



## The Antioxidant Study of an Ayurvedic Medicine, Balarishtam

Sivasankar Reddy Konda<sup>1</sup>, Mudiganti Ram Krishna Rao<sup>2\*</sup>, Minu Priya<sup>3</sup>, K. Prabhu<sup>4</sup>, V. S. Kalaivani<sup>5</sup>, D. Kumaran<sup>6</sup>, Ayub Alam<sup>7</sup>, Kumari Sangeeta Singh<sup>8</sup>, Lakshmi Sundaram<sup>9</sup>

<sup>1</sup>Post Graduate Student, Sree Balajee Medical College and Hospital, Chennai, Tamil Nadu, India.

<sup>2</sup>Professor, Department of Industrial Biotechnology, Bharath University, Chennai, Tamil Nadu, India.

<sup>3</sup>Associate Professor, Department of Gynecology, Sree Balajee Medical College and Hospital, Chennai, Tamil Nadu, India.

<sup>4</sup>Associate Professor, Department of Anatomy, Sree Balajee Medical College and Hospital, Chennai, Tamil Nadu, India.

<sup>5</sup>Professor, Department of Biochemistry, Sree Balajee Medical College and Hospital, Chennai, Tamil Nadu, India.

<sup>6</sup>Student, Sree Balaji Medical College and Hospital, Bharath University, Chennai, Tamil Nadu, India.

<sup>7</sup>Student, Department of Industrial Biotechnology, Bharath University, Chennai, Tamil Nadu, India.

<sup>8</sup>Student, Department of Industrial Biotechnology, Bharath University, Chennai, Tamil Nadu, India.

<sup>9</sup>Scientist, V Clin Bio, Central Research Facility, Sri Ramachandra University, Chennai, Tamil Nadu, India.

\*Corresponding author's E-mail: [mrk Rao1455@gmail.com](mailto:mrk Rao1455@gmail.com)

Received: 13-07-2016; Revised: 30-11-2016; Accepted: 18-12-2016.

### ABSTRACT

Balarishtam is an Ayurvedic formulation used for the treatment of diseases caused due to Vata imbalance such as brain related dysfunctions. The present study deals with the antioxidant activities such as DPPH assay, FRAP assay and Hydrogen Peroxide Scavenging Activities of Balarishtam. The results indicated that Balarishtam has very good antioxidant properties which could be one of the mechanisms responsible for the treatment of such diseases.

**Keywords:** Balarishtam, DPPH, FRAP, Hydrogen Peroxide Scavenging, Antioxidant.

### INTRODUCTION

Ayurveda and sidha are traditional medical practices of India which are an age old and time tested modes of treatment.

Modern medical practice is beset with problems like, Adverse Drug Reactions (ADR), major side effects and multidrug resistant pathogenic (MDR) strains.

Thus there is an urgent need to re-evaluate the traditional medical systems to come to rescue for the safe treatment of mankind.

The traditional systems of medicine are good candidates to serve the purpose being safe, more natural, easily available and affordable. But this system requires thorough efficacy evaluation in the light of modern medical standards and parameters to prove or disprove the claims that they are safe with less side effects. It is heartening that since last decade or two many reports in this regard have come and this is welcome sign.<sup>1-5</sup>

Balarishtam is an Ayurvedic preparation made out of nine different plants and jaggery and is used for the treatment of diseases caused due to Vata imbalance in the body, like neuralgia, hemiplegia, paraplegia, arthritis, spondylosis etc. and works as a nerve tonic.

This is also used for strengthening muscles and bones.

This medicine finds reference in the Ayurvedic treatise, Bhaishaja Ratnavali Vatavydhi-569-572.

Balarishtam is taken 12-24 ml once or twice daily after food or as per the advice of prescribed Ayurvedic medicine practitioner.

**Balarishtam ingredients: Balarishtam is prepared by the following ingredients.**

Bala (*Sida cordifolia*) – root / whole plant – 4.8 kg

Ashwagandha (*Withania somnifera*) – root – 4.8 kg

Water for decoction – 49.152 liters, boiled and reduced to 12.288 liters.

