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Review on Uraria Picta - A Traditionally Medicinal Plant of India: A Herbal Benefaction

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ARTICLE INFO	ABSTRACT
Article history:	In present times, focus on herbal research has increased worldwide. Urariapictais an
Received: 25 April 2017;	important medicinal herb that is extensively used in dasamula and is becoming popular.
Received in revised form:	Flavanoids, alkaloids and pterocarpans are the key constituents of <i>Urariapicta</i> and mainly
15 May 2017;	alleged for its broad beneficial actions. Other than for given treatments, the herb is
Accepted: 25 May 2017;	suggested as remedy for a variety of other ailments. The present review is an effort to
	 provide complete information on Phytochemicals screening, traditional uses and

Keywords

Uraria picta, Description, Phytochemicals, Pharmacology profiles. pharmacology relating to preclinical studies.

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1. Introduction

Urariapicta of Family Leguminosae Sub-Family Fabaceae Is A Well-Liked Ayurvedic Healthful Plant That Additionally Glorious By Name Prsniparni In Sanskrit. It's One Amongst A Part Of Dasamula That Precisely Means That Ten Roots And Is A Well Recognized Ayurvedic Drug Of Indian System Of Medicines Used For Treating General Fatigue, Antioxidant, Analgesic And Anti-Inflammatory Like Medical Conditions [1]. Urariapictaalso Used In Other Ayurvedic Formulations Like Abana, Amrutharishta, Angamardaprashamanakashayachurna, Dasamulataila, Vyaghritaila, Madhyama Narayana Taila, Dasamularishta And Shirashuladivajra Rasa [2, 3]. Other Vernaculars Were Given To Urariapictasuch As Citraparni, Prthakparni, And Simhapuchi Etc. Ayurvedic Texts Clearing Up Its Morphological Characteristics, Attributes And Benefits [4].

Urariapictais A Local Of Tropical Zone Including Nepal, Srilanka, Northern Australia, China, And Burma. This Suffructiose Herb Which Grows Up To 1.5 Meters Tall Is Originate In Dry Grassland, West Places, And Open Deciduous Forests And In All Plains Of India Extending From Himalyas To Ceylon, Malayasia And Phillipines [5]. Urariapicta Consists Of Different Phytoconstituents Present In Different Extracts Exhibit A Number Of Biological Profiles And Guard From Most Of The Chronic Diseases [6, 7].

2. Botanical Description [8]

It Is An Vertical, Small Branched, Perpetual Herb, 90 -180 Cm Tall, Stems With Short, Rough Hairs, Leaves Imparipinnate with 5-9 Leaflets (Lowermost Leaves Often 1-3-Foliolate); Leaflets Narrowly Lanceolate, 7-25 Cm Long (Lowermost Smaller), Often Variegated, Shiny And Hairless Above, Rough Hairy Below; Margins Entire, Inflorescence A Long Terminal Densely Many-Flowered Spike-Like Raceme, Up To 55 Cm Long, Covered In Long Whitish Hairs,

Tele: 0876874707 E-mail address: azmilubna@gmail.com © 2017 Elixir All rights reserved Flowers Pink, Bluish Or Reddish, Fruit 5-9 Mm Long, Folded Into 3-6 Segments, Brown To Black, Turning Greyish-White When Old.



Fig 1. Urariapicta Whole Plant.







Fig 3. Urariapicta Plant Leaves.



Fig 4. Urariapicta Plant Roots. Table 1 Scientific Classification [9]

a	ble 1. Scienti	fic Classification	[9]
	Kingdom	Plantae	
	Subkingdom	Viridaeplantae	
	Super Order	Rosanae	
	Order	Fabales	
	Class	Magnoliopsida	
	Subclass	Rosidae	
	Family	Leguminosae	
	Sub-Family	Fabaceae	
	Genus	Uraria	
	Species	Picta - (Jacq.) Dc	
T	able 2. Verns	acular Names [10]	1

Table 2. Vernacular Names [10].

