



A Review on Pharmacological Actions of Herbs Used In Siddha Formulations

G. Ramanan ^{*1}, L. Abinash manimaran ², S.Sonitha ³, S. Mathukumar ⁴

*^{*1&2} UG Scholar, Sri Sairam Siddha Medical College and Research Centre, Chennai – 600 044, Tamil Nadu, India*

³ Lecturer, Sri Sairam Siddha Medical College and Research Centre, Chennai – 600 044, Tamil Nadu, India

⁴ Professor, Sri Sairam Siddha Medical College and Research Centre, Chennai – 600 044, Tamil Nadu, India

ABSTRACT

Siddha system of medicine is a distinct therapeutic science with many single or compound formulations. In Siddha system medicine is classified into 32 internal and 32 external types of medicines. In that medicines were predominantly made up of herbs. These herbal and herbo-mineral medicines has a great therapeutic value against various ailments because of their naturally occurring active principles, Many pharmacological and phytochemical studies were proven that the following herbs Andrographis paniculata (Nilavembu), Cyperus rotundus (Koraikkizhangu), Sida acuta (vattatiruppi), Rothea serrata (siruthekkku), Anacyclus pyrethrum (akrakaram) has various chemical constituents such as alkaloids, flavonoids, amides, terpenoids, steroids etc., these biologically active substance cures various ailments such as fever (suram), leucoderma (ven-kustam), sinusitis (pinisam), Arthritis (keel vatham) when it was added in the siddha medicinal formulations in an appropriate ratio. In this article discussing about the pharmacologically active substance of siddha herbs and their bioactivity on the living organisms and herbal siddha preparation were also listed with a proper references.

KEY WORDS: *Siddha medicine, Siddha herbs, Pharmacological actions, Bioactivity*

Corresponding Author: G. Ramanan, Sri Sairam Siddha Medical College and Research Centre, Chennai – 600 044, Tamil Nadu, India

1. Introduction

Siddha system of medicine is the oldest traditional system of healing that originate in southern part of India. It is based on the combination of medicinal practice and spiritual disciplines. Siddha medicine were derived from the nature sources i.e either herbs or metalo-mineral forms, Primarily herbal preparation were predominantly used to heal the disease. The World Health Organization (WHO) has listed 21,000 plants, which are used for medicinal purposes around the world. Among these 2500 species are in India, out of which 150 species are used commercially on a fairly large scale. India is the largest producer of medicinal herbs and is called as botanical garden of the world [1].

The current review focuses on pharmacological actions of herbs used in siddha formulations. These herbs were used in the preparation of herbal and herbo-mineral medicines with great therapeutic value.

2. Materials and Methods

2.1. Study design

A literature review on pharmacological actions of herbs used in siddha formulations.

2.2. Data collection

The information were collected by referring the authorised articles published in well classed journal's and siddha literatures by using electronic search.

2.3. Words used for citation

Botanical names of the corresponding plants, distribution, pharmacological actions, constituent etc

2.4. Review of herbs

Medicinal plants are considered a repository of numerous types of bioactive compounds possessing varied therapeutic properties. According to the World Health Organisation survey report approximately 80% of the people in developing countries depend on traditional herbal related formulations to cure diseases [2]

Table 1: Herbs used in siddha formulations

S.no	Botanical name	Tamil name	Family	Part used
1.	<i>Andrographis paniculata</i>	Nilavembu	Acanthaceae	Whole plant
2.	<i>Cypreus rotundus</i>	Koraikkizhangu	Cyperaceae	Rhizome
3.	<i>Sida acuta</i>	Vattatiruppi	Malvaceae	Root
4.	<i>Rothea serrata</i>	Siruthekku	Verbenaceae	Root
5.	<i>Anacyclus pyrethrum</i>	Akarakaram	Asteraceae	Root

2.5. *Andrographis paniculata*

Kingdom : Plantae

Division : Tracheophyta

Class : Magnoliopsida

Order : Lamiales

Family : Acanthaceae

Genus : *Andrographis*

Species : *Andrographis paniculata*

A. Paniculata (Acanthaceae) is known as Nilavembu in siddha is a herb reaching upto a height of 30-110 cm with glabrous leaves and white flowers with purple spotted petals. It is seen as a common weed in South India and also present in states of Assam, Missoram and Himachal Pradesh [3]. It is used to cure malaria, leucoderma, jaundice, abscess, wounds and eczema [4]. Diterpenoids andrographolide are major bioactive components. The compounds from the plant have been reported to have anti-inflammatory [5], anti-cancer [6], antimicrobial [7] and hepatoprotective [8], anti-viral activities [9].