Jaggery – 14.400 kg

Dhataki (*Woodfordia fruticosa*) – flower – 768 g

Payasya (*Ipomea digitata*) – root / whole plant 96 g

Eranda (*Ricinus communis*) – root – 96 g

Rasna (*Pluchea lanceolata*) – root -48 g

Ela (Cardamom) (*Elettaria cardamomum*) – 48 g

Prasarini (*Paederia foetida*) – root – 48 g

Usheera (*Vetiveria zizanioides*) – 48 g

Gokshura (*Tribulus terrestris*) – whole plant / fruit – 48 g

### Method of Manufacturing

Coarse powders of Bala and Ashwagandha are added to water, boiled and filtered. To this Kashayam, jaggery is added, filtered for impurities. Rest of the ingredients is added and kept closed in an air tight container for a month time.



After fermentation, the liquid is filtered and stored in air tight container. The standard manufactures of Balarishtam are Dabur, Baidyanath, Arya vaidya Sala, AVP etc.

Although it is a standard medicine, not much scientific documentation about its efficacy in terms of pharmacology, pharmacokinetics, toxicology etc. is available.

Rajalakshmy and Sindhu, 2011, have reported the primary phytochemical and antioxidant activity of Balarishtam.<sup>6</sup> Tiwari, 2014 has given a comparative account of the antihyperlipidemic activities of Balarishtam prepared by traditional and modern methods.<sup>7</sup> The present study deals with the antioxidant activity of Balarishtam by some standard methods. This is the first step of our work to prove the scientific efficacy of this medicine.

A general account of the medicinal values of constituent ingredients plants of Balarishtam is mentioned below.

#### **Bala (*Sida cordifolia*)**

This is known as Bala in Ayurvedic literature and is supposed to control all the three doshas namely, Vata, Pitta and Kapha, although it works predominantly on Vata. This plant contains Ephedrine, hypaphorine, vasicinone, choline, betaine, phytosterol etc. and the roots are rich source of  $\beta$ - sitosterols, known for their immunomodulatory role.

Sharma, 2013 and Jain have reviewed the medicinal values of this plant.<sup>8,9</sup> The CNS pharmacological effect of *Sida* extracts was reported by Franco.<sup>10</sup> Its anti-inflammatory and analgesic activities were studied by Franzotti.<sup>11</sup> Mediros have studied the role of *Sida* on the cardiovascular system.<sup>12</sup> Sharma have worked on the hepato-protective activity of *Sida cordifolia*.<sup>13</sup>

#### **Aswagandha (*Withania somnifera*)**

This is considered to be a wonder drug in Ayurveda for its numerous medicinal values. It has activities like immunomodulatory, cardioprotective, anticancer and as a rasayana (rejuvenant).<sup>14-21</sup>

#### **Dhataki (*Woodfordia fruticosa*)**

It was reported by Dubey that the presence of therapeutically potent antimicrobial compounds against MDR bacteria in *Woodfordia fruticosa* and the crude leaf extract had no host toxicity on human lymphocytes.<sup>22</sup>

The n-butanol fraction of the extract was the most suitable bio-active fraction. The terpenes isolated were, phenol, 5-methyl-2-(1-methylethyl)-, phenol, 2-methoxy-4-(2-propenyl)-, 2, 6-octadien-1-ol, 3, 7-dimethyl-(E)-, 2, 6-octadienal, 3, 7-dimethyl-, cyclohexanol, and 2-methylene-5-(1-methylethenyl).

The leaves have sedative properties and the juice of its fresh flowers, when applied on the head, supposed to reduce headache.