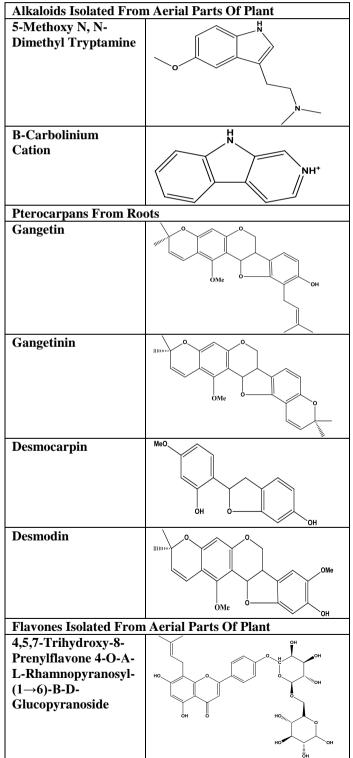
Regions/ Language	Names
Sanskrit	Citraparni, Kalasi, Dhavani,
	Prthakparni, Srgalavinna
Bengali	Salpani, Chhalani, Chakule
Gujarat	Pithavan
Hindi	Pithavan, Dabra
Telugu	Murelehonne, Ondele
	Hone, Prushniparni
Malyalam	Oril
Marathi	Pithavan, Prushniparnee
Oriya	Prushniparnee, Shankarjata
Punjabi	Detedarnee
Tamil	Orpai
Kannadaa	Kolakuponna

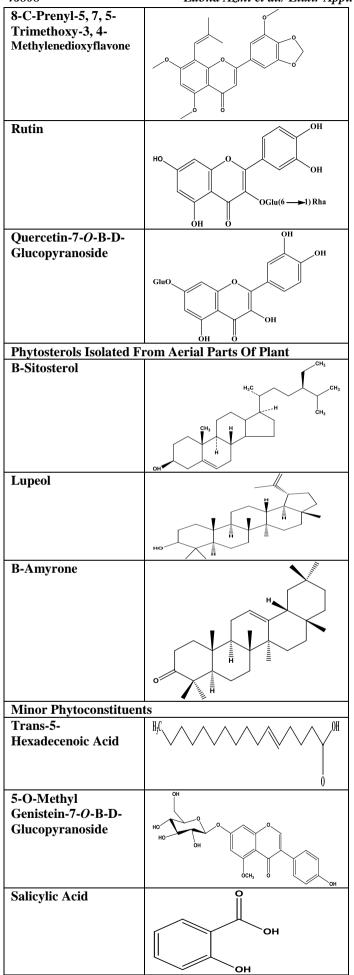
Table 3. Active Constituents [11].

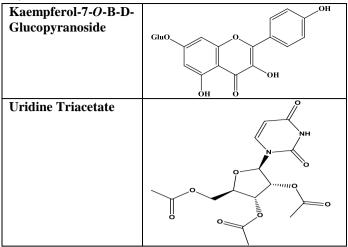
Chemical Nature Of Ingredients	Parts Of Plant
Flavanoids	
Flavones	Stem
Flavonols	Whole Plant
Isoflavones	Aerial Parts
Pterocarpans	Whole Palnt
Alkaloids	
Indole-3-Alkylamines	Whole Plant, Stem
Amide Alkaloids	Whole Plant
Phenylethylamine Alkaloids	Leaf, Stem
Terpenoids	Whole Plant
Steroids	Seed
Phenolic Acid	Aerial Parts
Others	Aerial Parts
Volatile Oils	Roots And Whole Plant
Phenylpropanoids	Aerial Part, Root

Phytochemical Screening Reveals U.Pictais Loaded With Flavonoids, Alkaloids, Steroids, Terpenoids, Phenylpropanoids, Pterocarpans, Coumarins And Volatile Oil [11]. Among The Isolated Compounds Flavonoids, Alkaloids And Pterocarpans Are Considered As Key Bio-Actives. Alkaloids For Example 5-Methoxy N, N-Dimethyl N-Bmethyl-H-4-Harman, **B**-Carbolinium Tryptamine, Cation, Indole-3-Alkyl-Amines Have Been Secluded From Aerial Parts Of The Plant [12]. Pterocarpans For Example Gangetin, Gangetinin, Desmodin, And Desmocarpin Were Reported To Be Present In Roots [13]. Recently A Novel Pterocarpan, Gangetial, Have Been Secluded From The Chloroform Extract Of The Roots Of U. Picta [14]. Flavones Like 4,5,7-Trihydroxy-8-Prenylflavone, 4-0-A-L-Rhamnopyranosyl- $(1\rightarrow 6)$ -B-D-Glucopyranoside, 8-C-Prenyl-5,7,5-Trimethoxy-3,4-Methylenedioxyflavone, Rutin Quercetin-7-O-B-D-Glucopyranoside And Were Also Reported From The Aerial Parts. Phytosterols Viz. B-Sitosterol, B-Amyrone, Lupeol Plus Its Acetate, Stigmastrol Have Been Secluded From Aerial Parts. Additionally, Aminoglucosylglycerolipid Was Reported For The First Time From Seed [15]. More, Minor Phytoconstituents Viz. Trans-5- Hexadecenoic Acid, Salicylic Acid, 5-O-Methylgenistein-7-O-B-D-Glucopyranoside, 3,4- Dihydroxy Benzoic Acid, Kaempferol-7-O-B-D-Glucopyranoside, And Uridine Triacetate Were Reported Additionally [16].