Major Siddha formulations using *A. paniculata* as an ingredient includes kabasura kudineer, Nilavembu kudineer, vathasura Kudineer [10].

Taste: Bitter [65]

Therapeutic uses: Arthritis, fever, sinusitis, syncope [65]

Table 2: Pharmacological actions of *Andrographis paniculata*

S.no	Constituents	Class	Bioactivity	Reference
1.	Bis-andrographolide	Terpene	Anti-HIV	11
2.	Ninandrographolide	Terpene	Immuno stimulant	12
3.	Oxygenated flavones	Flavonoid	Anti-bacterial	13
4.	OroxylinA	Flavonoid	Anti-cancer	14

5.	Andrographolide	Diterpene	Anti-inflammatory, anti-cancerous, antimicrobial and hepatoprotective	5,8
----	-----------------	-----------	---	-----

2.6. *Cyperus rotundus*

Kingdom : Plantae

Division : Tracheophyta

Class : Magnoliopsida

Order : Poales

Family : Cyperaceae

Genus : *Cyperus*

Species : *Cyperus rotundus*

C. rotundus (Cyperaceae) is a perennial sedge plant and is known as Koraikkizhangu in Siddha. This is seen distributed all over India [15]. The drug is used as anti microbial, cytotoxic, larvicidal [16], anti-inflammatory [17] and anti-malarial activities [18]. This drug also possess analgesic, antispasmodic, astringent, diaphoretic, diuretic properties and is used as tonic and vermifuge [15]. Major Siddha formulations using *C. rotundus* as an ingredient includes Athimathura mathirai, Adathodai chooranam, sivathai chooranam, sukkutailam, Sanjeevi theener, Chandraprakasa mathirai, kabasura kudineer, Thathu busti kuligai, Parangipattai chooranam, Milagu thailam [10].

Taste : Bitter [65]

Therapeutic uses : Hypertension, fever, thirst [65]

Table 3: Pharmacological actions of *Cyperus rotundus*

S.no	Constituents	Class	Bioactivity	Reference
1.	Vitexin	Flavonoid	Anti-viral, anti-cancerous	19,20
2.	Cyperene	Sesquiterpene	Apoptotic, anti-oxidant, anti-bacterial	21,22
3.	Kobusone	Sesquiterpene	Anti-inflammatory, analgesic	16
4.	Cyperenone	Sesquiterpene	Antiulcer	17
5.	Eugenol	Ether-alcohol	Antiseptic	18

2.7. *Sida acuta*

Kingdom : Plantae

Division : Tracheophyta

Class : Magnoliopsida

Order : Malvales

Family : Malvaceae

Genus : *Sida*

Species : *Sida acuta*

S. acuta (Malvaceae) and is known as Vattatiruppi in siddha. This is a common weed of waste plains and grows gregariously and is present in the tropical regions in India. The whole plant is effective in treating snake bites and haemorrhagic effects of *Bothrops atrox* venom [23], and is also used for the treatment of urinary infections [35]. The drug possess antimalarial activity [24], anti-ulcer [25], hepatoprotective activity [26], cardiovascular activity [27], anticancer [28] and anti-inflammatory activities [29].

The phytochemical present in this species are vasicine, ephedrine and cryptolepine, ecdysterone, β -sistosterol, stigmaterol, campesterol, tannins, phenolic compounds, evofolin-A and B, scopoletin, loliolid and 4-ketopinonesinol, polyphenol, sesquiterpene and flavonoids [30-33]. Tannin obtained from this plant is used to cure ailments like leucorrhoea, rhinorrhea and diarrhea [34].

