

British Journal of Pharmaceutical Research 7(2): 63-75, 2015, Article no.BJPR.2015.092 ISSN: 2231-2919



SCIENCEDOMAIN international

www.sciencedomain.org

A Systematic and Comprehensive Review on Withania somnifera (L.) Dunal- An Indian Ginseng

Anisha Bano¹, Navdeep Sharma¹, Harcharan S. Dhaliwal¹ and Vivek Sharma^{1*}

¹Department of Biotechnology, Akal College of Agriculture, Eternal University, Baru Sahib-173101 (H.P.) India.

Authors' contributions

This work was carried out in collaboration between all authors. Author AB designed the study, wrote the protocol, and wrote the first draft of the review manuscript. Author NS managed the literature searches with author AB. Authors HSD and VS managed the process of review article and identified the plant species. All the authors read and approved the final manuscript for publication.

Article Information

DOI: 10.9734/BJPR/2015/17102

Editor(s):

(1) Ali Nokhodchi, Professor of Pharmaceutics and Drug Delivery, School of Life Sciences, University of Sussex, UK.

Reviewers:

(1) Anonymous, National Research Centre, Egypt.

(2) Slobodan Jankovic, Faculty of Medical Sciences, University of Kragujevac, Serbia.

(3) Anonymous, University of Naples Federico II, Italy.

Complete Peer review History: http://www.sciencedomain.org/review-history.php?iid=1177&id=14&aid=9465

Review Article

Received 26th February 2015 Accepted 24th April 2015 Published 28th May 2015

ABSTRACT

Present review article reveals the importance of species *Withania somnifera* (L.) Dunal, distributed in India and other parts of the world, this extensive research information on this species is highly significant for future researchers worldwide. In this article cytomorphological, phytochemical and biological activities inputs have been extensively recorded and discussed. As a part of our investigation on cytomorphological and phytochemical aspects for important medicinal plants from India, the aim of this pioneer attempt is to provide precise, truthful and detailed information of *W. somnifera* (L.). As per our knowledge, there is not even a single, combined, constructive review report available about this species, evaluated by using cytomorphological, phytochemical and biological activities based aspects.

Keywords: Ashwagandha, biological activities; cytomorphology; indian ginseng; phytochemistry; Withania somnifera (L.) Dunal.

1. INTRODUCTION

Plants are one of the most important sources of medicines in world and the writing indicate their medicinal uses as old as 4000-500 B.C. Chinese were the first to use the herbal preparations as medicines. In India, Rig-Veda (written between 3500-1600 B.C.) have earliest references of use of plants as medicine and later the properties and therapeutic uses of medicinal plant were recorded in Ayurveda [1]. World Health organization has listed more than 20,000 species of medicinal plants which are used globally [2]. According to WHO, for primary health care more than 80% of World's population relies on traditional herbal medicines [3]. Various drugs and chemicals derived from various parts of the plants and they continue to be possible sources of new drugs and chemicals [4,5]. Withania somnifera (L.) Dunal is an important medicinal plant and used in Ayurvedic medicines for the treatment of many diseases and it is also used in other parts of world [6,7]. In Ayurveda it is known as 'Rasayana' because it promote health and longevity, arrest ageing process, increase capability of individual to resist adverse environmental conditions [8]. The steroidal lactones (withanolides) obtained from its roots are mainly responsible for its therapeutic activities and general health maintenance like immunomodulation, combating infectious agents, anti-cancer, anti-epileptic, anti-ageing, antihypoglycemic, hypocholesterolemic oxidant. activities, memory enhancer and in common an effective adaptogen. These compounds also

work as "marker compounds/agents" for chemical standardization of Ashwagandhabased products [9-11]. W. somnifera (L.) Dunal commonly known as "Ashwagandha", "Asgandh" "Winter Cherry" belongs to family Solanaceae and widely distributed in warmer parts of the world. Genus Withania comprises 23 species including W. somnifera (L.) Dunal and W. coagulans (L.) Dunal having high medicinal value which are used as "Rasayana" in Ayurvedic formulations [12]. Ashwagandha attains the special name because its root smells like horse ("Ashwa") and believe to provide power like horse when consumed [11]. In Vedas it is described as herbal tonic and health food and considered as 'Indian Ginseng' because of its ginseng like health promoting effects [8,13]. Ashwagandha is as one of the ingredients in 74 Ayurvedic, 9 Siddha, 3 Unani and 126 herbal formulations [14]. Ashwagandha improves energy and also memory by enhancing the brain and nervous function; shows anxiolytic effects, has hepatoprotective property, raises hemoglobin level and red blood cell count, improve energy level; has potent antioxidant activity, improve the cell-mediated immunity; promotes vigor and vitality along with cheerful sexual life and reproductive equilibrium and act as powerful adaptogen [15-20]. As per our knowledge, there is not even a single, combined, constructive review report available about this highly recommended and utilized Indian species W. somnifera (L.) Dunal evaluated by using cytomorphological, phytochemical and biological activities based aspects from India and abroad.

1.1 Botanical Classification		1.2 Vernacular Names		
Kingdom:	Plantae	Arabic	:	Kaknaj-e-Hindi
Division:	Angiosperms	Bengali	:	Ashvaganda, Asvagandha
Class :	Dicotiledoneae	English	:	Winter cherry
Order :	Tubiflorae	Gujarati	:	Asan, Asana, Asoda, Asundha
				Ghodaasoda,
Family :	Solanaceae	Malayalam	:	Amukkiram, Pevetti
Genus :	Withania	Marathi	:	Askandha, Kanchuuki, Tilli
Species :	somnifera	Odiya	:	Asugandha
		Persian	:	Kaknaj-e-Hindi, Asgand Nagaori
		Sanskrit	:	Ashvagandha, Ash vakandika,
				Gandhapatri, Palashaparni
		Tamil :	:	Amukkira, Asubam, Asuvagandi
		Telugu	:	Asvagandhi, Penneru, Pennerugadda,
				Dommadolu
		Urdu :	:	Asgand, Asgand Nagori [21-23]

1.3 Common Names

Winter cherry (Eng.); Bitterappelliefie, Geneesblaarbossie, Koorshout (Afr.); Bofepha (Sotho); Ubuvuma (Xhosa); Ubuvimbha (Zulu); Ashwagandha (Hindi).

1.4 Conservation Status

Raimondo and co-workers, described the status of *W. somnifera* as 'Least Concern' as it is not a threatened species [24].

1.5 Geographical Distribution

W. somnifera is widely distributed around the world from Southern Mediterranean regions to the Canary Island and from South to East Africa; from Palestine to North India covering Israel, Jordan, Egypt, Sudan, Iran, Afghanistan, Baluchistan and Pakistan. In India the plant can be seen growing wild in the North Western regions extending to the mountainous region of Punjab, Himachal Pradesh and Jammu up to an altitude of 1,500 m [14].