Table 4. Structures Of Active Constituents.







Alkaloids Have Various Pharmacological Actions Like Antiarrhythmic, Anticholinergic, Analgesic, Antitumor, Antihypertensive, Antipyretics, Antimalarial, Stimulant, Anti-Hiv, Antileukemic And Many More And Often Used As Medications. [17]

Flavonoids Are The Mainly Widespread Group Of Polyphenolic Compounds In The Human Diet And Are Found All Over In Plants. Pharmacologically, Flavanoids Embrace Cns, Cardiotonic, Lipid Lowering, Antiulcer, Anti-Inflammatory, Hepatoprotective, Antineoplastic, Antimicrobial, Antioxidant And Hypoglycemic Activities. Relating To Diet, Flavanoids Containing Foods Potentially Lowers The Risk Of Certain Free Radical Related Pathophysiology [18]. Steroids And Triterpenoids Are Pharmacologically Vigorous Compounds And Show The Analgesic Action [19]. The Steroids Also Reveal Central Nervous System Activities. Terpenoids Tend To Decrease Glucose Level In Animals [20]. Cardiac Glycosides Contain Therapeutic Significance And Used In Treating Congestive Heart Failure And Cardiac Arrhythmia [21]. Phenols And Phenolic Compounds Have Remarkable Antimicrobial Potential. They Have Been Wide Utilized In Disinfections And Remained The Standards As Comparison For Other Bactericides [22]. They Point Out Signs Of Cellular Defense Mechanism In Atherogenesis And Cancer. A Wide Variety Of Phenolic Substances Show Strong Antioxidant And Antimutagenic Activities. Current Evidences, Phenolic Compounds Can Also Play An Essential Health Promoting Function [23]. Saponins Were Commercially Used As Dietary Supplements And Nutraceuticals In Traditional Medicine Preparations [24]. They Also Have Hypocholesterolemic And Antidiabetic Actions [25]. Certain Tannins (Ellagitannins From Lagerstroemia Speciosa) Arouse Glucose Uptake. They Reveal Insulin Like Activity Acting As Glucose Transport Activators Of Fat Cells [26]. 3. Traditional Uses

The Plant Traditionally Being Used As Antipyretic, Diuretic, Astringent (Used In Irritable Bowel Syndrome, Diarrhoea And Dysentery), Anticatarrhal, Diuretic, Anthelmintic, Laxative And Nervine Tonic Where As In China*urariapicta* Is Used As Folkloric Medicine Mainly To Treat Fever, Deactivate Toxins, Inhibit Pain, Stimulate Blood Circulation, Suppress Cough And Improve Dyspnea [27].

Table 5. Traditional Uses.

Table 5. Traditional Oses.			
Traditional Use			
Premature Ejaculation [28]			
Treat Mouth Ulcer [29]			
Hair Falling [16]			

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Vera	
Leaves Paste	Eczema Infection, Dermal Disorders [30]
Water Decoction	Type 2 Diabetes Mellitus [31]
Root Powder Boiled	Flatulence [32]
With Milk	
Leaf Decoction	Diarrhea, Dysentery [33]
Leaf Paste	Piles [33]
Root Extract	Whooping Cough [34]
Fresh Leaves Juice	Scabies, Ringworm [35]
Root Extract	Diarrhea, Sedative Agent (Children) [36]
Root And Leaf Pastes	Toothache And Headache [27]
Water Decoction Of	Antipyretic, Anti-Inflammatory And
Root And Aerial Parts	Antinociceptive [36]
Roots Decoction	Asthma And Bronchial Complications
	[37]
Root And Powder	Typhoid Fever, Cerebrospinal Meningitis
Paste	And Antidote Of Snake Venom [38]

4. Pharmacological Profiles

4.1 Antioxidant Profile

Ethanolic Extract Of U. Picta Were Tested In-Vitro To Establish The Antioxidant Potential. The Ic50 Values In Dpph, Abts, O-Phenanthroline, Lipid Peroxidation And Superoxide Scavenging Models Were Found. These Findings Revealed The Antioxidant Profile Of The Extract Against Injury Mediated Through Several Reactive Oxygen Species.

The Antioxidant Profile Of Ethanolic Extract Of U. Pictawas Evaluated In-Vitro. The Results Showed The Presence Of Phenolic, Flavanoids, Sterol And Terpene Derivatives [39].