Major Siddha formulations using *S. acuta* as an ingredient is Kabasura Kudineer [10].

Taste : Astringent [65]

Therapeutic uses : Arthritis, diarrhea, fever, itching, scabies [65]

Table 4: Pharmacological actions of *Sida acuta*

S.no	Constituents	Class	Bioactivity	Reference
1.	Vasicine	Alkaloid	Antibacterial	35
2.	Sistosterol	Steroid	Cytotoxic, anti-microbial	36
3.	Evofolin	Phenyl propene	Anti-microbial	37
4.	4-ketopinonesinol	Lignan	Anti-oxidant	38
5.	Loliodid	Monoterpenoid hydroxylactone	Oxidative stress protection, anti melanogenic	39

2.8. *Rotheca serrata*

Kingdom : Plantae
 Division : Tracheophyta
 Class : Magnoliopsida
 Order : Lamiales
 Family : Lamiaceae
 Genus : Rotheca
 Species : Rotheca serrata

R. serrata (syn. *Clerodendrum serratum*), (Verbenaceae) is a shrub and is known as Ciruteku in Siddha. It is found upto an altitude of 1200m in lower Himalayas distributed in Kumaun, West Bengal and Bihar [40]. The major chemical constituents present are D-Mannitol, gamma-sitosterol, stigmasterol, glucose, oleanolic, queretaroic and serratagenic acid [41-45]. It shows antiasthmatic, antispasmodic [46], antiinflammatory and antipyretic activities [47].

Major Siddha preparations using *R. serrata* as an ingredient includes Sarabungavilvathi ilakam, Rasakanthi melugu, kabasura Kudineer , Notchi thailam, Parankipattai rasayanam, vadha sura Kudineer [10].

Taste : slightly bitter [65]

Therapeutic uses : Asthma, fever, myalgia, sinusitis [65]

Table 5: Pharmacological actions of *Rotheca serrata*

S.no	Constituents	Class	Bioactivity	Reference
1.	D-Mannitol	Alcohol	Diuretic	48
2.	Serratagenic acid	Triterp enoid	Antibacterial	49
3.	Stigmasterol	Steroid alcohol	Anti-inflammatory	50
4.	Oleanolic acid	Triterp enoid	Anti-tumour, anti-fungal, anti-inflammatory	51
5.	Verbacoside	Sterol	Anti proliferative	52

2.9. *Anacyclus pyrethrum*

Kingdom : Plantae
 Division : Tracheophyta
 Class : Magnoliopsida
 Order : Asterales
 Family : Asteraceae
 Genus : Anacyclus
 Species : Anacyclus pyrethrum

A. Pyrethrum (Asteraceae) is known as Akarakaram in siddha. It is perennial procumbent

herb native to North Africa and is cultivated at the elevation of 900m in Jammu and Kashmir [53]. The phyto constituents present in this drug belongs to the class of amides, isoflavones and alkaloids [54-57]. The drug possesses immunomodulatory [58], anticonvulsant [59], antidepressant [60], antidiabetic [61] and antibacterial activities [62].

Major Siddha formulations using *A. pyrethrum* as an ingredient includes Sarabungavilvathi ilakam, kabasura Kudineer, korosanai mathirai, nanthi mezhugu, thuthulai nei , vasanthakusmagaram mathirai [10].

Taste : sweet [65]

Therapeutic uses : Arthritis, dental problem, dryness of tongue, epilepsy, fever, tonsillitis [65]

Table 6: Pharmacological actions of *Anacyclus pyrethrum*

S.no	Constituents	Class	Bioactivity	Reference
1.	Pellitorine	Alkaloid	Anti-diabetic, anti-cancerous, anti-bacteria, anti-inflammatory	61
2.	Anacycline	Amide	Anti-inflammatory	56
3.	Sesamin	Amide	Anti-cancerous	56
4.	Genistein	Isoflavone	Anti-cancerous	63
5.	Biochanin	Isoflavone	Anti-inflammatory, neuroprotective	64

3. Conclusion

According to the findings, herbs taken to the study has a diverse pharmacologically active substances and those components has a marked value of bioactivity in human being. Either the herb used singly or in a formulation has a wide range of therapeutic value in curing diseases