2. CYTOLOGICAL STATUS

W. somnifera is reported to have intraspecific variations chromosomal having 2n=2x=24 [25], tetraploid, 2n = 4x = 48 [26] and hexaploid, 2n=6x=72 [27] cytotypes located in different parts of India. An aneuploid cytotype with 2n=42 from a callus regenerated plant is also mentioned in some reports [28]. The plant shows more than 97% incompatibility between wild and cultivated accessions which may be due to self-pollinating nature of the plant [29]. This plant show high degree of variability with respect to growth, habit and morphological characteristics in different parts of India and five distinct morphotypes of this plant have been identified from India [30]. The comparative detailed cytological studies on different accessions from North India have also been reported [31].

3. MORPHOLOGICAL DESCRIPTION

W. somnifera is an erect, green, branched or unbranched herb with height up to 1.25m. The aerial parts like stem, leaves and calyx are covered with fine hairy tomentum. Its branches are rounded; leaves are simple, petiolate, ovate, entire, shiny smooth and opposite; flowers are inconspicuous, greenish or yellow, in axillary umbellate cymes, bisexual; fruit is a berry in persistent calyx and seeds are small, flat, yellow,

reniform, very light [14,32]. The plant prefers a sunny situation, seeds are sown during June or July and seedlings are transplanted at a distance of 60 cm x 30 cm [33]. It is susceptible to bacterial, fungal, viruses, phytoplasms and pests infections [34]. Comparative morphological evaluations from wild accessions have also been extensively covered to find out new cytomorphotypes from North Indian wild germplasm [31].

4. CHEMICAL CONSTITUENTS

Withanolides and alkaloids are the major secondary groups characterized from W. somnifera and are of great medicinal interest [35]. Large numbers of withanolides have been isolated from its roots and leaves which attribute the medicinal property of this plant. Withaferin A represented the first natural lactone of the withanolide series isolated from its shoots [36]. Most of the pharmacological activities of this plant are due to two main withanolides, withaferine A and withanolide D [37]. Its roots are the major source of desired phytochemicals. Withanolides are characterized by the presence of C-28 basic skelton with a nine carbon atom side chain in which C-22 and C-26 oxidized to form a six membered lactone ring (CS-8). According to biogenetic point of view, withanolides are considered to have a cholestane type structure with an extra methyl group at C-24 and various oxygenated groups or double bonds placed at different sites of the skelton [14]. Five distinct chemotype of this plant, three from Israel and one from South Africa and India, were reported [38,39]. High performance liquid chromatography (HPLC) investigation of various organs of W. somnifera show a gradual decrease in content of withanolide A from aerial parts i.e., from young leaves to the root [40]. High-performance thin-layer chromatography (HPTLC) quantification of major bioactive withanolides like withaferine A, 1,2 deoxywithastramonolide, withanolide A and withanolide B from the root has also been done for rapid validation of Ayurvedic product [41]. GCMS analysis of roots and callus confirm the presence of 17 alkaloids out of which maximum present in roots followed by callus [42]. Metabolic profiling of chemotype variant in W. somnifera fruits using GC-MS and NMR spectroscopy has identified 82 chemically diverse metabolites consisting of fatty acids, organic acids, aliphatic and aromatic acids, polyols, sterols, sugars, tocopherols, phenolic acids and withanamides. Squalene and tocopherol has been identified for the first time in the fruit and rated as the most potent naturally compounds with antioxidant properties [43]. Some research work on *W. somnifera* has already been done in our laboratories to identify cytomorphotypes and chemotypes by morphometric parameters, cytological studies and (HPTLC) High Performance Thin Layer Chromatography, (HPLC) High Performance Liquid Chromatographic techniques [13,44].

5. PHARMACOLOGICAL AND BIO-ACTIVITIES

W. somnifera (L.) (Ashwagandha) is widely used in Ayurvedic medicines and is an important ingredient prescribed for а variety musculoskeletal conditions like arthritis. rheumatism and also used as general tonic to increase energy, improve overall health and longevity. Many pharmacological studies like immunomodulatory, cardioprotective, protective, anti-ageing and anti-oxidant etc. have been conducted to investigate the properties of Ashwagandha [45]. The information about various pharmacological activities have also been reviewed and given by few workers in their respective reviews [46,47]. It is one of the most important plants in Indian and pharmacopoeia. The main aim of the present review is to collect all the possible information on W. somnifera so that it will become easy for other workers to retrieve the truthful information from this review.

5.1 Anti-cancerous Activity

Cancer is a hyper proliferative disorder that results in apoptosis, transformation metastasis [48]. Millions of people suffer with various kind of cancer and die each year [49]. Ashwagandha a proud herb of Ayurveda has great anti-tumorigenic activity against various cancer cell lines due to the presence of withaferin A (WFA), a withanolide derived from this medicinal plant [50]. To investigate the anticancerous activity of W. somnifera various studies have been conducted. A study conducted on five human cancer cell lines of four different tissues that is PC-3, DU-145 (prostrate), HCT-15 (colon), A-549 (lung) and IMR (neuroblastoma) showed 0-98% in vitro cytotoxicity of 50% ethanol extract of root, stem and leaves depending on cell lines but 50% ethanol extract of leaves of W. somnifera showed maximum activity. Ethanol extract of leaves obtained from treatment T_2 , T_3 , T_4 and T_5 showed 80-90% growth inhibitor activity against PC-3 and HCT-

15 which is a strong activity, while 50% ethanol extract of leaves from T₁ treatment show minimal 39% growth inhibition against HCT-15 and T3 showed a maximum of 98% growth inhibition against HCT-15 [51]. Water extract of Ashwagandha leaves (ASH-WSX) both by in vitro and in vivo assays show anti-cancerous activity and triethylene glycol (TEG) was identified as active anti-cancerous component. It was also found that ASH-WSX was selectively cytotoxic to cancer cells and results in in vivo tumor suppression. Molecular analysis showed activation of tumor suppression protein p53 and pRB in ASH-WEX and TEG treated cancer cells [52]. W. somnifera (leaves) flavonoid compounds show anti-cancerous activities on cell lines MCF-7, A549 and PA1. These readings showed that W. somnifera (leaves) shows efficient cytotoxicity on MCF-7 (10±1µg) than PA1 (13±1µg) and A459 (11±1µg) cancer cell lines [53]. A long term tumor genesis study on Swiss albino mouse model shows that roots of W. somnifera inhibit the benzo(a)pyrene-induced forestomach papillomagenesis and results in 60 and 92% inhibition in tumor incidence and multiplicity. Similarly it also inhibits the 12-dimethylbenzanthracene-induced skin papillomagenesis by showing 45 and 71% inhibition in tumor incidence and multiplicity [54]. Squamous cervical cell line, SiHa treated with ethanolic extract of roots of W. somnifera show apoptosis which is an important step in cancer biology [55]. Withanolides are important for the treatment of cancer cells by suppressing the expression of oncoprotein Skp2 which is a member of the F-box family of substrate recognition subunits of SCF ubiquitin-protein ligase complexes which implicate in the ubiquitinmediated degradation [56]. Similarly in another study it was observed that a bioactive withanolide tubocapsanolide A inhibits the proliferation of human lung cancer cells via repressing Skp2 expression [57]. Withaferin A enhances radiation-induced apoptosis human renal cancer cells (Caki) through induction of ROS, Bcl-2 down regulation and Akt dephosphorylation [58]. Yu and co-workers, conducted an in vitro and in vivo study on pancreactic cancer cell to reveal that withaferin show in vitro antiproliferative activity against pancreactic cancer cell lines Pac-1, MiaPaCa and BxPc3 with IC₅₀ value of 1.24, 2.93, 2.78μ respectively. The study also investigated that withaferin A binds to Hsp90, inhibits its chaperone activity through an ATP-independent mechanism, results in its client protein degradation and exhibits in vivo anticancer

