thus, the patients felt better, headaches disappeared, sleep became normal and they recovered their working capacity.

Several clinical studies of the dynamics of asthenic symptoms have showed that *Rhodiola rosea* extract was able to diminish or eliminate depressive symptoms. For example, Mikhailova reported that an open label clinical trial of *Rhodiola* extract (15 drops three times daily for a month or up to four months in some cases) in 58 patients with stress induced depression (asthenia of exogenous organic genesis), showed a remarkable symptomatic improvement was observed [178]. Their general weakness or feeling of being worn out in the morning with a high fatigability, and hypersomnia during the day (without disturbance of the following night's sleep) either disappeared or became considerably reduced. In addition, no side effects were observed except in one patient who showed sleep disturbance. In this case, the dose was reduced to 6 drops twice a day and the adverse symptoms immediately disappeared. Indeed, most patients (39/58) suffering from pre-and intra somnia (*i.e.*, difficulties in

falling asleep, waking up during the night) noted sleep improvements. However, it should be noted that sleep normalisation and reduction of other asthenic symptoms depended on the intensity of these disorders [178].

Furthermore, it was seen that patients in depressive states of different genesis normally treated with tricyclic antidepressants reduced the time spent in hospital whilst using them in combination with *Rhodiola rosea* [179]. The authors observed adjuvant therapy with *R. rosea* increased general activity, intellectual and physical productivity of patients, and decreases side effects induced by the tricyclic anti-depressants.

It was reported that *Rhodiola rosea* extract was successfully used in the alleviation of side effects after psychotropic therapy in schizophrenic patients [176]. In this short communication is mentioned that 31 patients suffering from pronounced clinical symptoms of extrapyramidal syndrome, which resulted from therapy by neuroleptics, received a high dose of *Rhodiola rosea* extract (25–40 drops; 2–3 times daily) for a duration of 1–1.5 months. In addition, 19 of these patients received a combination of a high dose of *Rhodiola rosea* extract and cholinolytic Romparkin (Trihexylphenidyl), as this preparation alone failed to eliminate or soothe the clinical symptoms of the side effects in this group of patients. The results of this study showed that *Rhodiola rosea* extract had the most pronounced therapeutic effect on symptoms of Parkinsonism and asthenia.

In conclusion, the introduction of *Rhodiola rosea* extract in the USSR was based on the following findings: (i) as stimulants for essentially healthy people in a state of fatigue and for patients with asthenic states during the rehabilitation period following somatic or infectious diseases. (ii) for essentially healthy people *Rhodiola rosea* extract should be taken several days before the expected strain and during the whole period of increased mental exertion as a prophylaxis. (iii) to recover mental and physical working capacity during and after long periods of intensive mental and physical work. (iv) and sexual disorders related to impotence in males. (v) in psychiatric practice as an adjuvant to counteract side effects of psychopharmacological therapy [28].

Furthermore, side effects of therapy with *R. rosea* extract are relatively rare. Although, some cases of individual sensitivity to the preparation have been observed, such as excitement, irritability, insomnia, and headache.

4.2.2. *Schisandra chinensis*

Schisandra chinensis extracts (e.g., tincture, decoction and tablets) have been reported to be effective in the treatment of general asthenia, exhaustion and reduced physical and mental performance in open label studies on a population of 2,000 poorly characterized mixed patients. It was reported that the extracts showed improvement after 2–10 weeks of therapy [165,166]. Furthermore, the treatment showed a particularly remarkable effect on a group of patients (n = 200) with nervous disorders where an increase in general well-being and working capacity, as well as a decrease in sleepiness and exhaustion [165,166]. Another study [180], which was based on 13 patients, concluded that two administrations of *S. chinensis* extract per day was optimal and that prolonged treatment may cause negative effects. The author pointed out that the duration of treatment should be decided on an individual patient basis. Taken together, the results of the treatment showed considerably improved results in short term patients, in comparison to chronic patients [180].

A comparative clinical pilot study of *S. chinensis* seeds (tincture) in neurasthenic patients (n = 95) showed that there was a high efficiency present in the therapy (dosage: 15 drops during 25–28 days). The results showed that general weakness, poor sleep and appetite, high irritability and headaches disappeared almost completely in the treatment group, whilst 55% of the control group complained of these symptoms. In addition, the muscular force of the hands increased around 2.5 times in the treatment group, in comparison to the control group. It was also seen that the gain of vital lung capacity reached 19% in the treatment group, in comparison to 3% in the control group [181].

Several studies have reported a positive therapeutic effect of *S. chinensis* preparations on asthenic and asthenodepressive states (particularly in stress induced depressions) and schizophrenia [184,185,186,187,188,189,190,191,

192]. For example, Leman observed a stimulatory effect of a *S. chinensis* preparation on the CNS after 16–40 days of treatment in 40 patients with asthenia and depressions of psychogenic or somatic origin [164]. There was also an improvement in dark vision found in almost all patients. Twenty two patients felt pleasantly warm, became more energetic and physically active, whilst hunger and fatigue disappeared. In addition, their mood improved and night sleep was normalised. The authors concluded that a therapy with *Schisandra chinensis* can be indicated in asthenia and depression of psychogenic etiology, so called “exogenous” (stress induced) depression, or states related to excessive fatigue, somatic and nervous exhaustion. However, they also reported that the therapy can only relieve symptoms in “endogenous” depression (organic etiology), asthenia, narcoleptic and amnestic syndromes. An advantage of treatment with *S. chinensis* is the absence of serious side effects. Therefore, the tolerance to *S. chinensis* extract is several times higher than the tolerance to caffeine or phenamine, and the effect is not reduced following a long treatment period.

Another study reported total recovery of psychosis following treatment with *S. chinensis* extract [182]. The authors treated 36 patients (19 with schizophrenia; six with reactive psychosis; four with alcoholic psychosis; three with depression; and four with psychopathy) with ADS three times daily with *S. chinensis* preparations. All in all, total recovery was observed in patients suffering from psychosis, while no effects were seen in patients with diagnosed psychopathy.

Furthermore, a tincture of *S. chinensis* was used in treatment of schizophrenia (79 patients) and chronic alcoholism (118 patients and 41 healthy subjects) in an open label observational trials [184]. The author measured pupillary and vascular (alterations of palm vessel volume) tentative reactions, and refractory reactions in order to evaluate the effects on the CNS. These reactions were comparatively suppressed in schizophrenia and chronic alcoholism. However, it was shown that treatment with *S. chinensis* tincture normalised the reactions in the patients under study and activated reactions in healthy subjects. As a result of the treatment, patients became calm, sociable, and active, free of emotional tension and anxiety, and willing to work. Indeed, they showed excellent well-being, which was associated with a good mood. The author reported an optimal dose of 15–25 drops in simple and hallucinogenic-paranoid schizophrenia, 5–15 drops in paranoid schizophrenia, and 5 drops in catatonic schizophrenia [184].

Furthermore, two open, uncontroled studies [184,185] reported a somewhat unexpected finding in that there was a stimulatory effect in schizophrenic patients and chronic alcoholics. This was due to the ability of the *S. chinensis* extract to increase the reactivity to insulin, sulphadiazine and apomorphine. Thus, administration of a *S. chinensis* preparation, which was used in conjunction with apomorphine, frequently eliminated or decreased addiction to apomorphine. The authors noted development of a stable conditional emetic reflex in most of the chronic alcoholics at the end of the treatment [184,185,186]. It was also found that *S. chinensis* treatment increased associative processes in the subjects and improved the quality of associations, as demonstrated by the increased number of higher verbal responses and a decrease in the number of lower responses [187].