5. References

- Seth S.D., Sharma B. Medicinal plants of India. Indian J. Med. Res. 2004;120:9–11.
- World Health Organisation. Health of indigenous peoples. Geneva, Switzerland: Factsheets N0 326; 2007.
- Samy RP, Thwin MM, Gopalakrishnakone P. Phytochemistry, pharmacology and clinical use of *Andrographis paniculata*. Natural Product Communications. 2007;2(5):1934578X0700200519.
- Panossian A, Davtyan T, Gukassyan N, et al. Effect of andrographolide and Kan Jang—fixed combination of extract SHA-10 and extract

- SHE-3—on proliferation of human lymphocytes, production of cytokines and immune activation markers in the whole blood cells culture. *Phytomedicine*. 2002;9(7):598-605. Doi:10.1078/094471102321616409
5. Kumar S, Patil HS, Sharma P, Kumar D, Dasari S, Puranik VG, et al. Andrographolide inhibits osteopontin expression and breast tumor growth through down regulation of PI3 kinase/Akt signaling pathway. *Curr Mol Med*. 2012;12(8):952-66. Doi: 10.2174/156652412802480826. PMID: 22804248.
 6. Levita J, Nawawi A, Mutalib A, Ibrahim S. Andrographolide: A review of its anti-inflammatory activity via inhibition of NF-kappaB activation from computational chemistry aspects. *Int J Pharmacol*. 2010;6(5):569-76.
 7. Shen KK, Liu TY, Xu C, Ji LL, Wang ZT. Andrographolide inhibits hepatoma cells growth and affects the expression of cell cycle related proteins. *Yao Xue Xue Bao*. 2009;44(9):973-9. PMID: 20055171.
 8. Singha PK, Roy S, Dey S. Antimicrobial activity of *Andrographis paniculata*. *Fitoterapia*. 2003;74(7-8):692-4. Doi:10.1016/s0367-326x(03)00159-x.
 9. Panraksa P, Ramphan S, Khongwichit S, Smith DR. Activity of andrographolide against dengue virus. *Antiviral Res*. 2017;139:69-78. Doi:10.1016/j.antiviral.2016.12.014
 10. Siddha formulary of India Part II, The controller of Publications, Delhi. 2011
 11. Reddy VL, Reddy SM, Ravikanth V, Krishnaiah P, Goud TV, Rao TP, et al. A new bis-andrographolide ether from *Andrographis paniculata* nees and evaluation of anti-HIV activity. *Nat Prod Res*. 2005;19(3):223-30. Doi: 10.1080/14786410410001709197. PMID: 15702635.
 12. Puri A, Saxena R, Saxena RP, Saxena KC, Srivastava V, Tandon JS. Immunostimulant agents from *Andrographis paniculata*. *J Nat Prod*. 1993;56(7):995-9. Doi:10.1021/np50097a002
 13. Xie Y, Yang W, Tang F, Chen X, Ren L. Antibacterial activities of flavonoids: Structure-activity relationship and mechanism. *Curr Med Chem*. 2015;22(1):132-49. Doi:10.2174/0929867321666140916113443
 14. Li HN, Nie FF, Liu W, Dai QS, Lu N, Qi Q, et al. Apoptosis induction of oroxylin a in human cervical cancer HeLa cell line in vitro and in vivo. *Toxicology*. 2009;4,257(1-2):80-5. Doi: 10.1016/j.tox.2008.12.011. Epub. 2008 Dec 24. PMID: 19135124.
 15. Samraj K, Thillaivanan S, Kanagavalli K. An update on siddha herb Korai (*Cyperus rotundus*, L.): A review. *IJP*. 2014;1(4):233-42.
 16. Al-Massarani S, Al-Enzi F, Al-Tamimi M, Al-Jomaiah N, Al-Amri R, Roaa BK, et al. Composition and biological activity of *Cyperus rotundus* L. tuber volatiles from Saudi Arabia. *Natural Volatiles and Essential Oils*. 2016;3(2):26-34
 17. Subramoniam A. Quality Standards of Indian Medicinal Plants. ICMR, NewDelhi. 2003;1:89-94.
 18. Gupta MB, Palit TK, Singh N, Bhargava KP. Pharmacological studies to isolate the active constituents from *Cyperus rotundus* possessing anti-inflammatory, anti-pyretic and analgesic activities. *Indian J Med Res*. 1971;59(1):76-82
 19. Thebtaranonth C, Thebtaranonth Y, Wanauppathamkul S, Yuthavong Y. antimalarial sesquiterpenes from tubers of *Cyperus rotundus*: structure of 10,12-peroxy-calamenene, a sesquiterpene endoperoxide. *Phytochemistry*. 1995;40(1):125-8. Doi:10.1016/0031-9422(95)00260-e
 20. National Center for Biotechnology Information (2022). PubChem Bioassay Record for Bioactivity AID 366284 – SID 103582302, Bioactivity for AID 366284 – SID 103582302, Source: ChEMBL. Retrieved July 6, 2022 from <https://pubchem.ncbi.nlm.nih.gov/bioassay/366284#sid=103582302>.