activity against pancreatic cancer [59]. A novel bioactive compound withanolide sulfoxide obtained from the methanol extract has been shown to W. somnifera roots inhibit nuclear transcription factor-kappa-B, cyclooxygenase and tumor cell proliferation against the human breast (MCF-7) cancer cell lines [60]. It has also shown that leaf extract of W. somnifera produce antiproliferative activity against MCF-7 human breast tumor cell lines [51]. Anticancerous activity of W. somnifera was also confirmed by various workers [61-64].

5.2 Immunomodulatory Activity

A series of studies conducted on animals showed that W. somnifera has a profound effect on hematopoietic system by acting as an immunoregulator and chemoprotective agent [1, 65]. Extract of this plant experimentally in normal mice records the increased cell mediated immunity (CMI) and root extract known to enhance the level of interferon gamma (IFN-y), interleukin (IL-2) and granulocyte macrophages colony stimulating factor (GM-CSF) in mice. This suggests their immunoprotective and myeloprotective effect. Ashwagandha increases the microbes killing power of these immune cells by enhancing nitric oxide synthatase activity of the macrophages [66]. For increasing the phagocytic activity, it activates and mobilizes macrophages, potentates the activity of lysosomal enzymes and acts as an anti-stress molecule and anti-inflammatory agent in mice and rat [67]. Ethanolic extract of W. somnifera enhances humoral immune response in Swiss albino mice on 7th day of administration by 12% as compare to control cyclophosphamide (54%). Cell mediated immune response was enhanced 19.27% in comparison with control cyclosporine (37.63%). In tumor bearing mice, the effect of methanolic extract on hematological parameter shows an increase in number of RBCs and a decrease in WBCs compared to the control mice [33]. Treatment with five doses of Withania root extract (20mg/dose/animal; i.p.) in Babl/c mice showed an enhancement of the total WBC count (17125 cells/mm on 10th day), increase in bone marrow cellularity (27x10 cells/femur) and significant increase (P<0.001) in α-esterase cell number (1800/4000 positive cells). Administration of Withania extract along with antigen (SRBC) showed enhancement in circulating antibody titer, maximum number of plaque forming cells PFC (985 PFC/10 on fourth day) in spleen and inhibition of delayed type of hypersentivity in mice. It also showed an

enhancement in phagocytic activity of peritoneal macrophages (76.5 pigmented cells/200) when compared to control (31.5/200cells) [68]. W. somnifera is an excellent immunomodulator because in the absence as well as in presence cyclophosphamide it influences production, enhances neutrophil counts and produces significant humoral response against sheep RBCs. The immunomodulatory activities may be due to withaferin A and withanolide D [69]. Extract of W. somnifera shows immunomodulatory effects on azoxymethane induced colon cancer in Swiss albino mice when treated with plant extract of 400 mg/kg body weight once a week for four weeks orally [64]. The levels of IFNy, IL-2 and GM-CSF was reversed to approximately normal in cyclophosphamide treated mice when administrated with W. somnifera extract. This also lower the levels of tumor necrosis factor- a and this indicate the immunopotentating and myeloprotective effect of W. somnifera [70].

5.3 Cardioprotective Activity

Myocardial infraction is the most lethal manifestation of cardiovascular diseases and it is one of the most important subjects of intense investigation by scientists [71]. Now a day there is an increased realization that herbs can maintain the balance of body and can influence heart diseases and its treatment by providing nutritional substances [72-73]. Although the therapeutic properties of this plant like immunomodulatory, adaptogenic, antioxidant, hypoglycemic and anti-cancerous are well known [74,75], but very few studies are available, which assess its cardioprotective potential [9]. Cardioprotective effect of hydroalcoholic extract of W. somnifera was studied on the basis of haemodynamic, histopathological and biochemical parameters in the isoprenaline-(isoproterenol) induced myocardial necrosis in rats and was compared with vitamin E, a known cardioprotective antioxidant. the drugs restore the myocardial antioxidant status and maintain membrane integrity by evidently reducing the malonyldialdehyde levels. Cardioprotective effect of these drugs was also confirmed by histopathological examinations. W. somnifera at 50 mg/kg dose shows maximum cardioprotective effect [76].

5.4 Neuroprotective Activity

Ashwagandha can be used for the treatment of neurodegenerative disorders like Alzheimer,

Parkinson. Huntington and neurodegenerative disorders at any stage of disease because it can significantly revers the neurotic atrophy, synaptic loss, along with GABA mimetic effect and promotes formation of dendrites due to therapeutic activity glycowithanolides withaferin A VII-X present in roots of ashwagandha [77-80]. Scientists of the institute of Natural Medicine at the Toyama Medical and Pharmaceutical University of Japan, by using the valid model of damaged nerve cell and impaired nerve signaling pathway, showed that the plant supports significant regeneration of the axon and dendrites of nerve cells. Its extracts supported the reconstruction of synapses or networks of the nervous system and it may act as potential treatment for neurodegenerative diseases such as Alzheimer, Parkinson etc. [18]. At the same institute during another study researcher found that Ashwagandha help in growth of nerve cell dendrites which suggests that Ashwagandha help in healing the brain tissue changes that accompany dementia [81]. Parkinson's disease 6-Hydroxydopamine rat models were pretreated with 100, 200 and 300 mg/kg dose of W. somnifera extract for 3 weeks. W. somnifera extract significantly reverse all the parameters (glutathione content, activities of glutathione-S-transferase, glutathione reductase, glutathione peroxidase etc.) to normal in dose dependent manner and demonstrate that it is helpful in protecting the neuronal injury in Parkinson's disease [80]. Ashwagandha root extract increases cortical muscarinic acetylcholine receptor capacity by affecting mainly the cortical and basal forebrain cholinergic signal transduction cascade which leads to cognition- enhancing and memory improving effect in animals and humans [77]. The pre-treatment of W. somnifera results in attenuation of cerebral ischemia-reperfusion and long-term hypoperfusion induced alterations in rats which confirm the neuroprotective effect of W. somnifera [82]. In another study female albino rats were subjected to immobilization stress for 14h and then treated with the root extract of W. somnifera which results in significant reduction (80%) in the number of degenerative cells in hippocampal sub-regions of rat [83]. Sankar and co-workers studied the influence of root extract of W. somnifera on Parkinsonism in MPTPintoxicated mice [84]. Oral treatment resulted in significant improvement in behavior, antioxidant status and a significant reduction in level of lipid peroxidation. Malhotra and co-workers have also studied the neuroprotective effect of this plant on the central nervous system [85].