The curative properties of *Woodfordia* are due to the presence of secondary metabolites like alkaloids, flavonoids, glycosides, phenols, saponins, sterols etc. Grover and Patni, 2013 have identified 21 compounds in the GC MS analysis of *Woodfordia* leaf extracts with important medicinal properties.<sup>23</sup>

#### **Payasya or Vidari (*Ipomea digitata*)**

This plant is reported to have activities like hypolipidemic, cardioprotective, and antidiabetic. The root is used by natives for gynecological disorders.<sup>24-26</sup>

#### **Erand (*Ricinis communis*)**

The oil of Erand is commonly used in India as purgative for children and also as lamp oil. The medicinal role of Erand has been reported by many researchers. Rachhadiya have reported the cytoprotective role of the oil of Erand on gastric mucosa thus reducing the risk of ulcer formation.<sup>27</sup> Castor oil is shown to have lypolytic, antidiabetic, antibacterial, anti-inflammatory, wound healing, hepatoprotective and antioxidant activities.<sup>28-37</sup>

Darmanin have reported the cytotoxic and apoptotic activity of castor.<sup>38</sup>

#### **Rasna (*Pluchea lanceolata*)**

It was observed that *Pluchea lanceolata* is a very good antioxidant and immunosuppressant.<sup>39,40</sup>

#### **Ela (*Elettaria cardamomum*)**

Cardamom is another important culinary ingredient used for its characteristic aroma. Apart from the aroma it has medicinal value. Verma have reported blood pressure lowering, fibrinolysis enhancing and antioxidant activities of Cardamom.<sup>41</sup> Khan have shown the pharmacological basis of cardamom as medicine for asthma.<sup>42</sup>

#### **Prasarini (*Paederia foetida L.*)**

This plant is reported to have antimicrobial, thrombolytic, cytotoxic and antidiabetic activities.<sup>43-45</sup> The leaf extracts showed protective effect on induced colitis in animal model (Das).<sup>46</sup>

#### **Usheera - Vetiver (*Vetiveria zizanioides*)**

*Vetiveria zizanioides* is a densely tufted grass. Vetiver oil is supposed to have a nerve relaxant reducing mental stress. Chemical components of *Vetiveria* roots have very high fungicidal, bactericidal and insecticidal properties.<sup>47-50</sup> The oil is reported to be carminative in flatulence, colic and obstinate vomiting. It is regarded as a stimulant, diaphoric, refrigerant, astringent and antimicrobial. When applied externally it removes excess heat from the body and gives cooling effect. The decoction of the roots is believed to dissolve kidney stones and the paste made from pounded fresh roots is considered as an abortifacient. Medicinally this plant oil is reported to be used as a carminative in flatulence and colic, as antipyretic and as anthelmintic.<sup>51-52</sup> Bhushan and Sharma in their reviews have described the various medicinal properties of *V. zizanioides*.<sup>53,54</sup>



**Gokshura (*Tribulus terrestris*)**

*Tribulus* is known as Gokshura in Ayurveda. It is an ancient herb with immense medicinal qualities. *Tribulus*, in modern day, is used for body building, to relieve diseases of uro-genital system and as an aphrodisiac. Fatima have elaborated in their review on the various pharmacological activities of *Tribulus*<sup>55</sup>. This plant has various medicinal applications such as diuretic, antitumor, antibacterial and antifungal, antioxidant and hypoglycemic.<sup>56-61</sup>

**MATERIALS AND METHODS**

Balarishtam was procured from standard Ayurvedic shop from Chennai. Antioxidant studies, namely, DPPH assay, FRAP assay, Hydrogen Peroxide scavenging activity assay and were conducted by standard methods.

**Antioxidant Study**

Antioxidant study was performed by **DPPH Assay, FRAP Assay and Hydrogen Peroxide Scavenging Activity assay.**

**DPPH Assay (1,1-diphenyl-2-picrylhydrazyl) (Bliss, 1958)<sup>62</sup>**

The sample was dissolved in Ethanol in 1mg/ml concentration and used as stock. From the stock, various concentrations (100, 200, 300, 400mg) were taken for further analysis.

Respective solvents were taken as negative control.