21. Al-Snafi AE. A review on *Cyperus rotundus* a potential medicinal plant. IOSR J Phar. 2016;6(7):32-48.
22. Ahn JH, Lee TW, Kim KH, Byun H, Ryu B, Lee KT, et al. 6-Acetoxy Cyperene, a Patchoulane-type Sesquiterpene Isolated from *Cyperus rotundus* Rhizomes ,Induces Caspase-dependent Apoptosis in Human Ovarian Cancer Cells. *Phytother Res.* 2015;29(9):1330-8. Doi: 10.1002/ptr.5385. Epub 2015 Jun 10.
23. Otero R, Núñez V, Barona J, Fonnegra R, Jiménez SL, Osorio RG, et al. snakebites and ethnobotany in the northwest region of Colombia. Part III: Neutralization of the haemorrhagic effect of *Bothrops atrox* venom. *J Ethnopharmacol.* 2000;73(1-2):233-41. Doi: 10.1016/s0378-8741(00)00321-4.
24. Karou D, Savadogo A, Canini A, Yameogo S, Montesano C, Simpore J, et al. Antibacterial activity of alkaloids from *Sida acuta*. *Afr J Biotechnol.* 2006;5(2):195-200.
25. Malairajan P, Gopalakrishnan G, Narasimhan S, Veni K. Antiulcer Activity of *Sida acuta* Burm. *Nat Prod Sci.* 2006;12(3):150-2.
26. Sreedevi CD, Latha PG, Ancy P, Suja SR, Shyamal S, Shine VJ, et al. Hepatoprotective studies on *Sida acuta* Burm. F. *J Ethnopharmacol.* 2009;124(2):171-5. Doi:10.1016/j.jep.2009.04.055
27. Kannan RR, Vincent SG. *Cynodon dactylon* and *Sida acuta* extracts impact on the function of the cardiovascular system in zebrafish embryos. *J Biomed Res.* 2012;26(2):90-7. Doi:10.1016/S1674-8301(12)60017-7
28. Mallikarjuna G. Anticancer activity of *Sida acuta* Burm. F against Nitrosodiethylamine and CCl₄ induced hepatocellular carcinoma. *Indo American J of Pharm Research.* 2013;3(9):74-8.
29. Oboh IE, Onwukaeme DN. Analgesic, anti-inflammatory and anti-ulcer activities of *Sida acuta* in mice and rat. *Nigerian Journal of Natural Products and Medicine.* 2005;9(1):19-21.
30. Pandit SS, Naik SD, Jathar VS, Kulkarni AB. Insect molting hormone, ecdysterone from *Sida caprinifolia* Linn. *Indian J Chem.* 1976;14B:907-8.
31. Benjumea DM, Gómez-Betancur IC, Vásquez J, Alzate F, García-Silva A, Fontenla JA. Neuropharmacological effects of the ethanolic extract of *Sida acuta*. *Revista Brasileira de Farmacognosia.* 2016;26(2):209-15.
32. Prakash A, Varma RK, Ghosal S. Alkaloid constituents of *Sida acuta*, *S. Humilis*, *S. rhombifolia* and *S. spinosa*. *Planta Med.* 1981;43(4):384-8. Doi:10.1055/s-2007-971529
33. Konaté K, Souza A, Coulibaly AY, Meda NT, Kiendrebeogo M, Lamien-Meda a et al. In vitro antioxidant, lipoxygenase and xanthine oxidase inhibitory activities of fractions from *Cienfuegosia digitata* Cav., *Sida alba* L. and *Sida acuta* Burn f. (Malvaceae). *Pak J Biol Sci.* 2010;13(22):1092-8. Doi: 10.3923/pjbs.2010.1092.1098. PMID: 21313883.
34. Blytt HJ, Guscar TK, Butler LG. Antinutritional effects and ecological significance of dietary condensed tannins may not be due to binding and inhibiting digestive enzymes. *J Chem Ecol.* 1988;14(6):1455-65. Doi:10.1007/BF01012417
35. The Ayurvedic pharmacopoeia of India, part 1, New Delhi: Government of India, Ministry of Health and family Welfare, Department of indian System of Medicine and Homeopathy. 2001;3:110-11.
36. Donia AE, Soliman GA, El-Sakhawy MA, Yusufoglu H, Zaghoul AM. Cytotoxic and antimicrobial activities of *Emex spinosa* (L.) Campd. Extract. *Pak J Pharm sci.* 2014;27(2):351-6.
37. Brader G, Bacher M, Hofer O, Greger H. Prenylated phenylpropenes from *Coleonema pulchellum* with antimicrobial activity. *Phytochemistry.* 1997;45(6):1207-12. [https://doi.org/10.1016/s0031-9422\(97\)00124-6](https://doi.org/10.1016/s0031-9422(97)00124-6)
38. Tebboub O, Cotugno R, Oke-Altuntas F, et al. Antioxidant Potential of herbal Preparations and Components from *Galactites elegans* (All.) nyman ex Soldano. *Evid Based*