It was also reported that *Schisandra chinensis* extract had a positive effect in an open, uncontroled study on 386 patients with nervous and/or mental disorders of borderline exogenous-organic genesis (stress induced mild depression) [188]. These positive therapeutic effects were the result of a stimulating action and a prevention, or reduction, of development of side effects associated with treatment with tranquilizers and anti-depressants. Furthermore, the combination of *Schisandra chinensis* extract with nootropic preparations (e.g., nootropil, pyracetam) was found to be effective in patients with pronounced mild cognitive (mnestic-intellectual) disorders. Although, in patients with agitative dementia the *Schisandra chinensis* therapy was less effective [188]. Another observation, which could be of practical future importance, is that a combination of *S. chinensis* therapy with tranquilizers or anti-depressants (e.g., amitriptyline, relanium) may be very effective in the elimination of the undesirable effects of these drugs. For example, the development of side effects (e.g., headaches, dizziness, flaccidity, dryness in the mouth, and urination disorders) were observed in 23 out of 39 patients (53.5%) with

neuro-mental disorders of exogenous-organic genesis, which when treated with increased doses of amitriptyline (from 50 to 75 mg). However, patients treated with *S. chinensis* experienced a similar effect in only 4 out of 172 patients (1.9%). Thus, a combined administration of tranquilisers and adaptogens permitted the usage of optimal doses of these drugs in 96% of patients, whilst it was only possible in 16% of control patients ($p < 0.001$). It has been concluded that: (i)*S. chinensis* extract can be used in psychiatric practice as a symptomatic agent against astheno-depressive states independent of the nature of the disease.(ii)preparations of *S. chinensis* decrease fatigue, improve the general mood and appetite, and can be recommended as a tonic for healthy people in a state of fatigue. No negative effects were observed on the somatic state of patients.(iii)*S. chinensis* extract can be used in the treatment of psychoses as a stimulant without harmful side effects.(iv)the curative effect of *S. chinensis* preparations is pronounced in cases of asthenic and depressive syndromes.(v)the combination of *S. chinensis* therapy with tranquilisers or anti-depressants eliminates the side effects of these drugs and allows them to be employed at optimal doses [188].

We may suggest, based on the reported studies, that *Schisandra chinensis* have a place in the treatment of schizophrenia in reduction of catatonia and negative symptoms, relief of fatigue, and amelioration of side effects from neurotropic and other sedating medications. However, the evidence for the use of *S. chinensis* in the treatment of schizophrenia is not scientifically rigorous. In our view, studies which claim that *S. chinensis* significantly improves symptoms of schizophrenia must be validated by accepted diagnostic criteria (ICD-10), rigorous methodology, and validated assessment tools before any conclusions may be drawn.

5. Conclusions and Perspectives of Implementation

Recent pharmacological studies of some adaptogens give a rationale to their effects at the molecular level. It has been shown that the beneficial stress-protective effect of adaptogens is related to the regulation of homeostasis via several mechanisms of action, which are associated with the hypothalamic-pituitary-adrenal (HPA) axis and the regulation of key mediators of the stress response, such as molecular chaperons (e.g., Hsp70), stress-activated c-Jun N-terminal protein kinase (JNK1), Forkhead box O (FoxO) transcription factor, cortisol and nitric oxide (NO). In summary, adaptogens may be regarded as a novel pharmacological category of anti-fatigue drugs that: (i)induce increased attention and endurance in situations of decreased performance caused by fatigue and/or sensation of weakness.(ii)reduce stress-induced impairments and disorders related to the function of stress (neuro-endocrine and immune) systems.

It was suggested that adaptogens have not only specific therapeutic effects in some stress-induced and stress-related disorders, but will also have an impact on the quality of life of patients when implemented as adjuvants in the standard therapy of many chronic diseases and pathological conditions (e.g., post-surgery recovery, asthenia, congestive heart failure, chronic obstructive pulmonary disease). It may be suggested that adaptogens have potential use in age related disorders, such as neurodegenerative diseases, and cardiovascular diseases. Thus, elderly people may be able to maintain their health status on a normal level, improve their quality of life and may increase longevity. However, further research may be needed to evaluate the efficacy of adaptogens as geriatrics and to elucidate molecular mechanisms of action of these complex herbal extracts and their active principles.

References

1. Drake K.V.. On the pharmacology of . *Farmakol. Toksikol.*. 1942. 5:32-36
2. Astanin Yu. P., Mikhelson M.Ya., Egolinskij Ya.A.. The study of phenamine and some other stimulants effect on sport running. *Farmakol. Toksikol.*. 1943. 6:60-65
3. Varlakov M.N.. On the stimulating effect of . *Pharmacia*. 1944. 6:31-34