Conc.	= Concentration of the sample
OD	= OD of the sample
Neg. Control	= The Solvent
Activity	= Neg. Control – OD / Neg. Control
% of Activity	= Activity/100
IC50	= 50 – c value / m value

**RESULTS AND DISCUSSION****Table 1:** Indicates the Results of DPPH Assay with Ethanol for Balarishtam

S. No.	Solution	Conc.	OD	Neg. Control	% Activity	m value	C value	IC50	IC50/ml
1	Ethanol	100	0.506	0.989	48.83721	0.1221	3.1951	383.3325	127.7775
2		200	0.370		62.58847				
3		300	0.350		64.81072				
4		400	0.213		78.46309				

From the results it shows that IC50/ml was lowest value (127.7775) indicating highest activity.

FRAP test Results are mentioned in Table 2.

**Table 2:** Indicates the FRAP Assay Patterns of Balarishtam in Ethanol Solution

Solvent	Conc.	OD	M Value	C Value	mM	Fe(II)/mg	Mean	STDEV
Ethanol	10	0.278	0.0274	0.1086	6.182482	61.8248152		
	10	0.267	0.0274	0.1086	5.781022	57.8102689		
	10	0.256	0.0274	0.1086	5.379562	53.7956204	57.81	4.01

IC50/ml = IC50/3 (3 ml of DPPH for the assay. To find the activity in 1 ml, the value had been divided by 3).

**FRAP Assay (Ferric Reducing/Oxidant Power) (Pulido, 2000)<sup>63</sup>**

Balarishtam was dissolved in Ethanol. Triplicates had been put for all the Processes.

Conc.	= Concentration of the sample
OD	= OD of the sample
Linearity (y)	= mx + c
M	= Slope
C	= The point x crosses y axis
X	= OD – c value / m value
mM Fe/mg	= X value / concentration x 1000
Mean	= Average of mM Fe/mg
STDEV	= Standard Deviation for mM Fe/mg.

**Hydrogen Peroxide Scavenging Activity (Ruch 1989)<sup>64</sup>**

Balarishtam was dissolved in Ethanol.

Triplicates had been put for all the Processes.

Conc.	= Concentration of the sample
OD	= OD of the sample
Neg. control	= The solvent
Activity	= Negative control – OD / Negative control
% of activity	= Activity / 100
Mean	= Average of % of Activity
STDEV	= Standard Deviation of % of Activity
Graph	= (For Mean of % of Inhibition vs samples) Drawn using 2D clustered column.



From the Table 2 It is clear that ethanol solution of Balarishtam indicated antioxidant activity (57.81).

Hydrogen peroxide scavenging assays results of Balarishtam are mentioned in Table 3.

**Table 3:** Indicates the Hydrogen Peroxide Scavenging Activity of Balarishtam

Solvent	Conc.	OD	Neg. Control	% Activity	Mean	STDEV
Ethanol	100	0.587	0.748	21.52406		
	100	0.487	0.748	34.89305		
	100	0.429	0.748	42.64706	33.02	10.69

From Table 3 it is clear that Balarishtam has antioxidant activity as averaged to 33.02 % with regard to Hydrogen peroxide scavenging along with other medicinal functions.

The antioxidant results of Balarishtam obtained in this report conforms to such activities by the constituent plant parts. It is a known fact that the role of Reactive Oxygen Species in the manifestation of a disease is very high.

The ROS is generated in the body due to a number of reasons like life style, eating habits, lack of proper exercise, intoxication, pollution, allergies etc. It is thus quite obvious to have high amounts of ROS in the body. Balarishtam seems to play a vital role in reducing the ROS in the body, as suggested in this report, which could be one important mechanism of cure of diseases for which this medicine is prescribed.

## CONCLUSION

From the results obtained it could be concluded that Balarishtam has very good antioxidant activity which could be one of the curative properties of this medicine. Further work to understand the mechanism of action is needed.