- Complement Alternat Med. 2018;9294358. Doi:10.1155/2018/9294358
39. Park SH, Choi E, Kim S, Kim DS, Kim JH, Chang S, et al. Oxidative Stress-protective and Anti-Melanogenic Effects of Loliolide and Ethanol Extract from fresh Water Green Algae, *Prasiola japonica*. Int J Mol Sci. 2018;19(9):2825. <https://doi.org/10.3390/ijms19092825>
40. Quality Standards of Indian Medicinal Plants. ICMR, New Delhi. 2003;3:167-76.
41. Sachadev KS, Banerjee SK, Chakravarthi RN. Chemical examination of the root bark of *Clerodendrum serratum*. Bull Calcutta Sch Trop Med. 1965;17-8.
42. Verma SCL, Garg VP, Gupta SS. A source of mannitol. Crr Sci. 1967;36:126-7.
43. Vasavada SA, Banerjee SK, Chakravarti RN. Gamma—Sitosterol from *Clerodendron serratum*. Bull Calcutta Sch Trop Med. 1967;15(2):61.
44. Rangaswami S, Sarangan S. Sapogenins of *Clerodendrum serratum*. constitution of pentacyclic triterpene acid- serratagenic acid. Tetrahedron. 1969;25(17):3701-5.
45. Ramachandran Nair AG, Vedantham, TNC, Sankara SS. Crystalline components of *Clerodendrum serratum*. Curr Sci. 1967;45:391.
46. Sachadev KS, Vasavada SA, Joseph AD. Antihistamic activity of *Clerodendrum serratum* (Linn) Moon. Indian J Pharm. 1964;26:105-6.
47. Wang J, Ren H, Xu QL, et al. Antibacterial oleanane-type triterpenoids from pericarps of *Akebia trifoliata*. Food Chem. 2015;168:623-9. Doi:10.1016/j.foodchem.2014.07.105
48. Sachadev KS, Vasavada SA, Joseph AD. Antihistamic activity of *Clerodendrum serratum* (Linn) Moon. Indian J Pharm. 1964;26:105-6.
49. Wang J, Ren H, Xu QL, et al. Antibacterial oleanane-type triterpenoids from pericarps of *Akebia trifoliata*. Food Chem. 2015;168:623-9. Doi:10.1016/j.foodchem.2014.07.105
50. Antwi AO, Obiri DD, Osafo N, Essel LB, Forkuo AD, Atobiga C. Stigmasterol alleviates Cutaneous Allergic Responses in Rodents. Biomed Res Int. 2018;3984068. Published 2018 Jul 24. Doi:10.1155/2018/3984068
51. Fujii Y, Hirose S, Fujii T, Matsumoto N, Agematu H, Arisawa A. Hydroxylation of oleanolic acid to quercetaroic acid by cytochrome P450 from *Nonomuraea recticatena*. Bioscience, Biotechnology and Biochemistry. 2006;70(9):2299-302.
52. Ram A, Joseph DA, Balachandar S, Singh VP. Medicinal plants from Siddha system of medicine useful for treating respiratory diseases. Int J Adv Pharm anal. 2009 Jan 1;1(2):20.