In-Vivo Free Radical Scavenging Potential Of Aqueous Extract Ofu. Picta Root Was Analysed By Inducement Of Oxidative Stress In Ischemic Reperfused Rat Heart Model. The Reading Supported Antioxidant Capacity Of U. Picta As Compared To Standard Drug Verapamil Against Revascularization Damage [40, 41].

4.2. Anti-Inflammatory And Anti-Nociceptive Profiles

Aqueous Decoction Of Roots And Aerial Parts Ofu. *Picta* Showed Anti-Inflammatory And Anti-Nociceptive Profiles In Dosing Manner. The Inhibition Of Swelling Observed Against Carrageenan Was Equivalent To Cotton Pellet Granuloma. In Addition, Protection Was Observed Against A Significant Increase In Analgesio-Meter-Induced Force And Acetic Acid Induced Writhing Respectively [42].

Whole Plant Juice Of *U. Picta* Possesses Anti-Rheumatic And Anti-Osteo Arthritic Profile By Anti-Inflammatory Profile. The Profile May Be Associated With Several Phytoconstituents Like Polyphenolics, Pterocarpinoid (Gangetin) [43].

4.3. Anti-Leishmanial And Immunomodulatory Profiles

Glyco-Lipids Like Aminoglucosylglycerolipid And Glycosphingolipid, Isolated From The Roots Of *U. Picta* Showed Effectiveantileishmanial And Immunomodulatory Profiles By Increasing Nitric Oxide (No) Production And Provided Resistance Against Infection Recognized In Peritoneal Macrophages By The Protozoan Parasite *Leishmania Donovani* [15].

Additionally, Ethanolic Extract Of *U. Picta* Was Screened Chemoprophylactically And Chemotherapeutically In Opposition To Experimental Visceral Leishmania In Hamsters. Results Revealed Highest Prophylactic **Efficacy** In N-Butanol Fraction And Fairefficacy In Ethanol Extract [44]. **4.4. Cardio-Protective Profile**

Methanolic Extract Of *U. Picta* Roots Defend Mitochondrial Respiratory Enzymes And Thus Defending Rat Heart Against Oxidative Stress Induced Via Reperfusion Injury [45]. Extract Mediates Cardio Protection In Ischemic Reperfusion Injury Model In Isolated Frog Heart. The Effect Was Mediated By Stimulating The G-Protein Coupled Receptors Alike To The Action Of Acetylcholine; The Studies Were Compared Amid Verapamil And Standard Cardioprotective Drug [46]. Pre-Treatment With Aqueous Extract Of *U. Picta* Showed Abridged Cholesterol Level And Free Radical Scavenging Potential Against Isoproterenol Induced Myocardial 46809Nfracted Rats. These Conclusions Were Associated With Cardio-Protective Profile Of The Plant [47].

Ethyl Acetate Extract Of *U. Picta* Root Indicated Potent Cardio-Protection Against Ischemia Reperfusion-Induced Oxidative Stress Models. The Extract Decrease Tbars In Myocardium Together With Better The Recovery Of Antioxidant Enzymes From The Attack Of Ischemic Reperfusion Injury. The Effects Of The Extract May Be Related To The Inhibition Of Lipid Peroxidation [48]. Methanol Extract Of *U. Picta* Root Showed Myocardial Protection In Rat Ischemic Reperfusion Injury Model By Invigorating Muscarinic Receptors. The Profile Might Be Because Of The Decrease In Calcium Overload And Free Radical Release And Enhanced Recovery Of Antioxidant Enzyme Towards Myocardium [49].

4.5. Anti-Ulcer Profile

Ethanolic Extract Of U. Picta When Taken Orally Showed Effective Anti-Ulcerogenic Property In Sprague Dawley Rats And Guinea Pigs. A Significant Defense Against Cold Resistant, Alcohol, Aspirin, Pyloric Ligation And Hst Induced Ulcer Models Were Observed. Fall In Acid Secretion And Increase In Mucin Secretion Was Also Recorded. Results Show Cytoprotective Effect With Anti-Secretory Profile Of U. Picta Which May Be Responsible For Its Anti-Ulcer Property [50].

Ethanolic Extract Of *U. Picta* Root When Taken Orallyshowssignificant Decrease In Ulcer Index And Lesion Number In A Dose Dependent Manner Against Ethanol Induced Acute Gastric Ulcer In Mice. Extract Provokes A Noticeable Increase In Protein And Glutathione Levels, In Comparison To Control [51].