4. Yefimov V.V., Vlasova L.N.. The effect of Schizandra-stimulant on nervous-muscular excitability in man. *Biul. Eksp. Biol. Med.*. 1945. 19:29-30
5. Zhestyanikov V.D.. Some data on the effect of the Far East Schizandra on the CNS. *Scientific Papers of 3rd Year Students of the Naval Medical School*. 1945. 3:24-29
6. Karo V.I.. The ergographic study of Schizandra stimulating effect. *Scientific Papers of 3rd Year Students of the Naval Medical School*. 1945. 3:30-33
7. Kokhanova A.N., Peresyphkina N.V., Balagurovskaya E.N., Kotel'nikova K.M., Grigorieva I.T., Nechaev S.K.. The effect of Schizandra on muscular fatigue in man. *Proceedings of the 4th Student Scientific Session of the Khabarovsk Medical Institute*. 1950. : 85-89
8. Fil'kin A.M.. About . *Aptech. Delo*. 1952. 2:46-48
9. Zemlinskij S.E., Zemlinskij S.E.. Schizandra chinensis (Turcz.) Baill. *Lekarstvennye Rasitenija SSSR*. 1958. :57-61
10. Gammerman A.F., Kadaev G.N.. *Medicianl Plants*. 1975. :142-143
11. Lazarev N.V.. General and specific in action of pharmacological agents. *Farmacol.Toxicol.*. 1958. 21:81-86
12. Lazarev N.V., Ljublina E.I., Rozin M.A.. State of nonspecific resistance. *Patol. Fiziol. Experim. Terapia*. 1959. 3:16-21
13. Selye H.. *Stress*. 1950
14. Brekhman I.I., Dardymov I.V.. New substances of plant origin which increase nonspecific resistance. *Ann. Rev. Pharmacol.*. 1968. 8:419-430 doi:10.1146/annurev.pa.09.040169.002223
15. Meerson F.. *Adaptation, Stress and Prophylax*. 1981
16. Brekhman I.I.. *Eleutherococc (Bibliographic index) 1958-1981*. 1982. :1-14
17. Wiegant F.A., Surinova S., Ytsma E., Langelaar-Makkinje M., Wikman G., Post J.A.. Plant adaptogens increase lifespan and stress resistance in *C. elegans*. *Biogerontology*. 2009. 10:27-42 doi:10.1007/s10522-008-9151-9
18. Wiegant F.A.C., Limandjaja G., de Poot S.A.H., Bayda L.A., Vorontsova O.N., Zenina T.A., Langelaar Makkinje M., Post J.A., Wikman G., Lukyanova L., Takeda N., Singal P.K.. Plant adaptogens activate cellular adaptive mechanisms by causing mild damage. *Adaptation Biology and Medicine: Health Potentials*. 2008. Volume 5:319-332
19. Jafari M., Felgner J.S., Bussel I.I., Hutchili T., Khodayari B., Rose M.R., Vince-Cruz C., Mueller L.D.. Rhodiola: a promising anti-aging Chinese herb. *Rejuvenation Res.*. 2007. 10:587-602 doi:10.1089/rej.2007.0560
20. Schriener S.E., Avanesian A., Liu Y., Luesch H., Jafari M.. Protection of human cultured cells against oxidative stress by Rhodiola rosea without activation of antioxidant defenses. *Free Radic. Biol. Med.*. 2009. 47:577-584 doi:10.1016/j.freeradbiomed.2009.05.025
21. Panossian A., Hambartsumyan M., Hovanissian A., Gabrielyan E., Wikman G.. The Adaptogens Rhodiola and Schizandra Modify the Response to Immobilization Stress in Rabbits by Suppressing the Increase of Phosphorylated Stress-activated Protein Kinase, Nitric Oxide and Cortisol. *Drug Targets Insights*. 2007. 1:39-54
22. Panossian A., Wikman G., Kaur P., Asea A.. Adaptogens exert a stress protective effect by modulation of expression of molecular chaperons. *Phytomedicine*. 2009. 16:617-622 doi:10.1016/j.phymed.2008.12.003
23. Panossian A., Wikman G.. Evidence-based efficacy of adaptogens in fatigue, and molecular mechanisms related to their stress-protective activity. *Current Clin.Pharmacol.*. 2009. 4:198-219 doi:10.2174/157488409789375311
24. Panossian A., Wikman G., Bonn K.. Evidence-based efficacy of adaptogens in fatigue and molecular mechanisms related to their stress-protective activity. *International Evidence-Based Complementary Medicine Conference*. 2009. :10 doi:10.2174/157488409789375311
25. Committee on Medicinal Products
26. Panossian A., Wikman G.. Pharmacology of . *J. Ethnopharmacol.*. 2008. 118:183-212 doi:10.1016/j.jep.2008.04.020
27. Dardimov I.V.. *Ginseng, Eleutherococcus. On the Mechanism of Biological Activity*. 1976. :1-189
28. Saratikov A.S., Krasnov E.A.. *Rhodiola rosea (Golden root)*. 2004. :1-286
29. Lebedev A.A.. *Schizandra*. 1971. :1-98
30. Lupandin A.V., Lapaev I.I.. *Schisandra*. 1981. :1-89
31. Farnsworth N., Kinghorn A.D., Soejarto D.D., Waller D.P., Wagner H., Hikino H., Farnsworth N.R.. Siberian Ginseng (. *Economic and Medicinal Plant Research*. 1985. Volume 1:156-209
32. Brekhman I.I.. *Eleutherococcus*. 1968. :1-168
33. Brown R.P., Gerbarg P.L., Graham B.. *The Rhodiola revolution*. 2004. :1-260
34. Winston D., Maimes S.. *Adaptogens. Herbs for strength, stamina and stress relief*. 2007. :1-324

35. Kurkin V.A., Zapesochneya G.G.. Chemical composition and pharmacological properties of *Rhodiola rosea* L. *Khim.Farm. Zhurnal*. 1986. 20:1231-1244
36. Khanum F., Bawa A.S., Singh B.. *Compr. Rev. Food Sci. Food Safety*. 2005. 4:55-62 doi:10.1111/j.1541-4337.2005.tb00073.x
37. Kelly G.S.. *Rhodiola rosea*: a possible plant adaptogen. *Altern. Med. Rev.*. 2001. 6:293-302
38. Zapesochneya G.G., Kurkin V.A., Boyko V.P., Kolkhir V.K.. Phenylpropanoids—promising biologically active compounds of medicinal plants. *Khim.Farm. Zh.*. 1995. 29:47-50
39. Aksenova R.A., Zotova M.I., Nekhoda M.F., Cherdintsev S.G., Saratikov A.S.. Comparative characteristics of the stimulating and adaptogenic effects of . *Stimulants of the central nervous system*. 1968. Volume 2:3-12
40. Panossian A., Nikoyan N., Ohanyan N., Hovhannisyan A., Abrahamyan H., Gabrielyan E., Wikman G.. Comparative study of *Rhodiola* preparations on behavioral despair of rats. *Phytomedicine*. 2008. 15:84-91 doi:10.1016/j.phymed.2007.10.003
41. Barnaulov O.D., Limarenko A.Y., Kurkin V.A., Zapesochneya G.G., Shchavilinskij A.N.. A comparative evaluation of the biological activity of compounds isolated from species of *Rhodiola*. *Khim.Pharm. Zh.*. 1986. 23:1107-1112
42. Sokolov S.Y., Boyko V.P., Kurkin V.A., Zapesochneya G.G., Rvantsova N.V., Grinenko H.A.. A comparative study of the stimulant property of certain phenylpropanoids. *Khim. Pharm. Zh.*. 1990. 24:66-68
43. Sokolov S.Y., Ivashin V.M., Zapesochneya G.G., Kurkin V.A., Shavilinskiy A.N.. Study of neurotropic activity of new substances isolated from *Rhodiola rosea*. *Khim. Pharm. Zh.*. 1985. 19:1367-1371
44. Ovodov Y.S., Ovodova R.G., Solovyova T.F., Elyakov G.B., Kochetkov N.K.. Glycosides of *Eleutherococcus senticosus*. I. Isolation and some properties of Eleutherosides B and E. *Khim. Prir. Soedin.*. 1965. 1:3-7
45. Ovodov Y.S., Frolova G.M., Nefedova M.Y., Elyakov G.B.. Glycosides of *Eleutherococcus senticosus*. II. Structure of Eleutherosides A, B1, C and D. *Khim. Prir. Soedin.*. 1965. 1:63-64
46. Elyakov G.B., Ovodov Y.S.. The glycosides of Araliaceae. *Khim. Prir. Soedin.*. 1972. 6:697-709
47. Norr H.. Phytochemical and pharmacological Investigations of the Adaptogens: *Eleutherococcus senticosus*, *Ocimum sanctum*, *Codonopsis pilosula*, *Rhodiola rosea* and *Rhodiola crenulata*. *Ph.D. Dissertation*. 1993. :1-231
48. Wagner H., Amason J.T.. Immunostimulants and adaptogens from plants. *Phytochemistry of Medicinal Plants*. 1995. :1-18
49. Balandin D.A., Lazarev N.V.. Schizandrin—a new stimulant from *Schizandra* fruits. *Materials for the Study of Stimulants and Tonics from Ginseng and Schizandra Roots*. 1951. :45-50
50. Lebedev A.A., Lazarev N.V.. Some materials on the pharmacology of schizandrin. *Materials for the Study of Stimulants and Tonics from Ginseng and Schizandra Roots*. 1951. :109-117
51. Lebedev A.A.. *Schizandrine –a New Stimulant from Schizandra Chinensis Fruits—Summary of Thesis for a Candidate's degree in Medicine*. 1967. :1-21
52. Kochetkov N.K., Khorlin A.Y., Chizhov O.S., Sheichenko V.I.. Chemical investigation of . *Izv. Akad. Nauk. Ser. Khim.*. 1962. 5:850-856
53. Kochetkov N.K., Khorlin A.Y., Chizhov O.S., Sheichenko V.I.. Schizandrin, a lignan of unusual structure. *Tetrahedron Lett.*. 1961. 31:730-734
54. Kim S.R., Lee M.K., Koo K.A., Kim S.H., Sung S.H., Lee N.G., Markelonis G.J., Oh T.H., Yang J.H., Kim Y.C.. Dibenzocyclooctadiene lignans from *Schisandra chinensis* protect primary cultures of rat cortical cells from glutamate-induced toxicity. *J. Neurosci. Res.*. 2004. 76:397-405 doi:10.1002/jnr.20089
55. Wang B., Wang X.M.. Schisandrin B protects rat cortical neurons against Abeta1-42-induced neurotoxicity. *Pharmazie*. 2009. 64:450-454
56. Ko K.M., Chen N., Leung H.Y., Leong E.P., Poon M.K., Chiu P.Y.. Long-term schisandrin B treatment mitigates age-related impairments in mitochondrial antioxidant status and functional ability in various tissues, and improves the survival of aging C57BL/6J mice. *Biofactors*. 2008. 34:331-342 doi:10.1002/biof.5520340408
57. Liu X.N., Zhang C.Y., Jin X.D., Li Y.Z., Zheng X.Z., Li L.. Inhibitory effect of schisandrin B on gastric cancer cells in vitro. *World J. Gastroenterol.*. 2007. 13:6506-6511 doi:10.3748/wjg.13.6506
58. Chiu P.Y., Leung H.Y., Poon M.K., Mak D.H., Ko K.M.. (–)Schisandrin B is more potent than its enantiomer in enhancing cellular glutathione and heat shock protein production as well as protecting against oxidant injury in H9c2 cardiomyocytes. *Mol. Cell. Biochem.*. 2006. 289:185-191 doi:10.1007/s11010-006-9163-1
59. Chen N., Chiu P.Y., Ko K.M.. Schisandrin B enhances cerebral mitochondrial antioxidant status and structural integrity, and protects against cerebral ischemia/reperfusion injury in rats. *Biol. Pharm. Bull.*. 2008. 31:1387-1391 doi:10.1248/bpb.31.1387
60. Pao T.T., Hsu K.F., Liu K.T., Chang L.G., Chuang C.H., Sung C.Y.. Protective action of schizandrin B on hepatic injury in mice. *Chin Med J (Engl.)*. 1977. 3:173-179

