

***Boerhaavia diffusa*: Bioactive Compounds and Pharmacological Activities**

Harpreet Kaur

Botany Department, Government College for Girls, Patiala-147001, Punjab, India.

*Corresponding author E-mail: harpkaur1987@gmail.com

<http://dx.doi.org/10.13005/bpj/1797>

(Received: 01 July 2019; accepted: 05 November 2019)

***Boerhaavia diffusa* belonging to family Nyctaginaceae has a wide distribution, occurring on major part of the globe. It is known as Punarnava in Ayurveda and is a main ingredient in many formulations of Ayurveda. It is used as traditional medicine by indigenous people of many countries in the world for its protective role against inflammation, prostatic hyperplasia, diabetes, cancer, gastrointestinal problems, arthritis etc. The whole plant contains numerous bioactive compounds which are responsible for its pharmacological activities. Experiments are being done to evaluate full potential of the plant. The present review focuses on the bioactive compounds and pharmacological activities of *B. diffusa*. The study is carried out with the aim that it will be helpful for more research and wide acceptance of *B. diffusa* so that the plant which till now was used by indigenous people for its medicinal properties will become an ingredient of major mainstream medicines used to treat chronic human ailments.**

Keywords: *Boerhaavia diffusa*, Bioactive compounds, Pharmacological activities, indigenous, antioxidants.

Boerhaavia diffusa is a species of flowering plants which belongs to family Nyctaginaceae (Four o'clock family). The genus '*Boerhaavia*' is so named to honour Hermann Boerhaave who was a famous 18th century Dutch physician and the species is named '*diffusa*' due to the typical diffuse branching of the plant (Mishra *et al.* 2014). It is also known as spreading Hogweed, Windflower, Red Spiderling, Tarvine, Satha, Punarnava, Itcit, Kathilla and by many other regional names. In ancient medicine, *B. diffusa* is commonly called as Punarnava because of its regeneration properties in rainy season. It is creeping perennial weed, prostrate herb, can be upto 1m long and with spreading branches. It occurs abundantly in ditches, marshy places and waste places during rains (Pooja *et al.* 2017). It is widely distributed in many

countries located in tropical and subtropical regions like India, Sri Lanka, Egypt, Sudan, Ghana, South Africa, Nigeria, China, Australia, Philippines, Iran etc. (Chaudhary and Dantu 2018, Nayak and Thirunavoukkarasu 2016). Entire plant is used as vegetable at some places. In some parts of India like Assam and West Bengal the indigenous people cook leaves as a vegetable (Jana 2007).

Many bioactive compounds like tannins, flavonoids, alkaloids (punarnavine), glycosides, steroids, terpenoids, phenolic compounds, rotenoids (boeravinones A-O) etc. are reported in *B. diffusa* plants (Parmer *et al.* 2018, Shisode and Kareppa 2011, Krishnamoorthy *et al.* 2017). Due to its ethanopharmacological, chemical and therapeutic values, it is classified as rasayana in Ayurveda. Ayurvedic formulations such as Punarnavasava,

Sukumara ghritha, Punarnavadyarishta, Punarnavadi mandura, Sothagha Lepa, Maha Narayan Taila, Punarnavastaka kvatha curna, Punarnava guggulu, Punarnavadi kvatha curna and Varuni contain *B. diffusa* as main ingredient. These formulations are used to treat numerous ailments in humans like heart disease, sciatica, inflammation, diseases of abdomen, spleen and liver disorders, arthritis etc. (Mishra *et al.* 2014). It is used as ethnomedicine in other countries also like in Brazil it is used to treat hepatitis, renal disorders, liver disorders, edema etc; in Iran it is used to treat gonorrhea, joint pain, jaundice, intestinal gas etc; in Nigeria for epilepsy, asthma, fever etc; in Philippines as diuretic and purgative and in Ghana for asthma and boils (Nayak and Thirunavoukkarasu 2016).