5.5 Anti-oxidant / Anti-aging Activity

There are many herbal plants which contain the antioxidant compounds and protect the cells from degenerative effect of Reactive Oxygen Species (ROS) [86,87]. Now research is going on to revel potential phytochemical antioxidant because they are safer to health and have better antioxidant activity than synthetic antioxidants [88]. W. somnifera acts as a powerful antioxidant by increasing the level of three naturally occuring antioxidant enzymes like superoxide dismutase. catalase and glutathione peroxidase in the brain of rats [9]. Active principles present in the root of W. somnifera have powerful antioxidant effect like anti-stress, cognition-facilitating, antiinflammatory and anti-aging [89]. Antioxidant protects the body against free radical damage. An in vitro antioxidant activity of extract of different parts of W. somnifera shows that these are the potential scavengers of radicals and protector of lipid membrane in order of: leaves > fresh tubers > dry tubers. The antioxidant activity may be due to withanolides, glycowithanolides and sitoindosides VII-X. So the study indicates that Ashwagandha could be proved as natural source of safe anti-oxidative agent [90]. GENtreated rat showing nephrocytotoxicity when administrated with W. somnifera (500mg/kg) significantly reverses the signs of tubular necrosis. Overall results suggest nephroprotective effect of W. somnifera could be due to enhanced antioxidant activity with natural antioxidants and scavenging free radicals present in it [91]. Root powder of this plant prevents Cadmium- induced oxidative stress in chickens and lead induced oxidative damage in mouse [92-94]. A study was conducted to identify the antioxidant activity of extracts from W. somnifera by using various in vitro methods like 2, 2-diphenyl, 1-picryl hydrazyl (DPPH) radical scavenging activity, reducing competition with DMSO, hydroxy group reducing capacity, estimation of total phenol and estimation of ascorbic acid. The highest percentage (%) of DPPH (83.07) scavenging activity was found in polar flavonoid extract [87]. Similarly, free radical scavenging activity of ethanolic and aqueous extract of W. somnifera leaves was studied by DPPH and NBT methods and the results indicate that leaf possess antioxidant activity which may be due the presence of flavonoids and tannins in the leaf extract [95]. A double blind clinical trial conducted on 101 healthy male (50-59) for one year to test the anti-aging property of Ashwagandha showed a significant improvement

in hemoglobin, red blood cell count, hair melanin, seated stature, decreased serum cholesterol and preserved nail calcium in subjects. Sexual performance was also increased in 70% of subjects [37,89].

5.6 Antistress / Adaptogenic Activity

Traditionally, Ashwagandha is used to stabilize the mood of patients having behavioral disturbances and experimentally, it has also known to produce more anti-depressant and antianxiety effects as compared to drugs imipramine (anti-depressant) and lorazepam (anti-anxiety) [96]. Ashwagandha in India widely used as tranquillizer, improving reproductive and nervous system, rejuvenating body, improving vitality and recovery after chronic illness, so it hold an important position similar to ginseng in China [37,97]. Anti-stress activity of W. somnifera was conducted in rats using cold water swimming stress treatment and it was found that the drug treated animals show better stress tolerance [96]. Similarly, a withanolide free aqueous extract of roots show dose dependent anti-stress activity in Department of Pharmacology, mice [98]. University of Texas Health Science Center research conducted а on extract Ashwagandha and these extracts were known to produce GABA-like activity which may be responsible for its anti-anxiety effects [99]. GABA is an inhibitory neurotransmitter in brain, decrease neuron activity, inhibit nerve cells from over firing, thus produces a calming effect. Two new acylsterylglucosides, sitoindoside VII and VIII isolated from root of W. somnifera show antistress activity and the preliminary toxicity also indicate the low order of acute toxicity of compounds [97]. In a passive rat experiment model, animals were subjected to multiple stress of cold, hypoxia, restrain (C-H-R) and treated with aqueous root extracts of W. somnifera and an active compound X isolated from it. C-H-R stressed rats treated with W. somnifera extract and compound X can better withstand the stress than control group [100]. A standardize W. somnifera root and leaf extract (WSE) reduces stress related parameters in chronically stressed humans. It was found that WSE treatment show greater dose-dependent response in parameters (mHAM-A score, serum cortisol, serum Creactive protein, pulse rate, blood pressure and mean serum DHEAS and hemoglobin) and also show significantly greater responses in mean fasting blood glucose, serum lipid as compared to placebo [101]. To study the adaptogenic activity of W. somnifera, chronical stress (CS)

was induced in Winstar rats through mild, unpredictable foot shock stress procedure once daily for 21 days. The significant conditions induced by CS like hyperglycaemia, glucose intolerance, increase in plasma cortisol levels, gastric ulcerations, male sexual dysfunction, cognitive defects, immunesuppression and mental depression were attenuated by WS (25 and 50 mg/kg) and Panax ginseng (100 mg/kg) which were given 1h before foot shock for 21 days [16]. In another study adaptogenic activity of a novel with anolide free aqueous fraction from roots of W. somnifera was conducted on various parameters like hypoxia time, antifatique effect, swimming performance time, swimming induced gastric ulceration, hypothermia, immobilization induced gastric ulceration, autoanalgesia and biochemical changes in adrenal gland. The extract has shown activity in dose related manner [102].

5.7 Anti-diabetic Effect

Sarangi and co-workers conducted investigation to explore the possibilities of using leaf and root extracts of W. somnifera against diabetes mellitus (DM) and also to examine their hypoglycaemic and hypolipidaemic effects on streptozotonic-induced diabetic rats [103]. The extract possess hypoglycaemic and hypolipidaemic properties and hence useful in diabetes mellitus. Another study show significant positive anti-diabetic activity of W. somnifera on diabetic when compared with Glibenclamide standard drug. Anti-diabetic activity may be due to increase in hepatic metabolism, increased insulin release from pancreatic β -cells or insulin sparing effect [104]. W. somnifera root (WSREt) and leaf (WSLEt) extract show hypoglycaemic and hypolipidaemic effect on alloxan-induced diabetic rats [105]. Andallu and Radhika (2000), the hypoglycemic, diuretic hypocholesterolemic effect of roots of W. somnifera on six mild NIDDM and six mild hypercholesterolemic human subjects. Their studies indicate that the plant can be a potential source of hypoglycemic, diuretic and hypocholesterolemic drugs. No adverse effects were observed during clinical observations [106].