4.6. Cns Profile

Aqueous Extract Of *U. Picta* Showed Effective Anti-Writhing Profile In The Acetic Acid-Induced Abdominal Writhing Assess. It Also Exhibited Fair Cns Depressant Profile And The Effects Of Extract On Locomotion Were Compared With Standard Cns Drugs [52].

4.7. Antiamnesic (Nootropic) Profile

Aqueous Extract Of *U. Picta* Showed Effective Anti-Amnesic Effects In Mice Against Scopolamine Induced Interoceptive Behavioral Models. The Study Was Compared With Piracetama Standard Nootropic Agent [53]. Pre-Treatment With Aqueous Extract Of *U. Picta* For Seven Consecutive Days, Inverted Scopolamine Induced Amnesia In Mice. Study Revealed That The Plant Augmented Mice Brain Acetylcholine Content And Decreased Acetyl Cholinesterase Profile As Compared To Standard Cerebro-Protective Drug Piracetam. Therefore, Aqueous Extract Of *U. Picta* Can Be Used To Wait The Onset And Lessen The Severity Of The Symptoms Of Dementia And Alzheimer's Disease [54].

4.8. Antidiabetic Profile

Methanolic Extract Of Aerial Parts Of *U. Picta* Showed A Significant Antidiabetic Profile In Rats By Raising Insulin Secretion From Min6 And Pseudoislets Cells Of Pancreatic Islet. It Plays A Key Role To Sustain The Lipid Profile Of

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The Rats By Dropping Cholesterol And Triglycerides Levels And Rise In High Density Lipoproteins (Hdl). This Supports The Traditional Use Of *U. Picta* As Anti-Diabetic Drug [55].

4.9. Hepatoprotective Profile

Hepatoprotective Profile Of The Chloroform Extract Roots Of *U. Picta* Was Assessed Against Ccl4 Induced Liver Damage In Rat Models. The Study Concluded That Extract Caused A Rise In Serum Levels Of Total Proteins And Fall In Levels Of Bilirubin, Alanine Aminotransferase (Alt) And Aspartate Aminotransferase (Ast) In Pretreated Groups [56].

4.10. Nephroprotective Profile

On Oral Administration Of Aqueous Extract Of *U. Picta* Whole Plant Showed Clear Renal Protective Profile Due To The Presence Of Polyphenolic And Carbohydrate Compounds Against Acetaminophen Induced Nephrotoxic Rats [57].

4.11. Wound Healing Profile

Topical Application Of Aqueous Extract Showed Noticeable Wound Healing Potential In Wistar Rat Models. Results Indicated A Fall In Wound Closure Time And Rise In Wound Contraction. Furthermore, A Significant Increase In Proline Content Was Also Observed. All The Studies Were Compared With Standard Povidine Iodine Ointment [58].

5. Clinical Aspects

Majority Of The Studies On *U. Picta* Have Been Conducted With Alcoholic Or Aqueous Extracts. Both Preclinical And Clinical Experiments Have Concluded About The Quicker Healing Of Fractures Due To Early Accumulation Of Phosphorus And More Deposition Of Calcium. It Is Also Stated That The Decoction Is Prescribed For Cough, Chills And Fevers. The Leaves Are Considered Antiseptic And Used In Gonorrhea. The Roots And Pods Are Employed To Treat Prolapse Of Anus In Infants; The Pods Are Also Employed For The Treatment Of Sore-Mouth In Children. It Is Used For The Treatment Of Urinary Diseases, Tumors, Edema, Burning Sensation And Difficulty In Breathing. Its Paste, Mixed With Water, Is Used As An Antidote For Snake Bite [59].

6. Conclusion

The Beneficial Effects Of This Plant In Relation To Its Effectiveness And Flexibility Are Such To Facilitate Advancement In Research Appears Vital. Bearing In Mind Different Solvents Like Methanol, Ethanol, Chloroform And Aqueous Extract, Methanol Extract Are More Effective For Numerous Activities Of *Urariapicta*. Thus, *Urariapicta* Is Quite Gifted As A Multipurpose Medicinal Means.

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Conflicts Of Interests All Authors Have None To Declare 8. References

[1]Ved Dk, Goraya Gs. Demand and Supply of Medicinal Plants In India. In: Bishen Singh Mahendra Pal Singh, Editors. Publishers And Distributors Of Scientific Books. India: Dehradun; 2008. Pp. 9, 12.

[2] Anonymous, The Ayurvedic Formulary Of India, 2nd Ed. 2003. Pp. 55

[3] Sastry K. Charaka Samhita Of Agnivesa With Cakrapanidattatika Part I & Ii. Varanasi: Chaukhambha Sanskrit Sansthan; 1997.