61. Panossian A., Gabrielian E., Wagner H.. On the mechanism of action of plant adaptogens with particular references on cucurbitacin R diglucoside. *Phytomedicine*. 1999. 6:147-155 doi:10.1016/S0944-7113(99)80002-6
62. Panossian A., Gabrielian E., Wagner H.. Plant adaptogens II: Bryonia as an adaptogen. *Phytomedicine*. 1997. 4:83-97 doi:10.1016/S0944-7113(97)80033-5
63. Wang L.C., Lee T.F.. Effect of ginseng saponins on cold tolerance in young and elderly rats. *Planta Med.* 2000. 66:144-147 doi:10.1055/s-2000-11122
64. Hiai S., Yokoyama H., Oura H., Yano S.. Stimulation of pituitary-adrenocortical system by ginseng saponin. *Endocrinol. Jpn.* 1979. 26:661-665 doi:10.1507/endocrj1954.26.661
65. Hiai S., Yokoyama H., Oura H.. Features of ginseng saponin-induced corticosterone secretion. *Endocrinol. Jpn.* 1979. 26:737-740 doi:10.1507/endocrj1954.26.737
66. Kim D.H., Jung J.S., Suh H.W., Huh S.O., Min S.K., Son B.K., Park J.H., Kim N.D., Kim Y.H., Song D.K.. Inhibition of stress-induced plasma corticosterone levels by ginsenosides in mice: involvement of nitric oxide. *Neuroreport*. 1998. 9:2261-2264 doi:10.1097/00001756-199807130-00021
67. Kim D.H., Moon Y.S., Jung J.S., Min S.K., Son B.K., Suh H.W., Song D.K.. Effects of ginseng saponin administered intraperitoneally on the hypothalamo-pituitary-adrenal axis in mice. *Neurosci. Lett.* 2003. 343:62-66 doi:10.1016/S0304-3940(03)00300-8
68. Kim D.H., Jung J.S., Moon Y.S., Sung J.H., Suh H.W., Kim Y.H., Song D.K.. Inhibition of intracerebroventricular injection stress-induced plasma corticosterone levels by intracerebroventricularly administered compound K, a ginseng saponin metabolite, in mice. *Biol. Pharm. Bull.* 2003. 26:1035-1038 doi:10.1248/bpb.26.1035
69. Ghosal S., Lal J., Srivastava R., Battacharya R., Upadhyay S.N., Jaiswal A.K., Chattopadhyay U.. Anti-stress activity of sitoindosides IX and X, New C-27-glycowithanolides from . *Phytother. Res.* 1989. 3:201-209 doi:10.1002/ptr.2650030510
70. Lee K.-H., Kwak J.H., Lee K.-B., Song K.-S.. Prolyl endopeptidase inhibitors from caryophylli flos. *Arch. Pharmacol. Res. (Seoul)*. 1998. 21:207-211 doi:10.1007/BF02974029
71. Baek N.I., Han J.T., Ahn E.M., Park J.K., Cho S.W., Jeon S.G., Jang J.S., Kim Choong, K., Choi S.Y.. Isolation of anticonvulsant compounds from the fruits of *Schizandra chinensis* Baill. *Agric. Chem. Biotechnol.* 2000. 43:72-77
72. Munck A., Guyre P.M., Holbrook N.J.. Physiological functions of glucocorticoids in stress and their relation to pharmacological actions. *Endocr. Rev.* 1984. 5:25-44 doi:10.1210/edrv-5-1-25
73. Fink G.. *Encyclopedia of Stress*. 2000. Volumes 1–3
74. Chrousos G.P., Gold P.W.. The concept of stress system disorders: Overview of behavioral and physical homeostasis. *JAMA*. 1992. 267:1244-1252 doi:10.1001/jama.1992.03480090092034
75. Van Diermen D., Marston A., Bravo J., Reist M., Carrupt P.A., Hostettmann K.. Monoamine oxidase inhibition by *Rhodiola rosea* L. roots. *J. Ethnopharmacol.* 2009. 122:397-401 doi:10.1016/j.jep.2009.01.007
76. Saratikov A.S., Krasnov E.A., Chnikina L.A., Duvidson L.M., Sotova M.I., Marina T.F., Nechoda M.F., Axenova R.A., Tscherdinzeff S.G.. Rhodiolosid, a new glycoside from *Rhodiola rosea* and its pharmacological properties. *Pharmazie*. 1968. 23:392-395
77. Panossian A., Wagner H.. Stimulating effect of adaptogens: an overview with particular reference to their efficacy following single dose administration. *Phytother. Res.* 2005. 19:819-838 doi:10.1002/ptr.1751
78. Panossian A., Wikman G.. Effect of adaptogens on the central nervous system. *Arq. Bras. Fitomed. Cient.* 2005. 2:108-130
79. Cao L.L., Du G.H., Wang M.W.. Effect of salidroside on mitochondria injury induced by sodium azide. *Yao Xue Xue Bao*. 2005. 40:700-704
80. Cao L.L., Du G.H., Wang M.W.. Effect of salidroside on cell damage induced by glutamate and intracellular free calcium in PC12 cells. *J. Asian Nat.Prod. Res.* 2006. 8:159-165 doi:10.1080/1028602042000325645
81. Chen X., Liu J., Gu X., Ding F.. Salidroside attenuates glutamate-induced apoptotic cell death in primary cultured hippocampal neurons of rats. *Brain Res.* 2008. 1238:189-198 doi:10.1016/j.brainres.2008.07.051
82. Cai L., Wang H., Li Q., Qian Y., Yao W.. Salidroside inhibits H₂O₂-induced apoptosis in PC12 cells by preventing cytochrome c release and inactivating of caspase cascade. *Acta Biochim. Biophys. Sin. (Shanghai)*. 2008. 40:796-802 doi:10.1093/abbs/40.9.796
83. Zhang W.S., Zhu L.Q., Niu F.L., Deng R.C., Ma C.X.. Protective effects of salidroside on injury induced by hypoxia/hypoglycemia in cultured neurons. *Zhongguo Zhong Yao Za Zhi*. 2004. 29:459-462
84. Zhang L., Yu H., Sun Y., Lin X., Chen B., Tan C., Cao G., Wang Z.. Protective effects of salidroside on hydrogen peroxide-induced apoptosis in SH-SY5Y human neuroblastoma cells. *Eur. J. Pharmacol.* 2007. 564:18-25 doi:10.1016/j.ejphar.2007.01.089