Long term use of synthetic drugs for the treatment of various chronic diseases always creates some side effects. So there is need to move towards alternative drugs i.e. herbal drugs which have no side effects due to the presence of bioactive compounds (Pooja *et al.* 2017). But there is need to improve the acceptance of classical and ayurvedic medicines in the world market which can be done by advancements in ayurvedic research which should be evidence based (Chauhan *et al.* 2015). Phytochemicals need to be identified and quantified in plants important in traditional medicine. Dosage of the medicine should be determined by further research. This all would establish quality standards for efficacy and safety of the traditional medicines and they would get huge acceptance and market in the whole world. Due to wide occurrence and easy accessibility of *B. diffusa*, medicines prepared from it will be cheap and will be a boon for the health of people especially for the people in low and middle income countries. Researches done in the past and recent years relating to the analysis of phytochemicals and pharmacological activities in *B. diffusa* are thoroughly studied and analysed and presented in the present work in a comprehensive way.

Bioactive compounds

Phytochemical analysis of extracts prepared from *B. diffusa* roots by Shisode and Kareppa (2011), Parmer *et al.* (2018) and Pooja *et al.* (2017) has revealed the presence of flavonoids, phenolic compounds, saponins, glycosides, alkaloids, tannins, terpenoids and steroids. Ethylacetate, diethyl ether, ethanol,

aqueous, chloroform and methanol extracts prepared from *B. diffusa* leaves were subjected to method of Harborne to test the presence of different phytochemicals by Umamaheswari *et al.* (2010). The qualitative analysis revealed the presence of different phytochemicals such as phenols, quinones, proteins, amino acids, saponins, carbohydrates, sterols, furanoids, alkaloids, glycosides, triterpenoids, flavonoids and tannins. The different extracts showed different constituency relating to the number of above phytochemicals. Three rotenoids, namely boeravinone A, B and C and 3-*O*-(6'-palmitoyl- β -D-glucopyranosyl) sitosterol, were isolated and their structure was determined by Lami *et al.* (1990) and Kadota *et al.* (1989). Boeravinone B was identified and quantified by employing HPLC in hydroalcoholic extracts made from whole plant of *B. diffusa* and also in its polyherbal formulation (Singh *et al.* 2017). The results showed that boeravinone B in the extract of *B. diffusa* was 0.041% w/w and in the polyherbal formulation was 0.011%. It scientifically validated the ratio of *B. diffusa* (ingredient) in the finished drug as one fourth of the finished product consisted of the ingredient and complied with the API limit. So the research established a criterion for global acceptance of ayurvedic drugs. Boeravinone B was isolated, purified and characterized from methanolic extracts of *B. diffusa* by Krishnamoorthy *et al.* (2017) by employing TLC and reverse phase HPLC. Aviello *et al.* (2011) first prepared methanol extracts of *B. diffusa* roots. After it with the help of Kupchan partitioning of the methanol extract four different fractions were obtained i.e. chloroform, n-hexane, n-butanol and carbon tetrachloride (CCl₄). As the CCl₄ fraction showed high antioxidant activity, it was subjected to further separation. Three rotenoids namely boeravinone D, boeravinone G and boeravinone H were isolated by using sequential silica gel column chromatography and HPLC. Their spectral data was compared with that reported in literature and so their structures were determined. Phenolic composition of *B. diffusa* collected from different locations was identified by Ferreres *et al.* (2005) by using HPLC-PAD-MS/MS. They identified ten phenolic compounds: quercetin-3-*O*-rhamnosyl(1-6)galactoside, kaempferol 3-*O*-(2''-rhamnosyl)-robinobioside, 3,4-dihydroxy-5-methoxycinnamoyl-rhamnoside, 3,5,4'-trihydroxy-

6,7-dimethoxyflavone 3-*O*-galactosyl(1'2) glucoside, quercetin 3-*O*-(2"-rhamnosyl)-robinobioside, kaempferol 3-*O*-robinobioside, caffeoyltartaric acid, quercetin, euphalitin 3-*O*-galactoside and kaempferol. The major phenolic compound in leaves was quercetin 3-*O*-(2"-rhamnosyl)-robinobioside and in roots was caffeoyltartaric acid. The results showed that the phenolic composition was influenced by the geographic origin of the plants. In that case also the nature of soil was the main responsible factor.