[4] Sarma Pv. Dhanvantari-Nighantuh. In: Sharma Gp, Editor. Varanasi: Choukhambhaorientalia; 2005. Pp. 32.

[5] Gurav Am, Dhanorkar Vm, Dhar Bp And Lavekar Gs. In Vitro Propagation Of The Medicinal Plant Uraria Picta (Jacq.) Desv. Ex Dc. From Cotyledonary Node And Nodal Explants. Phcog Mag 2008; 4(16): 0973-1296.

[6] Sagwan S, Rao Dv, Sharma Ra. Phytochemical Evaluation And Quantification Of Primary Metabolites Ofmaytenusemarginata(Willd.) Ding Hou. J Chem Pharm Res 2010; 2(6): 46-50.

[7] Rajurkar Ns, Gaikwad K. Evaluation Of Phytochemicals, Antioxidant Activity And Elemental Content Of Adiantum Capillus Venerisleaves. J Chem Pharm Res 2012; 4(1): 365-374.

[8] Groom, A., Uraria Picta. Iucn Red List Of Threatened Species. 2012: International Union For Conservation Of Nature And Natural Resources, Iucn.

[9] Hari O, Anjana S, Naseer M, Santosh K. Phytochemical Screening And Elemental Analysis In Different Plant Parts Of Urariapictadesv: A Dashmul Species.J Chem Pharm Res 2014; 6 (5); 756-760.

[10] Anonymous. The Ayurvedic Pharmacopoeia Of India Part- I Volume- Iv, 1st Ed. New Delhi: Government Of India, Ministry Of Health And Family Welfare, Department Of Ayush; 2004. P. 99-101, 235.

[11] Bhattacharjee A, Shashidhara Sc, Saha S, Phytochemical And Ethno-Pharmacological Profile Of Desmodium Gangeticum (L.) Dc. A Review. Int J Biomed Res 2013; 4(10): 507-515.

[12] Abdullah Al H, Choudhury Mh, Zafrulazam Atm. Antimicrobial, Cytotoxic And Antioxidant Profiles Of Desmodiumheterocarpon. Bang Pharm J 2011; 14(1): 49-52.

[13] Subha R, Madan Mp, Ajay Ksr. An Ethnomedicinal, Phytochemical And Pharmacological Profile Of Desmodium Gangeticum (L.) Dc. And Desmodium Adscendens (Sw.) Dc. J Ethnopharmacol 2011; 136: 283-96.

[14] Varaprasad Mv, Balakrishna K, Sukumar E, Patra A. Gangetial, A New Pterocarpan From The Roots Of Desmodium Gangeticum. J Indian Chem Soc 2009; 86: 654–56.

[15] Mishra Pk, Singh N, Ahmad G, Dube A, Maurya R. Glycolipids And Other Constituents From Desmodium Gangeticum With Antileishmanial And Immunomodulatory Profiles. Bioorg Med Chem Lett 2005; 15: 4543-46.

[16] Anurag S, Singh Pk. An Ethnobotanical Study Of Medicinal Plants In Chandauli District Of Uttar Pradesh, India. J Ethnopharmacol 2009; 121: 324–29.

[17] Robert Am. Encyclopedia Of Physical Science And Technology – Alkaloids, 3rded, 2002.

[18] Duthie Gg, Duthie Sj, Kyle Am. Plant Polyphenols In Cancer And Heart Disease: Implications As Nutritional Antioxidants.Nutr Res Rev 2000; 13: 79-106.

[19] Malairajan P, Gopalakrishnan G, Narasimhan S, Veni Kjk. Analgesic Activity Of Some Indian Medicinal Plants. J Ethnopharmacol 2006; 19: 425-428.

[20] Luo J, Cheung J, Yevich E. Novel Terpenoid-Type Quinines Isolated From Pycnanthusangolensis Of Potential Utility In The Treatment Of Type 2 Diabetes.J Pharmacolexptl Therapeutics1999; 288: 529-534.

[21] Sharma Hl, Sharma Kk, Editors. Drug Therapy Of Heart Failure. In: Principle Of Pharmacology, 1st Ed. Hyderabad: Paras Publishers; 2007. P. 314 - 325. [22] Okwu De. Evaluation Of The Chemical Composition Of Indigenous Spices And Flavouring Agents. Global J Pure Applsci2001; 8: 455-459.

[23] Hayashi T, Maruyama H, Hatton K, Hazeki O, Yamasaki K, Tanaka T. Ellagitannins From Lagerstroemia Speciosa As Activators Of Glucose Transport In Fat Cells.Planta Med2002; 68(2): 173-175.