85. Jang S.I., Pae H.O., Choi B.M., Oh G.S., Jeong S., Lee H.J., Kim H.Y., Kang K.J., Yun Y.G., Kim Y.C., Chung H.T.. Salidroside from *Rhodiola sachalinensis* protects neuronal PC12 cells against cytotoxicity induced by amyloid-beta. *Immunopharmacol. Immunotoxicol.*. 2003. 25:295-304 doi:10.1081/IPH-120024498
86. Li H.B., Ge Y.K., Zheng X.X., Zhang L.. Salidroside stimulated glucose uptake in skeletal muscle cells by activating AMP-activated protein kinase. *Eur.J. Pharmacol.*. 2008. 588:165-169 doi:10.1016/j.ejphar.2008.04.036
87. Ma L., Cai D.L., Li H.X., Tong B.D., Wang Y., Pei S.P.. Protective effects of salidroside on oxidative damage in fatigue mice. *Zhong Xi Yi Jie He Xue Bao*. 2009. 7:237-241 doi:10.3736/jcim20090308
88. Zou Y.Q., Cai Z.Y., Mao Y.F., Li J.B., Deng X.M.. Effects of salidroside-pretreatment on neuroethology of rats after global cerebral ischemia-reperfusion. *Zhong Xi Yi Jie He Xue Bao*. 2009. 7:130-134 doi:10.3736/jcim20090207
89. Yu S., Liu M., Gu X., Ding F.. Neuroprotective effects of salidroside in the PC12 cell model exposed to hypoglycemia and serum limitation. *Cell. Mol. Neurobiol.*. 2008. 28:1067-1078 doi:10.1007/s10571-008-9284-z
90. Samuel S.M., Thirunavukkarasu M., Penumathsa S.V., Paul D., Maulik N.. Akt/FOXO3a/SIRT1-mediated cardioprotection by n-tyrosol against ischemic stress in rat in vivo model of myocardial infarction: switching gears toward survival and longevity. *J Agric. Food Chem.*. 2008. 56:9692-9698 doi:10.1021/jf802050h
91. Owasojo J.O., Neri D.F., Lamberth J.G.. Tyrosine and its potential use as a countermeasure to performance decrement in military sustained operations. *Aviat. Space Environ. Med.*. 1992. 63:364-369
92. Kelly G.S.. Nutritional and botanical investigations to assist with the adaptation to stress. *Altern. Med. Rev.*. 1999. 4:249-265
93. Ko K.M., Lam B.Y.. Schisandrin B protects against tert-butylhydroperoxide induced cerebral toxicity by enhancing glutathione antioxidant status in mouse brain. *Mol.Cell. Biochem.*. 2002. 238:181-186 doi:10.1023/A:1019907316129
94. Xue J.Y., Liu G.T., Wei H.L., Pan Y.. Antioxidant activity of two dibenzocyclooctene lignans on the aged and ischemic brain in rats. *Free Radic. Biol. Med.*. 1992. 12:127-135 doi:10.1016/0891-5849(92)90006-3
95. Chiu P.Y., Tang M.H., Mak D.H., Poon M.K., Ko K.M.. Hepatoprotective mechanism of schisandrin B: role of mitochondrial glutathione antioxidant status and heat shock proteins. *Free Radic. Biol. Med.*. 2003. 35:368-380 doi:10.1016/S0891-5849(03)00274-0
96. Chiu P.Y., Mak D.H., Poon M.K., Ko K.M.. Role of cytochrome P-450 in schisandrin B-induced antioxidant and heat shock responses in mouse liver. *Life Sci.*. 2005. 77:2887-2895 doi:10.1016/j.lfs.2005.03.026
97. Chiu P.Y., Leung H.Y., Poon M.K., Ko K.M.. Chronic schisandrin B treatment improves mitochondrial antioxidant status and tissue heat shock protein production in various tissues of young adult and middle-aged rats. *Biogerontology*. 2006. 7:199-210 doi:10.1007/s10522-006-9017-y
98. Chiu P.Y., Ko K.M.. Schisandrin B protects myocardial ischemia-reperfusion injury partly by inducing Hsp25 and Hsp70 expression in rats. *Mol. Cell. Biochem.*. 2004. 266:139-144 doi:10.1023/B:MCBI.0000049151.79238.30
99. Ip S.P., Che C.T., Kong Y.C., Ko K.M.. Effects of schisandrin B pretreatment on tumor necrosis factor-alpha induced apoptosis and Hsp70 expression in mouse liver. *Cell Stress Chaperones*. 2001. 6:44-48 doi:10.1379/1466-1268(2001)006<0044:E0SBPO>2.0.CO;2
100. Ko K.M., Chiu P.Y.. Structural determinants of schisandrin B which enhance mitochondrial functional ability and glutathione status as well as heat shock protein expression in rat hearts and H9c2 cells. *Mol. Cell. Biochem.*. 2005. 276:227-234 doi:10.1007/s11010-005-4539-1
101. Tang M.H., Chiu P.Y., Ko K.M.. Hepatoprotective action of schisandrin B against carbon tetrachloride toxicity was mediated by both enhancement of mitochondrial glutathione status and induction of heat shock proteins in mice. *Biofactors*. 2003. 19:33-42 doi:10.1002/biof.5520190105
102. Chiu P.Y., Leung H.Y., Ko K.M.. Schisandrin B Enhances Renal Mitochondrial Antioxidant Status, Functional and Structural Integrity, and Protects against Gentamicin-Induced Nephrotoxicity in Rats. *Biol. Pharm. Bull.*. 2008. 31:602-605 doi:10.1248/bpb.31.602
103. Chiu P.Y., Leung H.Y., Poon M.K., Lee S.S., Ko K.M.. Schisandrin B induced antioxidant response is partly mediated by cytochrome P-450E1 catalyzed reaction in mouse liver. *Mol. Cell. Biochem.*. 2006. 293:87-92 doi:10.1007/s11010-006-2957-3
104. Lee I.S., Jung K.Y., Oh S.R., Park S.H., Ahn K.S., Lee H.K.. Structure-activity relationships of lignans from *Schisandra chinensis* as platelet activating factor antagonists. *Biol. Pharm.Bull.*. 1999. 22:265-267 doi:10.1248/bpb.22.265
105. Mak D.H., Ip S.P., Li P.C., Poon M.K., Ko K.M.. Effects of Schisandrin B and alpha-tocopherol on lipid peroxidation, in vitro and in vivo. *Mol. Cell. Biochem.*. 1996. 16:161-165 doi:10.1007/BF00229479
106. Mak DH, Ko KM.. Alterations in susceptibility to carbon tetrachloride toxicity and hepatic antioxidant/detoxification system in streptozotocin-induced short-term diabetic rats: effects of insulin and Schisandrin B treatment. *Mol. Cell. Biochem.*. 1997. 175:225-232 doi:10.1023/A:1006883919687
107. Lu H., Liu G.T.. Effect of dibenzo[a,c]cyclooctene lignans isolated from *Fructus schizandrae* on lipid peroxidation and anti-oxidative enzyme activity. *Chem.Biol. Interact.*. 1991. 78:77-84 doi:10.1016/0009-2797(91)90104-F