Pharmacological activities

Cardioprotective effect

Modern drugs used to treat various diseases always have some side effects. Its example can be seen in dual, healing as well as harming, effects of anthracycline drug, doxorubicin. It is an anticancer drug used to treat ovarian cancer, acute leukemia's, breast cancer and Hodgkin and non-Hodgkin lymphoma (Kim *et al.* 2006). It kills cancerous cells by intercalating with DNA and inducing apoptosis of cells. It also results into generation of reactive oxygen species, to which particularly the anti-oxidant defenses of heart are susceptible. Scientists are making efforts to have some herbal remedies of the problem. *B. diffusa* has been traditionally used in Ayurveda to treat cardiac disorders (Kirtikar and Basu 1991). Studies were undertaken by Nimbal and Koti (2016) to investigate the preventive role of ethanolic extract made from *B. diffusa* whole plant against doxorubicin generated myocardial toxicity in rats. Administration of doxorubicin elevated the levels of biochemical parameters such as alanine aminotransferase, aspartate aminotransferase and alkaline phosphatase; biomarker enzymes like creatine kinase and lactate dehydrogenase whereas reduced the levels of antioxidant enzymes such as catalase, glutathione and superoxide dismutase. The levels of all the above biochemical parameters, biomarker enzymes and antioxidant enzymes were normalized in the rats which were pretreated with the ethanolic extract of *B. diffusa*. So the data evidently proved the cardioprotective effect of *B. diffusa*, which was attributed to its antioxidant activity.

Treatment of prostatic hyperplasia

Benign prostatic hyperplasia (BPH) is common in elderly men as its prevalence is estimated to be 85%. Affected men have enlarged

prostate gland and experience troubles during urination (Lee 2008). Investigation was carried out by Vyas *et al.* (2013) to study the effect of hydroalcoholic extracts prepared from roots of *B. diffusa* on BPH in rats. Testosterone dissolved in arachis oil was administered subcutaneously in male Wistar rats for 28 days to induce BPH in them. The model rats showed significant enhancement in prostate weight when compared with negative control rats. But the treatment with *B. diffusa* extract significantly reduced the prostate weight. Incubation with *B. diffusa* extract attenuated the contractile response of prostate gland and vas deferens which was elicited by exogenous application of noradrenaline. Histoarchitecture of prostate gland also showed improvements in rats treated with both testosterone and *B. diffusa* as compared to the model rats. It was concluded that anti-proliferative and anti-inflammatory properties of bioactive components in *B. diffusa* extracts might had relaxed prostatic smooth muscles and relieved the urinary symptoms of BPH disease.

Anti-inflammatory action

Inflammatory reactions are usually treated by taking nonsteroidal anti-inflammatory drugs. But the use of these preferred drugs causes unwanted side effects such as inhibiting the protective cyclooxygenase enzyme in gastric mucosa which results in gastrointestinal damage (Rang *et al.* 2008). So there is arising more interest in natural drugs which can treat inflammation without any side effects. Anti-inflammatory effect of *B. diffusa* was evaluated in rats by Sudhamadhuri and Kalasker (2014). Aqueous extracts were prepared from the plant leaves and their activity was determined on sub acute inflammation (cotton pellet induced granuloma) and acute inflammation (carrageenan induced paw edema) in rats. Pre-administration of the extracts (200 and 400 mg/kg) to the rats resulted in dose-dependent anti-inflammatory activity against sub-acute as well as acute inflammation. It was concluded that the above effects of the extracts were probably due to inhibition of chemical mediators of inflammation.

Anxiolytic activity

Anxiolytic activity of hydroalcoholic extract of *B. diffusa* leaves was evaluated in rats by Gadekar *et al.* (2011). Haloperidol Induced Catalepsy method, ketamine induced sleep, Elevated Plus Maze test and Hole Board Test were

used to study anxiolytic effect of the extract. The results revealed that hydroalcoholic extracts of *B. diffusa* had significant anxiolytic activity which was comparable to standard drug Diazepam (0.5 mg/kg).