## REFERENCES

- Rao MRK, Kumar MH, Amutha A, Prabhu K, Chatterjee B, Selva Kumar S. Phytochemical Analysis and Antioxidant Efficacy of the Resin of *Bombax ceiba* (Salmali). Int J Pharm Sci Rev Res, 30(1), 2015, 335-339.
- Rao MRK, Phillips S, Kumar MH, Saranya Y, Divya D, Prabhu K. GC MS analysis, antimicrobial, antioxidant activity of an Ayurvedic medicine, Salmali Nirayasa. Journal of Chemical and Pharmaceutical Research, 7(7), 2015, 131-139.
- Ravi A, Jai Prabhu SP, Rao MRK, Prabhu K, Kalaiselvi VS, Saranya Y. Identification of Active Biomolecules in Saraswatarishtam (An Ayurvedic Preparation) by GC-MS Analysis. Int J Pharm Sci Rev Res, 33(2), 2015, 58-62.
- Chandrasekar T, Rao MRK, Kumar RV, Prabhu K, Nandha Kumar S, Divya D. GC-MS analysis, antimicrobial, antioxidant activity of an Ayurvedic medicine, Nimbapatradi Chooranam. Journal of Chemical and Pharmaceutical Research, 7(8), 2015, 124-136.
- Mudiganti Ram Krishna Rao, Aparna Ravi, Shridhar Narayanan, K. Prabhu, V. S. Kalaiselvi, Shruthi Dinakar, Guru Rajan, N. Kotteswaran. Antioxidant Study and GC MS Analysis of an Ayurvedic Medicine 'Talisapatradi Chooranam'. Int. J. Pharm. Sci. Rev. Res., 36(1), 2016, 158-166
- Rajalakshmy MR, Sindhu A. Preliminary phytochemical screening and antioxidant activity of an ayurvedic formulation: Balarishtam. Int J of Res In Ayurveda and Pharmacy, 2(6), 2011, 1645-1647.
- Tiwari P. Antimicrobial activity of Balarista prepared by traditional and modern methods. Res J of Pharmacy and Technology, 7(7), 2014, 789-791.
- Sharma A K. Medicinal properties of BALA (*Sida cordifolia* LINN. and its species). Int. J. Ayur. Pharma Research, 1(2), 2013, 1-9.
- Jain A, Choubey S, Singour PK, Rajak H, Pawar RS. *Sida cordifolia* (Linn)–An overview. Journal of Applied Pharmaceutical Science, 01(02), 2011, 23-31.
- Franco CIF, Morais LC SL, Quintans-Junior LJ, Almeida RN, Antonioli AR. CNS pharmacological effects of the hydroalcoholic extract of *Sida cordifolia* L. leaves. J Ethanopharmacol, 98, 2005, 275-279.
- Franzotti EM, Santos CV, Rodrigues HM, Mourão RH, Andrade MR, Antonioli AR. Anti-inflammatory, analgesic activity and acute toxicity of *Sida cordifolia* L. (malva-branca). J Ethno pharmacol, 72(1-2), 2000, 273–278.
- Mediros IA., Santos MR., Nascimento NM. and Durate JC. Cardiovascular effects of *Sida cordifolia* L. leaves extracts in rats. Fitoterapia, 77, 2006, 19-27.
- Sharma A, Sangameswaran B, Mahajan SC, Saluja MS. Protective effects of *Sida veronicaefolia* against ethanol induced hepatotoxicity in experimental animals. Phytopharmacology, 3(1), 2012, 137-144
- Aggarwal R, Diwanay S, Patki P, Patwardhan B. Studies on immunomodulatory activity of *Withania somnifera* (Ashwagandha). J App Pharm Sci., 2(1), 2012, 170-175.
- Mohanty I, Arya DS, Dinda A, Talwar KK, Joshi S, Gupta SK. Mechanisms of cardioprotective effect of *Withania somnifera* in experimentally induced myocardial infarction. Basic Clin Pharmacol Toxicol., 94(4), 2004, 184-190.
- Singh N, Verma P, Pandey BR, Gilca M. Role of *Withania somnifera* in prevention and treatment of cancer. Int J of Pharmaceutical Sciences and Drug Research, 3(4), 2011, 274-279.
- 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. 79(4), 2010, 542-551.