5.8 Anti-inflammatory Activity

Anti-inflammatory activity of *W. somnifera* has been attributed to the naturally occurring steroids, of which withaferin A is a major component and as effective as hydrocortisone sodium succinate dose, an anti-inflammatory

drug [98]. Rats treated with powder of W. somnifera orally 1h before the injection of inflammatory agent for 3 days produces antiinflammatory responses which are comparable to hydrocortisone sodium succinate [107]. In another experiment it was found that WS causes dose-dependent suppression of a2macroglobulin which is an indicator for antiinflammatory drugs in the serum of rats inflamed sub-plantar injection of carrageenan suspension. The doses of WS root powder (500, 1000, 1500 or 1200 mg/kg) were given orally 3-4h prior to induction of inflammation and it was found that maximum effect (about 75%) was seen at 1000 mg/kg [108]. Withaferin A was found to suppress the arthritic syndrome effectively without any toxic effect. In arthritic syndrome animals treated with hydrocortisone show weight loss while animal treated with withaferin A show gain in weight [109,110].

6. CONCLUSION

W. somnifera (L.) Dunal (Ashwagandha) has several health benefits, so is the most important 'Rasayana' in Indian Ayurveda. The plant has the ability to boost the immune functions, enhance the longevity and facilitate the restoration of homeostasis by reducing the stress. The manifestations can be made on the basis of this comprehensive perusal of literature, that the W. somnifera is being used traditionally, due to their immense therapeutic potential to treat/cure diseases. Important withanolides particularly withaferin-A extracted from root and leaf of W. somnifera play an important role in inhibiting the growth of human cancer cells. The roots of this plant are effective against the entire health problems already discussed here. We can say that the consumption of various parts of this plant as a dietary supplement may prevent or decrease the growth of tumors in cancer patients but can also protect the body from health conditions. Cytomorphological reveals that there is an immense need to find out new cytomorphotypes for further germplasm maintenance and evaluation, because till today nobody is working on these important aspects. As per reported phytochemical data, it is concluded that there is a need to identify few more chemotypes for further herbal and allopathic drugs formations. There is huge need and possibilities to isolate new components from untouched cytomorphotypes of the Genus Withania from India. Many studies demonstrated significant anti-inflammatory, anticancer, anti-diabetic, immunomodulatory and

anti-bacterial activities etc. which are reported in the extracts of different parts and from its phytoconstituents. These pharmacological activities and identified compounds provide solid scientific evidence for some of the traditional therapeutically claims. A variety of phytoconstituents has been isolated from the different parts of various species. It could be rightly said that due to its wide pharmacological activities, W. somnifera is considered as an important multi-purpose medicinal agent and an important component of various polyherbal preparations. Thus the plant has immense practical applicability in biomedicine as well as veterinary medicine focusing its potent role in the maintaince of sound health but more clinical trials should be conducted to support its therapeutic uses. Thus, there remains a wide scope for further scientific exploration of W. somnifera to their therapeutic efficacy establish commercial exploitation. In this review article we have made a constructive, detailed survey of literature for the species W. somnifera from Indian origin, which will definitely help researchers from India and abroad.

CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.

ACKNOWLEDGEMENT

The authors are grateful to His Holiness Baba Iqbal Singh Ji President, The Kalgidhar Trust & Founder Chancellor of Eternal University (H.P.), Hon'ble Vice Chancellor, Eternal University, Baru Sahib (Himachal Pradesh) India and former Head, Dr. R. C. Gupta, Department of Botany, Punjabi University, Patiala (Punjab) India for providing necessary facilities and logistic support for carrying out the study.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

 Verma KS, Kumar A. Therapeutic uses of Withania somnifera (ashwagandha) with a note on withanolides and pharmacological

- actions. Asian J Pharm & Clin Res. 2011; 4(1):1-4.
- Pandey MM, Rastogi S, Rawat AK. Indian herbal drug for health care and overview. Int J Alt Med. 2008;6(1):1-10.
- Arun V, Liju VB, Reena JJV, Parthipan B, Renuka C. Traditional remedies of Kani tribes of Kottoor reserve forest, Agasthyabanam, Trivananthapuram, Kerela. Ind J Trad Know. 2007;6(4):589-594.
- Srivastav S, Singh P, Mishra G, Jha KK, Khosa RL. Achyranthes aspera-An important medicinal plant: A Review. Nat Prod Plant Resour. 2011;1(1):1-14.
- Tijani Y, Uguru MO, Salawu OA. Antipyretic, anti-inflammatory and antidiarrhoel properties of Faidherbia albida in rats. Afr J Biotechnol. 2008;7(6):696-700.
- Patwarthan B, Panse GT, Kulkarni PH. Ashwagandha a review. J Nat Integr Med Assoc. 1998:30:7.
- Alam N, Hossain M, Khalil MI, Moniruzzaman M, Sulaimam SA, Gan SH. High catechin concentrations detected in Withania somnifera (ashwagandha) by high performance liquid chromatography analysis. BMC Compl & Alter Med. 2011;11:65.
- 8. Sharma K, Dandiya PC. Withania somnifera Dunal: Present Status. Ind Drugs. 1992;29:247.
- Dhuley JN. Adaptogenic and cardioprotective action of Ashwagandha on rats and frogs. J Ethnopharmacol. 2000;70:57-63.
- Shenoy S, Chaskar U, Sandhu JS, Paadhi MM. Effects of eight- week supplement of Ashwagandha on cardiorespiratory endurance in elite Indian cyclists. J Ayurv Intergr Med. 2012;3:209-214.
- Tiwari R, Chakraborty S, Saminathan M, Dhama K, Singh SV. Ashwagandha (Withania somnifera): Role in safeguarding health, immunomodulatory effects, combating infections and theapeutic Applications: A Review. J Biol Sci. 2014;14:77-94.
- CSIR. Withania somnifera L. (Dunal) (Ashwagandha). The Wealth of India – Raw Materials. Vol. X. Council of Scientific and Industrial Research, New Delhi, India. 1976:81-85.
- Sharma V, Gupta AP, Bhandari P, Gupta RC, Singh B. A validated and densitometric HPTLC method for the quantification of withaferin-A and