108. Zhao B.L., Li X.J., Liu G.T., Jia W.Y., Xin W.J.. Scavenging effect of schizandrins on active oxygen radicals. *Cell Biol Int Rep.* 1990. 14:99-109 doi:10.1016/0309-1651(90)90027-v
109. Li X.J., Zhao B.L., Liu G.T., Xin W.J.. Scavenging effects on active oxygen radicals by schizandrins with different structures and configurations. *Free Radic. Biol. Med.* 1990. 9:99-104 doi:10.1016/0891-5849(90)90111-u
110. Chiu P.Y., Ko K.M.. Time-dependent enhancement in mitochondrial glutathione status and ATP generation capacity by schisandrin B treatment decreases the susceptibility of rat hearts to ischemia-reperfusion injury. *Biofactors.* 2003. 19:43-51 doi:10.1002/biof.5520190106
111. Pan S.Y., Han Y.F., Carlier P.R., Pang Y.P., Mak D.H., Lam B.Y., Ko K.M.. Schisandrin B protects against tacrine-and bis(7)-tacrine-induced hepatotoxicity and enhances cognitive function in mice. *Planta Med.* 2002. 68:217-220 doi:10.1055/s-2002-23145
112. Panossian A., Wikman G., Borelli F., Capasso F., Milic N., Russo A.. Plant adaptogens: historical overview and perspectives. *3rd International Symposium on Natural Drugs. Proceedings.* 2003. :205-209
113. Panossian A.. Adaptogens: Tonic herbs for fatigue and stress. *Alt.Comp.Therap.* 2003. 9:327-332 doi:10.1089/107628003322658610
114. Chernykh S.I., Lukhtanov V.A., Simonenko N.P.. Adaptation to damage in the silkworm . *Entomologicheskoye Obozrenye.* 1985. 2:267-272
115. Boon-Niermeijer E.K., van den Berg A., Wikman G., Wiegant F.A.C.. Phytoadaptogens protect against environmental stress-induced death of embryos from the freshwater snail . *Phytomedicine.* 2000. 7:389-400 doi:10.1016/S0944-7113(00)80060-4
116. Kunitomo S., Murofushi W., Kai H., Ishida Y., Uchiyama A., Kobayashi T., Kobayashi S., Murofushi H., Murakami-Murofushi K.. Steryl glucoside is a lipid mediator in stress-responsive signal transduction. *Cell Struct. Funct.* 2002. 27:157-162 doi:10.1247/csf.27.157
117. Lishmanov Yu.B., Krylatov A.V., Maslov L.N., Nariznaya N.V., Zamotinkii A.V.. Effect of Rhodiola rosea on the level of inducible Hsp-70 in miocard in stress. *Bull. Exp. Biol. Med.* 1996. 121:256-258 doi:10.1007/BF02446762
118. Prodius P.A., Manukhina E.B., Bulanov A.E., Wikman G., Malyshev I.I.. Adaptogen ADAPT modulates synthesis of inducible stress protein HSP 70 and increases organism resistance to heat shock. *Biull. Eksp. Biol. Med.* 1997. 123:629-633
119. Panossian A., Wikman G., Andreeva L., Boykova A., Nikiforova D., Timonina N.. Adaptogens exert a stress protective effect by modulation of expression of molecular chaperons. *Planta Med.* 2008. 74:1018
120. Lee F.T., Kuo T.Y., Liou S.Y., Chien C.T.. Chronic Rhodiola rosea extract supplementation enforces exhaustive swimming tolerance. *Am. J. Chin. Med.* 2009. 37:557-572 doi:10.1142/S0192415X09007053
121. Jung C.H., Jung H., Shin Y.C., Shin Y.C., Park J.H., Jun C.Y., Kim H.M., Yim H.S., Shin M.G., Bae H.S., Kim S.H., Ko S.G.. *J. Ethnopharmacol.* 2007. 113:183-187 doi:10.1016/j.jep.2007.05.023
122. Olsson E.M.G., von Schéele B., Panossian A.G.. A randomized double-blind placebo controlled parallel group study of SHR-5 extract of Rhodiola rosea roots as treatment for patients with stress related fatigue. *Planta Med.* 2009. 75:105-112 doi:10.1055/s-0028-1088346
123. Filaretov A.A., Bogdanova T.S., Mitushov M.I., Podvigina T.T., Srailova G.T.. Effect of adaptogens on the activity of the pituitary-adrenocortical system in rats. *Biull. Eksp. Biol. Med.* 1986. 101:573-574
124. Kimura Y., Sumiyoshi M.. Effects of various Eleutherococcus senticosus cortex on swimming time, natural killer activity and corticosterone level in forced swimming stressed mice. *J. Ethnopharmacol.* 2004. 95:447-453 doi:10.1016/j.jep.2004.08.027
125. Lishmanov Iu.B., Trifonova Zh.V., Tsibin A.N., Maslova L.V., Dement'eva L.A.. Plasma beta-endorphin and stress hormones in stress and adaptation. *Biull. Eksp. Biol. Med.* 1987. 103:422-424
126. Maslov L.N., Lishmanov Iu.B., Naumova A.V., Lasukova T.V.. Do endogenous ligands of peripheral mu-and delta-opiate receptors mediate anti-arrhythmic and cardioprotective effects of Rhodiola rosea extract?. *Biull. Eksp. Biol. Med.* 1997. 124:151-153
127. Maimeskulova L.A., Maslov L.N., Lishmanov Iu.B., Krasnov E.A.. The participation of the mu-, delta-, and kappa-opioid receptors in the realization of the anti-arrhythmia effect of Rhodiola rosea. *Eksp. Klin. Farmakol.* 1997. 60:38-39
128. Panossian A., Oganessian A., Ambartsumian M., Gabrielian E., Wagner H., Wikman G.. Effects of heavy physical exercise and adaptogens on nitric oxide content in human saliva. *Phytomedicine.* 1999. 6:17-26 doi:10.1016/S0944-7113(99)80030-0
129. Park E.K., Shin Y.W., Lee H.U., Kim S.S., Lee Y.C., Lee B.Y., Kim D.H.. Inhibitory effect of ginsenoside Rb1 and compound K on NO and prostaglandin E. *Biol. Pharm. Bull.* 2005. 28:652-666 doi:10.1248/bpb.28.652
130. Park Y.C., Lee C.H., Kang H.S., Kim K.W., Chung H.T., Kim H.D.. Ginsenoside-Rh1 and Rh2 inhibit the induction of nitric oxide synthesis in murine peritoneal macrophages. *Biochem. Mol.Biol. Int.* 1996. 40:751-757 doi:10.1080/15216549600201353
131. Abidov M., Crendal F., Grachev S., Seifulla R., Ziegenfuss T.. Effect of extracts from . *Bull. Exp. Biol. Med.* 2003. 136:585-587 doi:10.1023/B:BEBM.0000020211.24779.15