[24] HosseinzadehH, Nassir Am. Review Of Pharmacological Effects Of Glycyrrhiza Sp. And Its Bioactive Compounds. Phytother Res 2008; 22: 709-24.

[25] Rupasinghe Hp, Jackson Cj, Poysa V, Berardo Cd, Bewley Jd; Jenkinson J. Soyasapogenol A And B Distribution In Soyabean (Glycine Max L. Merr) In Relation To Seed Physiology, Genetic Variability And Growing Location. Jagric Food Chem2003; 51: 5888-5894.

[26] Liu F, Kim J, Li Y, X Liu, Li J, Chen X. An Extract Of Lagerstroemia Speciosa L.Has Insulin-Like Glucose Uptake– Stimulatory And Adipocyte Differentiation–Inhibitory Activities In 3t3-L1 Cells. Jnutr2001; 131(9): 2242- 2247.

[27] Xueqin M, Chengjian Z, Changling H, Khalid R, Luping Q. The Genus Desmodium (Fabaceae)-Traditional Uses In Chinese Medicine, Phytochemistry And Pharmacology. J Ethnopharmacol 2011; 138: 314-32.

[28] Tabuti Jrs, Lye Ka, Dhillion Ss. Traditional Herbal Drugs Of Bulamogi, Uganda: Plants, Use And Administration. J Ethnopharmacol 2003; 88: 19–44.

[29] Kosalge Sb. Fursule Ra. Investigation Of Ethnomedicinal Claims Of Some Plants Used By Tribals Of Satpuda Hills In India. J Ethnopharmacol 2009; 121: 456-61. [30] Abinash Ps, Venkat Kr, Pragya S, Pranab G, Utpal B. Ethnobotany Of Medicinal Plants Used By Assamese People For Various Skin Ailments And Cosmetics. J Ethnopharmacol 2006; 106: 149-57.

[31] Dilip Kek, Janardhan Gr. Ethno Botanical Polypharmacy Of Traditional Healers In Wayanad (Kerala) To Treat Type 2 Diabetes. Ind J Trad Knowl 2012; 11(4): 667-73.

[32] Shubhangi P. Indigenous Herbal Remedies Against Stomach Disorder From Jalgaon District (M.S.) India. Lifescileafl2012; 5: 66-70.

[33] Shanmugam S, Rajendran K, Suresh K. Traditional Uses Of Medicinal Plants Among The Rural People In Sivagangai District Of Tamil Nadu, Southern India. Asian Pac J Trop Biomed 2012; S429-S434.

[34] Rao Br, Sunitha S. Medicinal Plant Resources Of Rudrakod Sacred Grove In Nallamalais, Andhra Pradesh, India. J Biodiversity 2011; 2(2): 75-89.

[35] Vijay Vw, Ashok Kj. Traditional Herbal Remedies Among Bheel And Bhilala Tribes Of Jhabua District Madhya Pradesh. Int J Bio Tech 2010; 1(2): 20-24.

[36] Cheryl L. Ethnomedicines Used In Trinidad And Tobago For Reproductive Problems. J Ethnobiol Ethnomed 2007; 3(13):1-12.

[37] Jeyaprakash K, Ayyanar M, Geetha Kn, Sekar T. Traditional Uses Of Medicnal Plants Among The Tribal People In Theni Diatrict (Westernghats), Sourthern India. Asian Pac J Trop Biomed 2011; S20-S25.

[38] Richa Sc. Taxa Of Family Fabaceae: A Potential Of Local Medicinal Values In Vindhya Region Uttar Pradesh, India. Int J Pharma And Bio Sci 2010; 1(4): B46-B53.

[39] Patel Bd, Kamariya Yh, Patel Mb. Antioxidant Potential Of Aqueous Extract Of Entire Plant Of Uraria Picta Desv. Int J Pharm Res 2011; 3(4), 92-96.

[40] Kurian Ga, Yagnesh N, Kishan Rs, Paddikkala J. Methanol Extract Of Desmodium Gangeticum Roots Preserves Mitochondrial Respiratory Enzymes, Protecting Rat Heart Against Oxidative Stress Induced By Reperfusion Injury. J Pharm Pharmacol 2008; 60(4): 523-30.

[41] Kurian G, Paddikkala J. Role Of Mitochondrial Enzymes And Sarcoplasmic Atpase In Cardioprotection Mediated By Aqueous Extract Of Desmodium Gangeticum (L.) Dc Root On Ischemic Reperfusion Injury. Ind J Pharm Sci 2010; 72(6):745-49.