18. 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., 72(5), 2010, 659-663.
19. Koduru S, Kumar R, Srinivasan S, Evers MB, Damodaran C. Notch-1 inhibition by Withaferin-A: a therapeutic target against colon carcinogenesis. Mol Cancer Ther. 9(1), 2010, 202-210.
20. Singh N, Bhalla M, Jager P, Gilca M. An Overview on Ashwagandha: A Rasayana (Rejuvenator) of Ayurveda. Afr J Tradit Complement Altern Med. 8(5), 2011, 208-213.
21. Sharma V, Sharma S, Pracheta, Paliwal R. *Withania somnifera*: A Rejuvenating Ayurvedic Medicinal Herb for the Treatment of various Human ailments. International Journal of PharmTech Research, 3(1), 2011, 187-192
22. D Dubey, R Patnaik, G Ghosh, RN Padhy. *In Vitro* Antibacterial Activity, Gas Chromatography–Mass Spectrometry Analysis of *Woodfordia fruticosa* Kurz. Leaf Extract and Host Toxicity Testing With *In Vitro* Cultured Lymphocytes from Human Umbilical Cord Blood. Osong Public Health and Research Perspectives, 5(5), 2014, 298-312.
23. N Grover, V Patni. Phytochemical characterization using various solvent extracts and GC MS analysis of methanolic extract of *Woodfordia fruticosa* (L.) KURZ. Leaves. Int J Pharm Pharm Sci, 5(4), 2013, 291-295.
24. Muthu AK, Alagumanivasagam G, Satheesh Kumar D, Manavalan R. Effect of Methanolic Extract of Tuberous Root of *Ipomoea Digitata* (Linn) on Hyperlipidemia induced by rat fed with high fat diet. Research Journal of Pharmaceutical, Biological and Chemical Sciences, 2(3), 2011, 183-191.
25. Jain V, Verma SK, Katewa SS. Therapeutic validation of *Ipomoea digitata* tuber (Kheervidari) for its effects on cardiovascular risk parameters. Ind J of Trad Knowledge, 10(4), 2011, 617-623.
26. Minaz N, Rao NV, Nazeer A, Preeth M, Shobana J. Antidiabetic Activity of Hydro-Ethanollic Root Extracts of *Ipomoea digitata* in Alloxan Induced Diabetic Rats. International Journal of Research in Pharmaceutical and Biomedical Sciences. 1(2), 2010, 76-81.
27. MR Rachhadiya, PK Mahaveer, VS Rajkumar. Evaluation of antiulcer activity of castor oil in rats. International Journal of Research in Ayurveda & Pharmacy, 2(4), 2011, 1349-1353.
28. HME Lombard, G Pieroni. Lipolytic activity of ricin from *Ricinus sanguineus* and *Ricinus communis* on neutral lipids. Biochem J, 358, 2001, 773-781.
29. P Shokeen, P Anand, Y K Murali Y, V Tandon. Antidiabetic activity of 50% ethanolic extract of *Ricinus communis* and its purified fractions. Food and Chemical Toxicology, 46, 2008, 3458–3466.
30. Mathur A, Verma SK, Yousuf S, Singh SK, Prasad GBKS, Dua VK. Antimicrobial potential of roots of *Ricinus communis* against pathogenic microorganisms. International Journal of Pharma and Bio Sciences, 2(1), 2011, 545-548.
31. T Islam, H Bakshi, S Sam, E Sharma, B Hameed, B Rathore, A Gupta, S Ahirwar, M Sharma. Assessment of antibacterial potential of leaves of *Ricinus communis* against pathogenic and dermatophytic bacteria. International Journal of Pharma Research and Development, 1(12), 2010, 1-7.