Protective effect on gastrointestinal problems

The disruption of integrity of stomach mucosa results in peptic ulcer disease. It results from imbalance between factors responsible for gastroduodenal defense (bicarbonate secretions, mucosal blood flow, prostaglandins and mucus) and factors promoting mucosal damage (*Helicobacter pylori* infection, pepsin, gastric acid) (Devi and Jyothi 2015). The anti-ulcer activities of *B. diffusa* are contributed by several mechanisms such as its healing effects, regulation of gastric secretions, anti-inflammatory properties and mucus stimulatory effects (Scott and Ellis 1980). Healing effect of *B. diffusa* on gastric ulcers had been investigated by Devi and Jyothi (2015). Administration of root extract of *B. diffusa* alone and also along with Omeprazole efficiently resulted in reduction in development of gastric ulcer secretion in indomethacin induced gastric ulcers, pylorus ligation induced gastric ulcers and stress induced duodenal and gastric ulcers in rats. They concluded that the healing effect of root extract of *B. diffusa* was due to decrease in gastric secretion and also due to reduced gastric action.

Anticancer activity

Radiotherapy, surgery and chemotherapy methods used to treat cancer are highly effective but exert numerous side effects. Also the cancer cells gradually become resistant to the treatments. To circumvent it scientists are doing numerous efforts to find herbal solutions to cancer. Remya *et al.* (2018) carried out in vitro experiments to study cytotoxicity of decoction prepared from root of *B. diffusa* in MCF-7 (Michigan Cancer Foundation-7) breast cell line. During 48 hours of incubation in MCF-7 breast cell line, the test sample of *B. diffusa* at 800 µg/ml concentration showed cytotoxicity of about 65.1 ± 1.2 . It proved the anticancer property of *B. diffusa* extracts. Anticancer activity of ethanol extract of *B. diffusa* leaves against Dalton's ascitic lymphoma in mice is also reported by Kayande and Kushwah (2014).

Antimicrobial activity

Antibacterial activity of ethanolic extracts prepared from roots, stem and leaves of *B. diffusa*

was evaluated by Majgaine and Verma (2017). Tests were conducted on three bacteria types *Staphylococcus aureus*, *Escherichia coli* and *Salmonella typhi*. Reduction in colony diameter (zone of inhibition) was the criterion to measure anti-bacterial activity of the extracts. Roots extracts exhibited highest and stem extracts the least antibacterial activities. The extracts from the three parts of the plants were highest efficient against *S. typhi* followed by *S. aureus*. But no inhibitory effect of the extracts was noted against *E. coli*.

Umamaheshwari *et al.* (2010) qualitatively analyzed phytochemicals as well as antimicrobial functions of extracts of *B. diffusa* made in different solvents. The extracts were tested against gram positive bacteria like *Streptococcus faecalis*, *Bacillus subtilis*, *Micrococcus luteus* and *S. aureus* and also against gram negative bacteria like *Shigella flexneri*, *Proteus vulgaris*, *S. typhi*, *E. coli*, *Vibrio cholerae*, *Serratia marcescens*, *Klebsiella pneumoniae* and *Pseudomonas aeruginosa*. Inhibition zone diameter was calculated to determine the extract's antibacterial activity against the test organisms. This property was attributed to the presence of phytochemicals such as flavonoids, sterols, phenols, alkaloids, tannins, quinones, saponins etc. It was noted that the extracts such as methanol extract, ethylacetate extract, chloroform extract, aqueous extract and ethanol extract showed antibacterial activity.

B. diffusa provided protection against pathogenic fungi and bacteria of humans was investigated by Wagh and Vidhale (2010). Decoction prepared from *B. diffusa* roots was effective against all the gram negative bacteria tested i.e. *S. typhi*, *P. vulgaris*, *E. coli*, *Enterobacter aerogenes*, *P. aeruginosa*, *Salmonella typhimurium* and *K. pneumoniae*. *Enterococcus faecalis* was the only gram positive bacterium which was sensitive to the decoction. Fungus, *Candida glabrata* showed high sensitivity to the decoction. The studies showed broad spectrum antimicrobial efficacy of *B. diffusa*.

In vitro experiments were performed by Ramachandra *et al.* (2012) to study antibacterial potential of *B. diffusa*. Chloroform, petroleum ether and methanol extracts were prepared from *B. diffusa* roots and aerial parts. The antibacterial efficacy of the extracts were tested against six bacteria types like *Agrobacterium tumefaciens*,

B. subtilis, *E. coli*, *P. aeruginosa*, *S. aureus* and *K. pneumoniae* by using agar well plate method. In case of the extracts prepared from the aerial parts, the methanol extract had strong antibacterial property and the most susceptible bacteria were *E. coli*. Methanol extract prepared from the roots was most effective against *S. aureus*. So the experiments revealed that the whole plant of *B. diffusa* possess antibacterial properties.