- withanolide-A in different plant parts of two morphotypes of *Withania somnifera*. Chromatographia. 2007;66:801-804.
- 14. Singh S, Kumar S. *Withania somnifera*: The Indian Ginseng, Ashwagandha. Central Institute of Medical and Aromatic Plants, Lucknow, India;1998.
- 15. Scarftiotti C, Fabris F, Cestaro B, Giuliani. Free radicals, atherosclerosis, ageing and related dysmetabolic pathologies: Pathological and clinical aspects. Eur J Cancer Prevention. 1997;6:S31-S36.
- Bhattacharya SK, Muruganandam AV. Adaptogenic activity of Withania somnifera: an experimental study using a rat model of chronic stress. Pharmacol Biochem Behav. 2003;75(3):547-555.
- 17. Arora S, Dhillon S, Rani G, Nagpal A. The in vitro antibacterial/synergistic activities of *Withania somnifera* extracts. Fitoterapia. 2004;75(3-4):385-388.
- Kuboyama T, Tohda C, Komatsu K. Neuritic regeneration and synaptic reconstruction induced by withanolide A. Br J Pharmacol. 2005;144(7):961:71.
- Harikrishnan B, Subramanian P, Subash S. Effect of Withania somnifera root powder on the levels of circulatory lipid peroxidation and liver marker enzymes in chronic hyperammonemia. E-J Chem. 2008;5:872-877.
- Sandhu JS, Shah B, Shenoy S, Chauhan S, Lavekar GS, Padhi MM. Effects of Withania somnifera (Ashwagandha) and Terminaalia arjuna (Arjuna) on physical performance and cardiorespiratory endurance in healthy young adults. Int J Ayurveda Res. 2010;1:144-149.
- 21. Anonymous. The Unani pharmacopoeia of India. Part I, Vol. I, depatt. of ayush, Ministry of health and family welfare, govt. of India, New Delhi. 2007;7-8.
- 22. Chopra RN, Nayar SL, Chopra IC. Glossary of Indian Medicinal Plants. Council of Scientific & Industrial Research, New Delhi. 1980;191:258.
- 23. Kirtikar KR, Basu BD. Indian Medicinal Plants. 2nd ed. Vol. III, Lalit Mohan Basu, Allahabad, India. 1980;1774-1777.
- 24. Raimondo D, Von Staden L, Foden W, Victor JE, Helme NA, Turner RC, Kamundi DA, Manyama PA. Red list of South African Plants, (eds.) Strelitzia 25. South African National Biodiversity Institute, Pretoria; 2009.

- Mohan Ram HY, Kamini. Embryology and fruit development in *Withania somnifera* Dunal. Phytomorph. 1964;14:574-587.
- 26. Iqbal M, Datta AK. Cytogenetic studies in *Withania somnifera* (L.) Dun. (Solenaceae). Cytologia. 2007;72:43-47.
- Bir SS, Neelam. SOCGI plant chromosome number reports LXII. Cytol.1984;19:113-114
- Ray S, Jha S. Regeneration of Withania somnifera plants. J Trop Med Plants. 2002; 3:89-95.
- Kaul MK, Kumar A, Sharma A. Reproductive biology of Withania somnifera (L.) Dunal. Curr Sci. 2005; 88(9):1375-1377.
- Atwal CK, Schwarting. intraspecific variability in Withania somnifera— preliminary survey. Llyodia. 1962;25:78-88
- 31. Sharma V, Dhaliwal H S, Gupta RC, Singh B. Comparative evaluation of cytomorphological studies on 27 accessions of "Indian Ginseng" *Withania somnifera* (L.) Dunal from North India. Braz J Bot. 2014;37(4):583-596.
- Atal CK, Gupta OP, Raghunathan K, Dhar KL. In: Pharmacognosy and Phytochemistry of Withania somnifera (linn.) Dunal (Ashwagandha). Central Council for Research in Indian Medicine and Homeopathy, New Delhi, India; 1975.
- 33. Verma SK, Shaban A, Purohit R, Chimata ML, Rai G, Verma OP. Immonomodulatory activity of withania somnifera (L.). J Chem & Pharm Res. 2012;4(1):559-561.
- 34. Alwadi HM, Baka ZA. Microorganisms associated with *Withania somnifera* leaves. Microbiol Res. 2001;156:303.
- Tripathi AK, Sukla YN, Kumar S. Ashwagandha [(Withania somnifera, Dunal (Solanaceae)]: A status report. J Med & Arom Plant Sci. 1996;8:46-62.
- Nigam KB, Kandalkar VS. Ashwagandha, In: Chadha KL, Rajendra G (eds) Advances in Horticulture Vol. 11-Medicinal and Aromatic Plants. Malhotra Publishing House, New Delhi, India. 1995;337-359.
- Singh G, Sharma PK, Dudhe R, Singh S. Biological activities of Withania somnifera. Annals Biol Res. 2010;1(3):56-63.
- Abraham A, Kirson I, Lavie D, Glotter E. The withanolides Withania somnifera chemotypes I & II. Phytochem. 1975;14:189-194.
- Kirson I, Glotter E, Lavie D, Abraham A. Constituents of Withania somnifera Dun.

- Part III. The withanolides of an Indian chemotype. J Chem Soc (London). 1971;11:2032-2044.
- 40. Praveen N, Naik PM, Manohar SH, Muthy HN. Distribution of withanolide a content in various organs of *Withania somnifera* (I.) Dunal. Int J Pharm & Bio Sci. 2010;1(3).
- 41. Devakar ST, Badhe YS, Jagtap SD, Hegde. Quantification of major bioactive withanolides in *Withania somnifera* (Ashwagandha) roots by HPTLC for rapid validation of Ayurvedic products. J Planar Chromatogr- Modern TLC. 2012;25(4).
- Sharma RA, Goswami M, Yadev A. GC-MS screening of alkaloids of Withania somnifera L. in vivo and in vitro. Ind J Appl Res. 2013;3(8).
- 43. Bhatia A, Bharti SK, Tewari SK, Sindhu OP, Roy R. Metabolic profiling for studying chemotype variations in *Withania somnifera* (L.) Dunal fruits using GC-MS and NMR spectroscopy. Phytochem. 2013:93:105-115.
- 44. Sharma V. HPLC-PDA Method for quantification of Withaferin- A and Withanolide- A in diploid (n = 12) and tetraploid (n = 24) cytotypes of Indian Ginseng *Withania somnifera* (L.) Dunal from North India. Int J Indigen Med Plants. 2013;46(2):2051-4263.
- 45. Devi PU, Sharada AC, Solomon FE, Kamath MS. In vivo growth inhibitory effect *Withania somnifera* (Ashwagandha) on a transplantable mouse tumor, sarcoma 180. Indian J Exp Bio. 1992;30:169-172.
- Aggrawal BB, Ichikawa H, Gorodia P, Weerasinghe P, Sethi G, Bhatt ID, Pandey MK, Shshodia S, Nair MG. From traditional Ayurvedic medicine to modern medicine: Identification of therapeutic targets for suppression of inflammation and cancer. Expert Opin Ther Target. 2006;10(1):87-118.
- Mishra LC, Singh BB, Dagenais S. Scientific basis for the therapeutic use of Withania somnifera (ashwagandha): a review. Alter Med Rev. 2000;5(4):334-346.
- 48. Gupta G, Rana AC. *Withania somnifera* (ashwagandha): a review. Pharmaco Rev. 2007;1(1):129.
- World Health Organization. "WHO calls for prevention of cancer through healthy workplaces" Press release. 2007;10-13.
- Mayola E, Gallerne C, Esposti DD, Martel C, Pervaiz S, Larue L, Debuire B, Lemoine A, Brenner C, Lemire C. Withaferin A induced apoptosis in human melanoma