32. Saini AK, Goyal R, Gauttam VK, Kalia AN. Antifertility activity of methanol extracts of three different seed varieties of *Ricinus communis* Linn. Journal of Chemical and Pharmaceutical Research, 2(5), 2010, 690-695.
33. Valderramas AC, Moura SHP, Couto M, Pasetto S, CChierice O, Augusto S, Guimarães AC, de Paula Zurron ACB. Anti-inflammatory activity of *Ricinus communis* derived polymer. Braz J Oral Sci, 7(27), 2008, 1666-1672.
34. Prasad MK, Rachhadiya RM, Shete RV. Pharmacological investigation on the wound healing effects of castor oil in rats. International Journal of Universal Pharmacy and Life Sciences, 1(1), 2011, 21-28.
35. Natu MV, Agarwal S, Agarwal SL, Agarwal S. Protective Effect of *Ricinus communis* leaves in Experimental Liver Injury. Indian Journal of Pharmacology, 9(4), 1977, 265-268.
36. Singh RK, Gupta MK, Singh AK, Kumar S. Pharmacognostical investigation of *Ricinus communis* stem. International Journal of Pharmaceutical Sciences and Research, 1(6), 2010, 89-94.
37. Oloyede Ganiyat KG. Antioxidant activities of methyl ricinoleate and ricinolic acid dominated *Ricinus* seed extract using lipid peroxidation and free radical scavenging methods. Research Journal of Medicinal Plant, 6(7), 2012, 511-520.
38. Darmanin S, Wismaver PS, Camilleri Podesta MT, Micallef MJ, Buhagiar JA. An extract from *Ricinus communis* L. leaves possesses cytotoxic properties and induces apoptosis in SK-MEL-28 human melanoma cells. Nat Prod Res, 23(6), 2009, 561-567.
39. Bhagwat DP, Kharya MD, Bani S, Kaul A, Kaur K, Chauhan PS, Suri KA, Satti NK. Immunosuppressive properties of *Pluchea lanceolata* leaves. Ind J of Pharmacol, 42(1), 2010, 21-26.
40. Sharma SK, Goyal N. *In Vitro* Antioxidant Activity of Root Extracts of *Pluchea lanceolata*. J of Pharma and Biomed Sci, 10(18), 2011, 1-3.
41. Verma SK, Jain V, Katewa SS. Blood pressure lowering, fibrinolysis enhancing and antioxidant activities of cardamom (*Elettaria cardamomum*). Indian Journal of Biochemistry and Biophysics., 46(6), 2009, 503-506.
42. Khan A, Khan QJ, Gilani A. Pharmacological basis for the medicinal use of cardamom in asthma. Bangladesh J Pharmacol, 6, 2011, 34-37.
43. Uddin B, Nahar T, Khalil MI, Hossain S. *In vitro* antibacterial activity of the ethanol extract of *Paederia foetida* L. (Rubiaceae) leaves. Bangladesh J. Life Sci., 19(2), 2007, 141-143.
44. Chanda S, Gehlot V, Das R, Mahant S, Das K, Singh K, Ahmad S. Antimicrobial activity of *Paederia foetida* extract against drug resistant *Helicobacter pylori* isolates from India. World J of Pharma Res, 3(4), 2014, 1234-1243.
45. Abu Ahmed AM, Islam Md.M, Rahman Md. A, Hossain Md. A. Thrombolytic, Cytotoxic and Antidiabetic Effects of