53. Usmani A, Khushtar M, Arif M, Siddiqui MA, Sing SP, Mujahid M. pharmacognostic and phytopharmacology study of *Anacyclus pyrethrum*: An insight. J Appl Pharm. 2016;6(03):144-50.
54. Gulland JM, Hopton GU. II-Pellitorine, the pungent principle of *Anacyclus pyrethrum*. J Chem Soc. 1930;6-11.
55. Burden RS, Crombie L. Amides of vegetable origin. Part XII. A new series of alka-2,4-dienoic tyramine-amides from *Anacyclus pyrethrum* DC.(Compositae). J Chem Soc. 1969;(19):2477-81.
56. Boonen J, Sharma V, Dixit VK, Burvenich C, DeSpiegeleer B. LC-MS N-alkylamide profiling of an ethanolic *Anacyclus pyrethrum* root extract. planta Med. 2012;78(16):1787-95. Doi:10.1055/s-0032-1315371
57. Kaur H. Estrogenic activity of some herbal galactagogue constituents. Indian J Anim Nutr. 1998;15(3):232-4.
58. Boonen J, Sharma V, Dixit VK, Burvenich C, DeSpiegeleer B. LC-MS N-alkylamide profiling of an ethanolic *Anacyclus pyrethrum* root extract. Planta Med. 2012;78(16):1787-95. Doi:10.1055/s-0032-1315371
59. Gautam OP, Verna S, Jain SK. Anticonvulsant and myorelaxation activity of *Anacyclus pyrethrum* DC. (Akarkara) root extract. Pharmacology Online. 2011;1(1):121-5.
60. Badhe SR, Badhe RV, Ghaisas MM, Chopade VV, Deshpande AD. Evaluation of antidepressant activity of *Anacyclus pyrethrum* root extract. Int J Green Pharm. 2010;4(2):79-82.

61. Tyagi S, Mansoori MH, Singh NK, Shivhare MK, Bhardwaj P, Singh RK. antidiabetic effect of *Anacyclus pyrethrum* DC in alloxan induced diabetic rats. *Eur J Biol Sci.* 2011;3(4):117-20.
62. Doudach L, Meddah B, Almnamer R, Chibani F, Cherrah Y. In vitro antibacterial activity of the methanolic and aqueous extracts of *Anacyclus pyrethrum* used in Moroccan traditional medicine. *Int J Pharmaceut Sci.* 2012;4(3):402-5.
63. Gulland JM, Hopton GU. II-Pellitorine, the pungent principle of *Anacyclus pyrethrum*. *J Chem Soc.* 1930;6-11.
64. Yu C, Zhang P, Lou L, Wang Y. Perspectives Regarding the Role of Biochanin A in humans. *Front Pharmacol.* 2019;10:793. Published 2019 Jul 12. doi:10.3389/fphar.2019.00793
65. Murugesamudaliyar KS. *Siddha Materia Medica* (Medicinal Plants Division).8th edition, Chennai: Directorate of Indian Medicine and Homeopathy. 2006;7-713.