[42] Rathi A, Rao Cv, Ravishankar B, Deb S, Mehrotra S. Anti-Inflammatory And Anti-Nociceptive Profile Of The Water Decoction Desmodiumgangeticum. J Ethnopharmacol 2004; 95: 259-63.

[43] Sharma K, Rani R, Dhalwal K, Shinde V, Mahadik K. Natural Compounds As Anti-Arthritic Agents- A Review. Pharmacog Rev 2009; 3(5): 22–8.

[44] Singh N, Mishra Pk, Kapil A, Arya Kr, Maurya R, Dube A. Efficacy Of Desmodiumgangeticumextract And Its Fractions Against Experimental Visceral Leishmaniasis. J Ethnopharmacol 2005; 98 (1-2): 83–8.

[45] Shabi Mm, Paddikkala J. Cardiotonic And Anti Ischemic Reperfusion Injury Effect Of Desmodiumgangeticumroot Methanol Extract. Turk J Biochem2010; 35 (2): 83–90.

[46] Kurian Ga, Srivats Rss, Gomathi R, Shabi Mm, Paddikkala J. Interpretation Of Inotropic Effect Exhibited By Desmo diumgangeticum chloroform Root Extract Through Gsms And Atomic Mass Spectroscopy: Evaluation Of Its Ant-Ischemia Reperfusion Property In Isolated Rat Heart. Asian J Biochem2010; 5(1): 23–32.

[47] Kurian Ga, Paddikkala J. Administration Of Aqueous Extract Of Desmodiumgangeticum(L) Root Protects Rat Heart Against Ischemic Reperfusion Injury Induced Oxidative Stress. Indian J Expbiol2009; 47(2): 129-135.

[48] Kurian Ga, Suryanarayanan S, Raman A, Padikkala J. Antioxidant Effects Of Ethyl Acetate Extract Of Desmodiumgangeticumroot On Myocardial Ischemia Reperfusion Injury In Rat Hearts. Chinese Med 2010; 5(1): 3-8.

[49] Gino Ak, Jose P. Methanol Extract Of Desmodiumgangeticumdc Root Mimetic Postconditioning Effect In Isolated Perfused Rat Heart By Stimulating Muscarinic Receptors. Asian Pac J Trop Med 2012; 5(6): 448-54.

[50] Dharmani P, Mishra Pk, Maurya R, Chauhan Vs, Palit G. Desmodiumgangeticum: A Potent Anti-Ulcer Agent. Indian J Expbiol2005; 43(6): 517-21.

[51] Ayyavu M, Robert J, Dowlathabad Mr, Devarajan T. Gastroprotective Effect Of Desmodiumgangeticumroots On Gastric Ulcer Mouse Models. Rev Bras Farmacogn2012; 22(5): 37-44.

[52] Jabbar S, Khan Mt, Choudhuri Ms. The Effects Of Aqueous Extracts Of Desmodiumgangeticumdc. (Leguminosae) On The Central Nervous System. Pharmazie 2001; 56(6): 506–8.

[53] JoshiH, ParleM. Antiamnesic Effects Of Desmodiumgangeticumin Mice. Yakugakuzasshi 2006; 126 (9):795-804.

[54] Hanumanthachar J, Milind P. Pharmacological Evidences For The Antiamnesic Effects Of Desmodiumgangeticumin Mice. Iran J Pharm Res 2010; 6(3):199–207.

[55] Govindarajan R, Asare-Anane H, Persaud S, Jones P, Houghton Pj. Effect Of Desmodiumgangeticumextract On Blood Glucose In Rats And On Insulin Secretion In Vitro. Planta Med 2007; 73(2): 427–32.

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[56] Prasad Mvv, Balakrishna K, Carey Mw. Hepatoprotective Profile Of Roots Of Desmodiumgangeticum(Linn.) Dc. Asian J Chem2005; 17(4): 2847-49.

[57] Kale Rh, Halde Uk, Biyani Kr. Protective Effect Of Aqueous Extract Of Urariapicta On Acetaminophen Induced Nephrotoxicity In Rats. Int J Res Pharm Biomed Sci 2012; 3(1), 110-113.

[58] Jain V, Prasad V, Pandey R. Wound Healing Profile Of Desmodiumgangeticumin Different Wound Models. J Plant Sci2006; 1(3): 247–53.

[59] Parotta Ja. Healing Plants Of Peninsular India, Usfda Forest Service, International Institute Of Tropical Forestry, Puerto, Usa. Uk: Cabi Publishing, 2001; Pp. 418-419, 393-394.