Protection against harmful radiations

B. diffusa has the potential for protection against damage induced by exposure to radiations. To investigate it Manu *et al.* (2007) prepared hydroalcoholic extracts of *B. diffusa* and tested their effects using in vivo mice models. Mice were given whole body exposure of single acute dose of gamma radiation (600 rads) and were treated intraperitoneally with the extract (20 mg/kg). Estimation of intestinal glutathione, maturing monocytes and bone marrow cellularity indicated that the extract provided considerable protection to most affected tissues (intestine and bone marrow). The extract administration also normalized the total white blood cell count which was drastically lowered down by the radiation exposure. Levels of liver and serum alkaline phosphatase in mice were elevated by the radiation exposure. Their levels were reduced after treatment with *B. diffusa* extract. The irradiated animals also had increased lipid peroxidation level in serum as well as in liver, but *B. diffusa* extract administration significantly lowered down the lipid peroxidation level. All the results proved that whole plant of *B. diffusa* has potential for protection against radiation damage which may be attributed to improved immune status of irradiated mice after treatment with *B. diffusa* extract (Manu *et al.* 2007).

Hepatoprotective activity

Madagundi *et al.* (2016) isolated the endophyte *Bacillus cereus* from the roots of *B. diffusa*, prepared ethyl acetate and chloroform extracts from it and studied their hepatoprotective activities in rats. The extracts were studied for free radical scavenging activities. Out of the two, the ethyl acetate extract had significant IC₅₀ value and was used for further in vivo studies. Liver disorders were induced in rats by oral administration of CCl₄ (hepatotoxic chemical) to them. CCl₄ administration increased the levels of total and direct bilirubin, serum alkaline phosphatase, triglycerides, serum

glutamic oxaloacetic transaminase and serum glutamate pyruvate transaminase in comparison to normal control. Treatment with the extracts of the endophytic bacteria reversed the elevated levels of all the above biochemicals. The extracts also maintained or increased the activity of enzymes like catalase and superoxide dismutase which play role in combating reactive oxygen species. All the results led to the conclusion that the extracts prepared from the endophytic bacteria of *B. diffusa* exhibit antioxidant and hepatoprotective functions.

Role of *B. diffusa* in curing hepatotoxicity induced by antituberculosis drug rifampicin in male albino Wistar rats was investigated by Muthulingam (2014). Tuberculosis is a major health problem and to treat it rifampicin therapy is given. Rifampicin cures tuberculosis but causes liver damage. To find a herbal cure for it Muthulingam (2014) first made methanolic extracts from leaves of *B. diffusa*. Hepatotoxicity was induced by administering rifampicin to the rats. It caused a substantial enhancement in the activities of aspartate aminotransferase, gamma-glutamyl transpeptidase, bilirubin, lactate dehydrogenase, alanine aminotransferase and alkaline phosphatase whereas it decreased the amount of protein in serum as compared to the control. When the rats were given the methanolic extract of *B. diffusa* orally, rifampicin toxicity was alleviated as was evident from the nearly normalized levels of the above biochemicals. The extract having concentration 200 mg/kg body weight was the most effective one. So the experiments provide evidence that *B. diffusa* extracts can heal liver problems which was supposed to be due to the presence of flavonoids in the plant which have antihepatotoxic characteristics.

Anti-arthritic activity

Parmer *et al.* (2018) evaluated anti-arthritic effects of chloroform, water, methanol and petroleum ether extracts prepared from roots of *B. diffusa* in arthritic rats. Arthritis was induced in rats by injecting them with Freud's Complete Adjuvant. Arthritic rats showed increased paw volume and decreased body weight. Arthritis also influenced hematological parameters such as increase in total WBC counts, increase in erythrocyte sedimentation rate and reduced RBCs and haemoglobin. Treatment of rats with the *B. diffusa* extracts decreased paw volume and increased body weight. Various

hematological parameters were also normalized like total WBC count became normal, erythrocyte sedimentation rate reduced and recovered RBCs and haemoglobin. Methanol extract gave the best results. Anti-arthritic activities of *B. diffusa* root extracts were correlated with its flavonoids constituents as revealed by phytochemical analysis.