132. Wagner H., Norr H., Winterhoff H.. Plant adaptogens. *Phytomedicine*. 1994. 1:63-76 doi:10.1016/S0944-7113(11)80025-5
133. Panossian A., Wikman G., Wagner H.. Plant adaptogens III: Earlier and more recent aspects and concepts on their modes of action. *Phytomedicine*. 1999. 6:287-300 doi:10.1016/S0944-7113(99)80023-3
134. Darbinyan V., Aslanyan G., Amroyan E., Gabrielyan E., Malmström C., Panossian A.. Clinical trial of . *Nord. J. Psychiatry*. 2007. 61:2343-2348 doi:10.1080/08039480701643290
135. Bocharov E.V., Kucherianu V.G., Bocharova O.A., Karpova R.V.. Neuroprotective features of phytoadaptogens. *Vestn. Ross. Akad. Med. Nauk*. 2008. 4:47-50
136. Fulder S.. The drug that builds Russians. *New Scientist*. 1980. 21:83-84
137. Barenboim G.M., Kozlova N.B., Barenboim G.M.. Application of Eleutherococcus extract for increasing of biological resistance of humans under influence of different environmental noxious factors. *Eleutherococc. Strategy of Use and New Data of Fundamental Investigations*. 1985. :7-20
138. Sarris J.. Herbal medicines in the treatment of psychiatric disorders: A systematic review. *Phytother. Res.*. 2007. 21:703-716 doi: 10.1002/ptr.2187
139. Bucci L.R.. Selected herbals and human exercise performance. *Am. J. Clin. Nutr.*. 2000. 72:624S-636S doi:10.1093/ajcn/72.2.624S
140. Walker T.B., Robergs R.A.. Does Rhodiola rosea possess ergogenic properties?. *Int. J. Sport Nutr. Exerc. Metab.*. 2006. 16:305-315 doi: 10.1123/ijsnem.16.3.305
141. Bu Y., Jin Z.H., Park S.Y., Baek S., Rho S., Ha N., Park S.K., Kim H., Sun Y.K.. Siberian ginseng reduces infarct volume in transient focal cerebral ischaemia in Sprague-Dawley rats. *Phytother. Res.*. 2005. 19:167-169 doi:10.1002/ptr.1649
142. Hartz A.J., Bentler S., Noyes R., Hoehns J., Logemann C., Sinift S., Butani Y., Wang W., Brake K., Ernst M., Kautzman H.. Randomized controlled trial of Siberian Ginseng for chronic fatigue. *Psychol. Med.*. 2004. 34:51-61 doi:10.1017/S0033291703008791
143. Darbinyan V., Kteyan A., Panossian A., Gabrielian E., Wikman G., Wagner H.. 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. 7:365-371 doi:10.1016/S0944-7113(00)80055-0
144. Spasov A.A., Wikman G.K., Mandrikov V.B., Mironova I.A., Neumoin V.V.. A double-blind, placebo-controlled pilot study of the stimulating and adaptogenic effect of . *Phytomedicine*. 2000. 7:85-89 doi:10.1016/S0944-7113(00)80078-1
145. Shevtsov V.A., Zhohus B.I., Shervarly V.I., Vol'skij V.B., Korovin Y.P., Khristich M.P., Roslyakova N.A., Wikman G.. A randomized trial of two different doses of a SHR-5 Rhodiola roseaextract versus placebo and control of capacity for mental work. *Phytomedicine*. 2003. 10:95-105 doi:10.1078/094471103321659780
146. Narimanian M., Badalyan M., Panosyan V., Gabrielyan E., Panossian A., Wikman G., Wagner H.. Impact of Chisan. *Phytomedicine*. 2005. 12:723-772 doi:10.1016/j.phymed.2004.11.004
147. Bogatova R.I., Malozemov V.V.. *Experimental Research on the Estimation of Influence of Single Dose of Phytoadaptogens on Short-Term Memory. Report on ADAPT 232*. 1994. :1-151
148. Bogatova R.I., Shlyakova L.V., Salnitsky V.P., Wikman G.. Evaluation of the effect of single administration of a phytoadaptogen on human subject work ability during long isolation. *Aerospace Environ. Med.*. 1997. 31:51-54
149. Berdyshev V.V., Breikhman I.I.. Some aspects of single doses of adaptogens (Eleutherococcus and Schizandra). *Valeology: Diagnosis, Means and Practice in Health Care, International Collection of Scientific Papers*. 1995. :105-117
150. Kaliko I.M., Tarasova A.A., Saratikov A.S.. Effect of Leuzea and Golden Root extracts on dynamic peculiarities of the highest neural performance. *Stimulants of the Central Nervous System*. 1966. :115-120
151. Spasov A.A., Mandrikov V.B., Mironova I.A.. The effect of the preparation rhodosin on the psychophysiological and physical adaptation of students to an academic load. *Eksp. Klin. Farmakol.*. 2000. 63:76-78
152. Roslyakova N.A., Bogatova R.I., Verishvili M.O., Wikman G.. The effect of single dose of Rodelim phytoadaptogen on the performance of operators under intensive activity. *Abstract Book of Scientific Conference Biologically Active Food Supplements and Natural Medicines in Prophylaxis, Treatment and Rehabilitation*. :157-160
153. Zotova M.I.. The effect of . *Collection of Reports at 3rd Scientific Conference of Physiologists*. 1965. :298-299
154. Komar V.V., Kit S.M., Sitschuk L.V., Sitschuk V.M.. The effect of Carpatian . *Farm. Zh.*. 1981. 4:62-64
155. Marina T.F., Mikhaleva L.K., Suslov N.I.. Comparative effects of para-tyrosol and Rhodiola extract on the central nervous system. *Proceedings of a Joint Plenary Session of Pathophysiologists and Pharmacologists of Siberia and the Far East on Mechanisms of the Development of Pathological Processes*. :66-68
156. Strokina T.I., Brekhman I.I. The changes in higher nervous activity in neurotic patients under treatment with Eleutherococcus. *Lekstvennie Sredstva Dalnego Vostoka*. 1967. Volume 7:201-111