- cells through generation of reactive oxygen species and down-regulation of Bcl-2. Apoptosis. 2011;16(10):1014-1027.
- Yadav B, Bajaj A, Saxena M, Saxena AK. In vitro anticancer activity of the root, stem and leaves of Withania somnifera against various human cancer cell lines. Indian J Pharm Sci. 2010;72(5):659-663.
- 52. Renu W, Rumani S, Ram G, Navjot S, Nashi W, Tomako N, Yashiyuki I, Keiji T, Sunil CK. Water extract of Ashwagandha leaves has anticancer activity: Identification of an active component and its mechanism of action. Pub Lib Sci. 2013;8(11).
- Nema R, Khare S, Jain P, Pradhan A. Anticancer activity of Withania somnifera (leaves) flavanoid compound. Int J Pharm Sci Rev & Res. 2013;19(1):103-106.
- 54. Padmavathi B, Rath PC, Rao AR, Singh RP. Roots of *Withania somnifera* inhibit forstomach and skin carcinogenesis in mice. eCAM. 2005;2(1):99-105.
- 55. Jha AK, Jha M, Kaur J. Ethanolic extract of Ocimum sanctum, Azadirachta indica and Withania somnifera cause apoptosis in SiHa cells. Res J Pharm, Biol & Chem Sci. 2012;3(2):557-562.
- 56. Sindhu K, Santhi N. Molecular Interaction of withanolides from *Withania somnifera* against the oncoprotein Skp2 using glide. Advanced Biotech. 2009;9(06):28-33.
- Chang HC, Chang FC, Wang YC, Pan MR, Hung WC, WU YC. A bioactive Tubocapsanolide A inhibits proliferation of human lung cancer cells via repressing Skp2 expression. Mol Cancer Ther. 2007;6(5):1572-1578.
- Yang ÉS, Choi MJ, Kim JH, Choi KS, Kwon TK. Withaferin A enhances radiationinduced apoptosis in Caki cells through induction of reactive oxygen species, Bcl-2 down regulation and Akt inhibition. Chem Bio Interact. 2011;190(1):9-15.
- Yu Y, Hamza A, Zhang T, Gu M, Zou P, Newman B, Li Y, Gunatilaka AA, Zhan CG, Sun D. Withaferin A targets heat shock protein 90 in pancreatic cancer cells. Biochem Pharmacol. 2010;79(4):542-551.
- Mulabagal V, Subbaraju GV, Rao CV, Sivaramakrishna C, Devitt DL, Holmes D, Sung B, Aggarwal BB, Tsay HS, Nair MG. Withanolide sulfoxide from Aswagandha roots inhibits nuclear transcription factorkappa-B, cyclooxygenase and tumor cell proliferation. Phytother Res. 2009;23(7): 987-992.

- Senthil V, Ramadevi S, Venkatakrishnan V, Giridharan P, Lakshami BS, Vishwakarma RA, Balakrishnan A. Withanolide induces apoptosis in HL-60 leukemia cells via mitochondria mediated cytochrome c release and caspase activation. Chemico-Biol Interact. 2007; 167:19-30.
- 62. Malik F, Kumar A, Bhushan S, Khan S, Bhatia A, Suri KA, Qazi GN, Singh J. Reactive oxygen species generation and mitochondrial dysfunction in the apoptotic cell death of human myeloid leukemia HL-60 cells by a dietary compound withaferin A with concomitant protection by N-acetyl cysteine. Apoptosis. 2007;12(11):2115-2133.
- 63. Koduru S, Kumar R, Srinivasan S, Evers MB, Damodran C. Notch-1 inhibition by Withaferin-A: a therapeutic target against colon carcinogenesis. Mol Cancer Ther. 2010;9(1):202-210.
- 64. Muralikrishnan G, Dina AK, Shakeel F. Immunomodulatory effects of *Withania somnifera* on azoxymethane induced experimental colon cancer in mice. Immunol Invest. 2010;37(7):688-698.
- 65. Kuttan G. Use of *Withania somnifera* Dunal as an adjuvant during radiation therapy. Indian J Exp Bio. 1996;34:854-856.
- Iuvone T, Esposito F, Capasso F, Izzo AA. Introduction of nitric oxide synthase expression by Withania somnifera in macrophages. Life Sci. 2003;72:1617-1625.
- 67. Rasool M, Varalakshmi P. Immuno-modulatory role of *Withania somnifera* root powder on experimental induced inflation: An in vivo and in vitro study. Vasc Pharmacol. 2006;44:406-410.
- 68. David L, Kuttan G. Immunomodulatory activity of *Withania Somnifera*. J Ethnopharmacol. 2000;71(1-2):193-200.
- 69. Grover A, Shandilya A, Punetha A, Bisaria VS, Sundar D. Inhibition of the NEMO/IKK β association complex formation, a novel mechanism associated with the NF- KB activation suppression by Withania somnifera's key metabolite wthaferin A. Biol. Med. Chem Genomics. 2010;4:S25-S36.
- Davis L, Kuttan G. Effect of Withania somnifera on cytokine production in normal and cyclophosphamide treatment mice. Immunopharmacol & Immunotoxicol. 1999;21(4):695-703.

- Bolli R. Mycardial ischemic metabolic disorder leading to cell death. Rev Post Cardiol. 1994;13:649-653.
- Dhar ML, Dhar MM, Dhawan BN, Mehrotra BN, Ray C. Screening of Indian plants for biological activity. J Exp Biol. 1968;6:232-247
- 73. Hertog MGL, Feskens EJM, Hollam PCH, Katan MB, Kromhout D. Dietary antioxidant flavonoids and risk of coronary heart diseases. The Zutphen Eldery Study. Lancet. 1993;342:1007-1020.
- Malhotra CL, Prasad KS, Das PK, Dhalla NS. Studies on Withania somnifera (part III). The effect of total alkaloid (ashwagandhandholine) on CVS and respiration. Ind J Med Res. 1961;49:449.
- Lavie D, Glotter E, Shro Y. Constituents of Withania somnifera. Dun IV. J Chem Soc. 1965:12:7517.
- Mohanty I, Arya DS, Dinda A, Talwar KK, Joshi S, Gupta SK. Mechanism of cardioprotective effect induced myocardial infarction. Bas & Clin Pharm & Toxicol. 2004;94(4):184-190.
- 77. Schliebs R, Liebmann A, Bhattacharya SK, Kumar A, Ghosal S, Bigl V. Systematic administration of defined extracts from *Withania somnifera* (Indian Ginseng) and Shilajit differentially affects cholinergic but not glutamatergic and GABAergic markers in rat brain. Neurochem Int. 1997;30:181-
- 78. Abbas SS, Singh V, Bhalla M, Singh N. Clinical study of organic Ashwagandha in case of parkinsonism, neuropathy, paralysis and uterine tumours (fibroids and other tumours) including cutaneous endodermal carcinoma, Proceeding of the National seminar on eco-friendly herbs of Ayurveda in healthcare of mankind: A strategy for scientific evaluation an uniform standardization, Lucknow. 2004;81.
- Abbas SS, Bhalla M, Singh N. A clinical study of organic Ashwagandha in some cases of uterine tumors (fibroids) and dermatofibrosarcoma. Proceedings of the workshop on essential medicines, Adverse Drug Reactions and Therapeutic Drug Monitoring, Lucknow. 2005;143-144.
- Ahmad M, Saleem S, Ahmad AS, Ansari MA, Yousuf S, Hoda MN, Islam. Neuroprotective effects of Withania somnifera on 6-hydroxydopamine induced Parkinsonism in rats. Human Exp Toxicol. 2005;137-147.