Anti-diabetic property

Indigenous system of medicine in India has recommended many herbal medicines including *B. diffusa* for curing diabetes (Madhumeha) (Nalamolu *et al.* 2004). With the aim to prove and support the traditional usage of *B. diffusa* for diabetes treatment, many research works have been done. Diabetic (Type I and Type II) patients with protein urea (\bar{A} 500 mg/day), were studied by Singh *et al.* (2010) for antiproteinuric effect of *B. diffusa*. The patients were given *B. diffusa* powder for six months. It was observed that there was some decrease in 24 hour urine protein in the patients after six months follow up.

Nalamolu *et al.* (2004) performed experiments on diabetic rats. Non-insulin dependent diabetes mellitus was induced in rats by injecting streptozotocin. Chloroform extracts of *B. diffusa* leaves were prepared and were orally administered (50, 100 and 200 mg/kg body weight) to rats daily for four weeks. Blood glucose was measured by glucose-oxidase method at weekly intervals for four weeks. Administration of the extracts of *B. diffusa* lowered down the blood glucose levels in the diabetic rats in a dose dependent manner. The extract anti-diabetic activity was comparable to that of the anti-diabetic drug glibenclamide. Streptozotocin induces a massive damage to β -cells of islets of Langerhans and lower down the synthesis and release of insulin. Anti-diabetic action of the extract was attributed to its role in rejuvenation of pancreatic β -cells. Anti-diabetic activity of *B. diffusa* has also been investigated by Chude *et al.* (2001) and Pari and Satheesh (2004).

Antihyperglycaemic and renoprotective effects of ethanolic extracts of *B. diffusa* were evaluated by Singh *et al.* (2011) in diabetic rats. Diabetic animals showed reduced $\text{Na}^+\text{-K}^+$ ATPase activity, altered ionic balance and glycaemic dysregulation. Administration of *B. diffusa* ethanolic extract maintained renal $\text{Na}^+\text{-K}^+$ ATPase activity and ionic balance and resulted in significant reduction in diabetic hyperglycaemia. The effects

were comparable to the effects of standard hypoglycaemic drug metformin.

CONCLUSION

The bioactive compounds like flavonoids, alkaloids, phenolic compounds, saponins, tannins, terpenoids, glycosides, steroids etc. are present in the extracts prepared from whole plant of *B. diffusa* or its individual parts like leaves, stem and roots. These bioactive compounds are responsible for the healing effects of *B. diffusa* against a number of human ailments like cancer, diabetes, inflammation, harmful radiations induced damage, gastrointestinal problems, microbial infections, prostatic hyperplasia, liver disorders, cardiac problems, anxiety problem etc. So, *B. diffusa* which is used in traditional medicines, now its pharmacological potential is validated by a number of scientific experiments. But, further extensive research is needed to increase the acceptance and inclusion of *B. diffusa* in mainstream medicines.

REFERENCES

1. Aviello G, Canadanovic-Brunet J. M, Milic N, Capasso R, Fattorusso E, Tagliamonte Scafati O, Fasolino I, Izzo A. A and Borrelli F. Potent antioxidant and genoprotective effects of boeravinone G, a rotenoid isolated from *Boerhaavia diffusa*. *PLoS ONE.*, **6**(5): e19628 (2011). doi:10.1371/journal.pone.0019628
2. Chaudhary G and Dantu P. K. Analysis of some important genes from the trichomes of *Boerhaavia diffusa* L. *Euro. J. Exp. Bio.*, **8**(4):25 (2018). doi: 10.21767/2248-9215.100066
3. Chauhan A, Semwal D. K, Mishra S. P and Semwal R. B. Ayurvedic research and methodology: Present status and future strategies. *AYU.*, **36**: 364-369 (2015).
4. Chude M. A, Orisakwe O. J, Afonne O. J, Gamaniel K. S, Vongtau O. H and Obi E. Hypoglycemic effect of the aqueous extract of *Boerhaavia diffusa* leaves. *Indian. J. Pharmacol.*, **33**: 215-216 (2001).
5. Devi K. M and Jyothi Y. Pharmacodynamic interaction of *Boerhaavia diffusa* with omeprazole in experimentally induced ulcers in rats. *Indian J. Pharm. Biol. Res.*, **3**: 56-63 (2015).
6. Ferreres F, Sousa C, Justin M, Valentao P, Andrade P. B, Llorach R, Rodrigues A, Seabra R M and Leitao A. Characterisation of the phenolic profile of *Boerhaavia diffusa* L. by HPLC PAD