157. Medvedev M.A., Brekhman I.I., Belikov I.F., Kurentsova G.E.. Effect of Ginseng and Eleutherococcus on working parameters of radio-telegraph operators. *Materials of Studies of Ginseng and other Medicinal Plants of Far East*. 1963. :237-239
158. Winther K., Ranlov C., Rein E., Mehlsen J.. Russian root (Siberian Ginseng) improves cognitive functions in middle-aged people, whereas . *J. Neurol. Sci.*. 1997. 150:90
159. Gubchenko P.P., Frumentov N.K., Bulanov A.E., Dardimov I.V., Li S.E.. A comparative study of stimulating effect of adaptogenic agents prepared from some far-eastern plants. *New Data on Eleutherococcus and Other Adaptogens*. 1986. :18-25
160. Gubchenko P.P., Frumentov N.K., Brekhman I.I., Dardimov I.V., Li S.E., Dobryakova A.I.. A comparative study of effectiveness of . *New Data on Eleutherococcus: Proceedings of the 2nd International Symposium on Eleutherococcus*. 1986. :171-178
161. Grigorenko G.F., Berdishev V.V.. The use of tonic drinks and drugs increasing working capacity of sailors during night shifts. *Abstract Book of the Scientific Practical Conference on Medical and Social Aspects of the "Man-Ocean" Problem*. :29-30
162. Kochmareva L.I., Belikov I.F., Brekhman I.I., Bykov V.T., Lazarev N.V., Serebryannik B.E., Sorokhtin G.N.. The effect of . *Materials for the Study of Ginseng and Schizandra*. 1958. :12-17
163. Galant I.B., Belikov I.F., Brekhman I.I., Bykov V.T., Lazarev N.V., Serebryannik B.E., Sorokhtin G.N.. Curative effect of . *Materials for the Study of Ginseng and Schizandra*. 1958. :198-208
164. Leman M.F.. Treatment of reactive and asthenic states of exogenous etiology using the Far East Schizandra. *J. Neuropathol.Psychiatry*. 1952. 52:67-70
165. Rossijskij D.M.. *Vrachebn. Delo*. 1952. 10:939-940
166. Rossijskij D.M.. New tonics and stimulants produced from local medicinal herbs. *Clin. Med.*. 1952. 30:66-69
167. Zakharova P.M., Krasnushkin E.K.. Experience of using . *Problems of Social and Clinical Psychiatry*. 1948. Volume 9:271-278
168. Herbert J., Goodyer I.M., Grossman A.B., Hastings M.H., de Kloet E.R., Lightman S.L., Lupien S.J., Roozendaal B., Seckl J.R.. Do corticosteroids damage the brain?. *J. Neuroendocrinol.*. 2006. 18:393-411 doi:10.1111/j.1365-2826.2006.01429.x
169. Rezvyi G., Oiesvold T., Parnikov A., Olstad R.. A comparative study of diagnostic practice in psychiatry in Northern Norway and Northwest Russia. *Soc. Psychiatry. Psychiatr. Epidemiol.*. 2005. 40:316-323 doi:10.1007/s00127-005-0894-1
170. Keith S.J., Regier D.A.. U.S. and Soviet perspectives on the diagnosis of schizophrenia and associated dangerousness. *Schizophr. Bull.*. 1989. 15:515-517 doi:10.1093/schbul/15.4.515
171. Miller M.A.. The theory and practice of psychiatry in the Soviet Union. *Psychiatry*. 1985. 48:13-24 doi:10.1080/00332747.1985.11024264
172. Smulevich A.B.. Sluggish schizophrenia in the modern classification of mental illness. *Schizophr. Bull.*. 1989. 15:533-539 doi:10.1093/schbul/15.4.533
173. Pharmacopoeia article PA 42-2163-96. *Extractum Rhodiolae fluidum*. 1996
174. Rhizome and roots of . *National Pharmacopoeia of the USSR*. 1990. Volume 2:317-319
175. Krasik E.D., Morozova E.S., Petrova K.P., Ragulina G.A., Shemetova L.A., Shuvaev V.P., Avrutskiy G.Y.. Therapy of asthenic conditions: clinical perspectives of application of . *Proceedings modern problems in psycho-pharmacology, Kemerovo*. 1970. :298-300
176. Krasik E.D., Petrova K.P., Rogulina G.A.. About the adaptogenic and stimulating effect of . *Proceedings of all-union and 5th Sverdlovsk area conference of neurobiologists, psychiatrists and neurosurgeons*. :215-217
177. Saratkov A.S., Marina T.F., Kaliko I.M.. The stimulating effect of . *Vestn. Sibirsk. Otd. USSR Acad. Sci.*. 1965. 8:120-125
178. Mikhailova M.N., Goldsberg E.D.. Clinical and experimental substantiation of asthenic conditions therapy using . *Current problems of psychiatry*. 1983. :126-127
179. Brichenko V.S., Kupriyanova I.E., Skorokhodova T.F., Goldsberg E.D.. The use of herbal adaptogens together with tricyclic antidepressants in patients with psychogenic depressions. *Modern problems of pharmacology and search for new medicines*. 1986. Volume 2:58-60
180. Zakharov A.A.. On the problem of Schizandra tincture effect on the higher nervous activity in cases of asthenia of different origin. *Proceedings of Omsk Medical Institute*. 1956. Volume 20:93-94
181. Farutina M.V., Lazarev N.V., Komarov's V.L.. Clinical investigation of tonic effects of pantocrine, and Ginseng and Schizandra tinctures. *Materials for the study of stimulants and tonics from Ginseng root and Schizandra*. 1951. Volume 1:131-136
182. Galant I.B., Kuznetsova A.I., Suvorina N.A., Shikhova S.S., Gutschina I.S.. The experience in using . *Clinical development and therapy of mental diseases and organisation of psychoneurological assistance-Abstract book*. 1957. :112-113
183. Jadad A.R., Moore R.A., Carroll D., Jenkinson C., Reynolds D.J., Gavaghan D.J., McQuay H.J.. Assessing the quality of reports of randomized clinical trials: is blinding necessary?. *Control. Clin.Trials*. 1996. 17:1-12 doi:10.1016/0197-2456(95)00134-4

184. Romas R.S.. About the effect of Schizandra chinensis on higher brain structures of schizophrenia patients and chronic alcoholics. *PhD Thesis*. 1967. :1-19
185. Romas R.S.. Effect of . *Vopr. Psykhonevrol.*. 1962. 30:435-440
186. Romas R.S.. The effect of . *Voprosy Klin Nevrapat Psykhiatrii*. 1958. :211-216
187. Lastovetskij V.V., Romas R.S.. Special features in the interaction of signal systems under the influence of . *Zh. Vyssh. Nerv. Deiat. Im. I. P. Pavlova*. 1963. 13:604-609
188. Sudakov V.N., Savinykh A.B., Agapov Yu.K., Goldsberg E.D.. The role of adaptogens in the psychoprophylaxis of patients with borderline states of exogenous–organic genesis. *Modern problems of Pharmacology and Search for New Medicines*. 1986. :61-64
189. Staritsina O.N.. The experience of using Schizandra preparations against depressive states in psychiatry. *Zdravookhr. Kazakhstana (Alma-Ata)*. 1946. 6/7:42-44
190. Sivertsev I.I.. *Farmakol. Tokikol.*. 1946. 9:10-12
191. Sivertsev I.I.. About the effect of . *Izvetiya Akad. Nauk Kazakhskoy SSR Biol. Ser. (Alma-Ata)*. 1950. 91:33-36
192. Romas R.S.. Specific features of . *Materials of Science Conference of Ternopol State Medical Institute*. 1960. :50-51
193. Metscheryakova E.I., Mikhailiva N.M., Abrameitsev V.D.. A study of Rhodiola rosea extract effect using the MMPI method. *Problems of Rehabilitation of Patients with Nervous and Mental Diseases*. 1975. :180-182