- Paederia foetida* L. Leaf Extract. British Journal of Medicine & Medical Research, 4(5), 2014, 1244-1256.
46. Das S, Kanodia L, Mukherjee A, Hakim A. Effect of ethanolic extract of leaves of *Paederia foetida* Linn. on acetic acid induced colitis in albino rats. Pharmacol (series online), 45, 2013, 453-457.
47. Singh G, Singh BS, Kumar BRV. Antimicrobial activity of essential oils against keratinophilic fungi. Indian Drug, 16(2), 1978, 47-51.
48. Devi VS, Kumar KA, Umamaheswar MI, Shanmugam ATS, Anand AR. *In vitro* antibacterial activity of ethanolic extract of *Vetiveria zizanioides* root. Int. J. Pharma Sci. and Res, 1(9), 2010, 120-124.
49. Luqman S, Srivastava S, Darokar M, Khunja SPS. Detection of antimicrobial activities in spent roots of two genotypes of Aromatic grass *V. zizanioides*. Pharma Biol, 43(8), 2005, 732-236.
50. Luqman S, Kumar R, Srivastava S, Darokar M, Khunja SPS. Antioxidant potential of root of *Vetiveria zizanioides* (L) Nash. Ind J of Biochem Biophys, 46, 2009, 122-125.
51. Karan SK, Pal DK, Tarai DK, MishraSK. Analgesic and anthelmintic activity of *Vertiveria zizanioides* root." Journal of Pharmacy Research, 3(4), 2010, 893-894.
52. Narkhede MB, Amire PV, Wagh AE, Bhise MR, Mehetre GD, Patil HJ. An evaluation of antipyretic potential of *Vetiveria zizanioides* (Linn.) root. Res J of Pharmacognosy and Phytochemistry, 4(1), 2012, 11-18.
53. Bhushan B, Sharma SS, Singh T, Singh L, Arya H. *Vetiveria zizanioides* (L.) Nash: a pharmacological overview. Int Res J Pharma, 4(7), 2013, 18-20.
54. Sharma BK, Shamim A, Singh R, Verma RK, Kumar N. *Novel Science International Journal of Pharmaceutical Science*, 1(6), 2012, 308-312.
55. Fatima MSL, SSultana A, Ahmed, Sultana S. Pharmacological activities of *Tribulus terrestris* linn: A systemic review. World J of Pharmacy and Pharmaceutical Sciences, 4(2), 2015, 136-150.
56. Jabbar A, Nazir A, Nilus A, Javed F, Janjua KM. Effects of *Tribulus terrestris* to study on urine output and electrolytes in rabbits. Professional Med J, 19(6), 2012, 843-847.
57. Angelova S, GospodinovaZ, Krasteva M, Antov G, Lozanov V, Markov T. Antitumor activity of Bulgarian herb *Tribulus terrestris* L. on human breast cancer cells. J Bio Sci Biotech, 2(1), 2013, 25-32.
58. Kianbakht S, Jahaniani F. Evaluation of antibacterial activity of L. Growing in Iran. Int J Pharma Tech, 2, 2003, 22-24.
59. Bayati FAA, Al-mola HF. Antibacterial and antifungal activities of different parts of *Tribulus terrestris* L. growing in Iraq. Zhejiang Univ Sci, B, 9(2), 2008, 154-159.
60. Dimitrova DZ, Obreshkova D, Nedialkov P. Antioxidant activity of *Tribulus terrestris* – a natural product in infertility therapy. Int J Pharm Pharm Sci, 4(4), 2012, 508-511.
61. EL-Tantawy WH, Hassanin LA. Hypoglycemic and hypolipidemic effects of alcoholic extract of *Tribulus alatus* in streptozotocin-induced diabetic rats: A comparative study with *T. terrestris*. Indian J Exp Biol, 45(9), 2007, 785-790.
62. Blois MS, 1958. Antioxidant determinations by the use of a stable free radical. Nature, 29, 1958, 1199-1200.
63. Pulido R, Bravo L, Sauro-Calixto F. Antioxidant activity of dietary polyphenols as determined by a modified ferric reducing/antioxidant power assay. J Agri Food Chem, 48, 2000, 3396-3402.
64. Ruch RJ, Cheng SJ, Klaunig JE. Prevention of cytotoxicity and inhibition of intracellular communication by antioxidant catechins isolated from Chinese green tea. Carcinogenesis, 10, 1989, 1003-1008.

Source of Support: Nil, Conflict of Interest: None.