- MS/MS as a tool for quality control. *Phytochem. Anal.*, **16**: 451-458 (2005).
7. Gadekar D. H, Jain S and Malik J. K. Evaluation of anxiolytic activity of *Boerhaavia diffusa* hydro-alcoholic extract of leaves in rats. *Int. Res. J. Pharm.*, **10**: 90-92 (2011).
 8. Jana J. C. Use of traditional and underutilized leafy vegetables of sub-himalayan terai region of West Bengal. *Acta Hort.*; **752**: 571-575 (2007).
 9. Kadota S, Lami N, Tezuka Y and Kikuchi T. *Chem. Pharm. Bull.*, **37**: 3214-3220 (1989).
 10. Kayande N and P Kushwah P. In vitro and in vivo evaluation of anticancer activity of *Boerhaavia Diffusa* Linn in experimental animals. *PharmaTutor.*, **2**: 133-137 (2014).
 11. Kim D. S, Kim H. R, Woo E. R, Kwon D. Y, Kim M. S, Chae S. W and Chae H. J. Protective effect of calceolarioside on adriamycin-induced cardiomyocyte toxicity. *Eur. J. Pharmacol.*, **541**: 24-32 (2006).
 12. Kirtikar K. R and Basu B. D. Indian Medicinal Plants. Periodical Experts Book Agency, Delhi. (1991).
 13. Krishnamoorthy P. K. P, Muthukumaran S, Maheswaran A and Sukumaran P. Isolation, purification and characterization of boeravinone B from *Boerhaavia diffusa* Linn. *Int. Res. J. Pharm.*, **8**: 140-144 (2017).
 14. Lad V. The complete book of ayurvedic home remedies. New York: Three Rivers Press. (1999).
 15. Lami N, Kadota S, Tezuka Y and Kikuchi T. *Chem. Pharm. Bull.*, **38**: 1558-1562 (1990).
 16. Lee M. Management of benign prostatic hyperplasia. In: DiPiro JT, Pharmacotherapy: A Pathophysiologic Approach. 7th ed. South Carolina: McGraw Hill Companies, pp 1387-1398 (2008).
 17. Madagundi S. D, Kothli P, Habbu P. V and Kulkarni V. H. Evaluation of endophytic fractions of *Boerhaavia diffusa* Linn. roots for hepatoprotective activity in rats. *BLDE Univ. J. Health Sci.*, **1**: 113-119 (2016).
 18. Majgaine S and Verma D. L. Antibacterial activity of *Boerhaavia diffusa* L. (Punarnava) on certain bacteria. *IOSR J. Pharm.*, **7**: 1-13 (2017).
 19. Manu K. A, Leyon P. V and Kuttan G. Studies on the protective effects of *Boerhaavia diffusa* L. against gamma radiation-induced damage in mice. *Integr. Cancer Ther.*, **6**: 381-388 (2007).
 20. Mishra S, Aeri V, Gaur P. V and Jachak S. M. Phytochemical, therapeutic, and ethnopharmacological overview for a traditionally important herb: *Boerhavia diffusa* Linn. Hindawi Publishing Corporation BioMed Research International. 2014; Article ID 808302:1-19.
 21. Muthulingam M. Antihepatotoxic role of *Boerhaavia diffusa* (Linn.) against antituberculosis drug rifampicin induced hepatotoxicity in male albino Wistar rats. *J. Pharm Res.*, **8**: 1226-1232 (2014).
 22. Nalamolu R. K, Boini K. M and Nammi S. Effect of chronic administration of *Boerhaavia diffusa* Linn. leaf extract on experimental diabetes in rats. *Trop. J. Pharm. Res.*, **3**: 305-309 (2004).
 23. Nayak P and Thirunavoukkarasu M. A review of the plant *Boerhaavia diffusa*: Its chemistry, pharmacology and therapeutical potential. *J. Phytopharmacol.*, **5**: 83-92 (2016).
 24. Nimbal S. K and Koti B. C. Cardio protective effect of *Boerhaavia diffusa* against doxorubicin-induced myocardial toxicity in albino rats. *Sch. Acad. J. Biosci.*, **4**: 171-178 (2016).
 25. Pari L and Satheesh M. A. Antidiabetic activity of *Boerhaavia diffusa* L.: Effect on hepatic key enzymes in experimental diabetes. *J. Ethnopharmacol.*, **91**: 109-113 (2004).
 26. Parmar D, Jain N. K and Tomar V. Anti-arthritic evaluation of different extracts of *Boerhaavia diffusa* Linn. in FCA induced arthritis in rats. *J. Drug Deliv. Ther.*, **8**: 388-393 (2018).
 27. Pooja, Lal V. K and Verma A. GC-MS and phytopharmacological analysis of aqueous distillate of *Boerhavia diffusa* roots. *Int. J. Pharm. Pharm. Res.*, **10**: 374-391 (2017).
 28. Ramachandra Y. L, Ashajyothi C and Rai S. P. In vitro antibacterial potential of *Boerhaavia diffusa*. *Int. J. Adv. Pharm., Biol. Chem.*, **1**: 420-423 (2012).
 29. Rang H. P, Dale M. M, Ritter J, Flower R. J. Rang and dale's Pharmacology, 6th Ed. Elsevier Publication, pp 226-245 (2008).
 30. Remya M. J, Hameed A. S and Sujathan K. Cytotoxicity of Punarnava (*Boerhaavia diffusa* L.) in breast cell line. *Int. J. Ayur. Pharma Research.*, **6**: 1-5 (2018).
 31. Scott L. D and Ellis T. M. Small intestinal transit and myoelectric activity in diabetic rats. In: Gastrointestinal Motility. J Christensen (ed). New York, Raven Press, pp 395-399 (1980).
 32. Shisode K. S and Kareppa B. M. In-vitro antioxidant activity and phytochemical studies of *Boerhaavia diffusa* Linn. roots. *Int. J. Pharm. Sci. Res.*, **2**: 3171-3176 (2011).
 33. Singh A, Sharma H, Singh R, Pant P, Srikant N and Dhiman K. S. Identification and quantification of boeravinone-B in whole plant extract of *Boerhaavia diffusa* Linn and in its polyherbal formulation. *JNR.*, **17**: 88-95 (2017).
 34. Singh P. K, Baxi D, Doshi A and Ramachandran A. V. Antihyperglycaemic and renoprotective effect of *Boerhaavia diffusa* L. in experimental diabetic rats. *Journal of Complementary*