- Tohda C, Kuboyama T, Komatsu K. Dendrite extension by methanol extract of Ashwagandha (roots of Withania somnifera) in SK-N-SH cells. Neuroreport. 2000;11(9):1981-5.
- 82. Trigunayat A, Raghavendra M, Singh RK, Bhattacharya AK, Acharya SB. Neuroprotective Effect of *Withania somnifera* (WS) in cerebral ischemiareperfusion and long-term hypoperfusion induced alterations in rats. J Nat Remed. 2007;7(2):234-246.
- 83. Jain S, Shukla SD, Sharma K, Bhatnagar M. Neuroprotective effects of *Withania* somnifera Dunn. In hippocampal subregions of female albino rat. Phytother Res. 2001;15(6):544-548.
- 84. Sankar SR, Manivasagam T, Krishnamurti A, Ramanathan M. The neuroprotective effect of *Withania somnifera* root extract in MPTP-intoxicated mice: An analysis of behavioral and biochemical variables. Int Sci J. 2007;12:473-481.
- 85. Malhotra CL, Mehta VL, Das PK, Dhalla NS. Studies on Withania-ashwagandha, Kaul V. The effect of total alkaloids (ashwagandhadholine) on the central nervous system. Ind J Physiol Pharmacol. 1965;9:127-136.
- 86. Blois MS. Antioxidant determinations by the use of a stable free radical. Nat. 1958;181:1199-1200.
- 87. Aruoma O, Cuppett SL. Antioxidant methodology in vivo and *in vitro* concepts. AOCS Press Champaign.1997;41-172.
- 88. Ansari AQ, Ahmad SA, Waheed MA, Junned AS. Extraction and determination of antioxidant activity of *Withania somnifera*. Euro J Exp Biol. 2013;3(5):502-507.
- 89. Bone K. Clinical application of Ayurvedic and Chinese herbs. Phytotherapy Press, Queensland, Australia. 1996;137-141.
- Sumathi S, Padma PR, Gathampari S, Vidhya S. Free radical scavenging activity of Withania somnifera. Ancient Sci Life. 2007;26(3):30-34.
- 91. Jeyanthi T, Subramanian P. Nephroprotective effect of *Withania somnifera*: A dose-dependent study. Renal Failr. 2009;31(9):814-821.
- Chaurasia SS, Panda S, Kar A. Withania somnifera root extract in the regulation of lead-induced oxidative damage in the male mouse. Pharmacol Res. 2000;41:663-666.
- 93. Mahadik KR, Gopu CL, Gilda SS, Paradkar AR, Mahadik KR. Comparative evaluation

- of antioxidant potential of Ashwagandha arishta and self-generated alcoholic preparation of *Withania somnifera* dunal. Planta Med. 2008;74:288-288.
- Bharvi K, Reddy AG, Rao GS, Reddy AR, Rao SV. Reversal of cadmium-induced oxidative stress in chicken by herbal adaptogens Withania somnifera and Ocimum sanctum. Toxicol Int. 2010;17:59-63.
- Panchawat S. In vitro free radical scavenging activity of leaves extract of Withania somnifera. Rec Res Sci & Technol. 2011;3(11):40-43.
- Archana R, Namasivayam A. Anti-stressor effect of Withania somnifera. J Ethanopharmacol. 1999;64(1):91-93.
- 97. Bhattacharya SK, Goel RK, Kaur R, Ghosal S. Anti-stress activity of VIII. sitoindosides VII and new acylsterylglucosides from Withania somnifera. 1987;1(1):32-37.
- 98. Khare CP. Indian medicinal plants-An illustrated dictionary. First Indian reprint, Springer (India) Pvt. Ltd., New Delhi. 2007;717-718.
- Mehta AK, Binkley P, Gandhi SS, Ticku MK. Pharmacological effect of W. somnifera roots extracts on GABA receptor complex Ind J Med Res. 1991;94:312-315.
- 100. Kaur P, Mathur S, Sharma M, Tiwari M, Srivastava KK, Chandra R. A biologically active constituent of *Withania somnifera* (ashwagandha) with anti-stress activity. Ind J Clin Biochem. 2001;16(2):195-198.
- 101. Auddy B, Hazra J, Mitra A, Abedon B, Ghosal S. A standardized Withania somnifera extract significantly reduces stress-related parameters in chronically stressed humans: A double-blind, randomized, placebo-controlled study. J Ame Nutr Assoc. 2008;11(1):50-56.

- 102. Singh B, Saxena A.K, Chandan BK, Gupta DK, Bhutani KK, Anand KK. Adaptogenic activity of a novel, with anolide-free aqueous fraction from the roots of Withania somnifera Dun. Phytotherapy Research. 2001;15(4):311-318.
- 103. Sarangi A, Jena S, Sarangi AK, Swain B. Anti-diabetic effect of Withania somnifera root and leaf extracts on streptozotocin induced diabetic rats. J Cell & Tissue Res. 2013;13(1):3597.
- 104. Navinder, Khatak M, Sehrawat R, Khatak S. A Comparative Study: Homoepathic medicine and a medicinal plant Withania somnifera for antidiabetic activity. J Pharma & Phytochem. 2013;2(3):109-112.
- 105. Udayakumar R, Kasthurirengan S, Mariashibu TS, Rajesh M, Anbazhagan VR, Kim SC, Ganapathi A, Choi CW. Hypoglycaemic and hypolipidaemic effects of Withania somnifera root and leaf extracts on alloxan-induced diabetic rats. Int J Mol Sci. 2009;10(5):2367-2382.
- 106. Andallu B, Radhika B. Hypoglycemic, diuretic and hypocholesterolemic effect of Winter cherry (Withania somnifera, Dunal) root. Ind J Exp Biol. 2000;38:607-609.
- Anbalagan K, Sadique J. Role of prostaglandins in acute phase proteins in inflammation. Biochem Med. 1984;19:245:24.
- 108. Begum VH, Sadique. Long term effect of herbal drug Withania somnifera on adjuvant induced arthritis in rats. Ind J Exp Biol. 1988;26:161-161.
- Rastogi RP, Mehrotra BN. Compendium of Indian medicinal plants, vol.6. Central Drug Research Institute, New Delhi;1998.
- 110. Narinderpal K, Junaid N, Raman B. A review on pharmacological profile of Withania somnifera (Ashwagandha). Res & Rev J Bot Sci. 2013;2(4):6-14.

© 2015 Bano et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:

The peer review history for this paper can be accessed here: http://www.sciencedomain.org/review-history.php?iid=1177&id=14&aid=9465