- and Integrative Medicine*. **8**(1) (2011) doi. org/10.2202/1553-3840.1533.
35. Singh R. G, Kumar G, Singh S. K, Tripathi Y. B, Singh R. H and Usha. Evaluation of antiproteinuric and renoprotective effect of Punarnava (*Boerhaavia diffusa* Linn.) in diabetic nephropathy. *J. Res. Educ. Indian Med.*; **16**: 45-58 (2010).
36. Sudhamadhuri A and Kalasker V. Evaluation of anti-inflammatory effect of aqueous extract of *Boerhaavia diffusa* leaves in rats. *Int. J. Res. Health Sci.*, **2**: 517-521 (2014).
37. Umamaheswari A, Nuni A and Shreevidya R. Evaluation of antibacterial activity of *Boerhaavia diffusa* L. leaves. *Int. J. Green Pharm.*, **4**: 75-78 (2010).
38. Vyas B. A, Desai N. Y, Patel P. K, Joshi S. V and Shah D. R. Effect of *Boerhaavia diffusa* in experimental prostatic hyperplasia in rats. *Indian J. Pharmacol.*, **45**: 264-269 (2013).
39. Wagh S and Vidhale N. N. Antimicrobial efficacy of *Boerhaavia diffusa* against some human pathogenic bacteria and fungi. *Biosci. Biotech. Res. Asia.*, **7**: 267-272 (2010).