International Journal of Reverse Pharmacology and Health Research (IJRPHR)

Review article



Review on uttāmaņi karukku a pediatric traditional medicine

Easwari D^{1*}, Sociya Parvin M¹, Soundararajan DKS²

1* PG Scholars, Department of Kuzhandhai Maruthuvam, Government Siddha Medical College and Hospital, Palayamkottai.

²Head, Department of Kuzhandhai Maruthuvam, Government Siddha Medical College and Hospital, Palayamkottai.

ABSTRACT

According to World Health Organization (WHO) more than 80% of the world's population relies on traditional practitioners and their armamentarium of medicinal plants in order to meet health care needs in spite of modern medicine which is well established worldwide. In *cittā* traditional system foremost awareness has been given to Pediatric population which is considered as the backbone of the developing nation. There are various medicinal formulation and lifestyle practices which are till date in practice. One among the pediatric disease *kaṇa māntam* is treated with a sastric formulation *uttāmaṇi karukku*. In spite of various medicines used in other system of medicine, till date in Tamil Nadu the practice of *Vacampu (piḷḷai maruntu)* has its own role, the raw drug has its own positive and negative effects it is overcome with its purification method.

Keywords:

cittā, ka \square *a māntam, uttāma* \square *i karukku*, Pediatric.

Address for correspondence:

Easwari D

PG Scholar,

Department of Kuzhandhai Maruthuvam

CODEN: IJRPHR

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-Non Commercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work noncommercially, as long as appropriate credit s given and the new creations are licensed under the identical terms.

 $\textbf{For reprints contact:} \ publisher@ijrphr.com$

To access this article online

Website: http://www.ijrphr.com/

DOI: 10.121/ijrphr/02.0203.350

Quick response code



How to cite this article:

Easwari D, Sociya Parvin M, Soundararajan DKS, Review on uttāmaṇi karukku a pediatric traditional medicine, International Journal of Reverse Pharmacology and Health Research, 2019, 2(3), 20-25.

Received: April, 2019.

Accepted: June, 2019.

INTRODUCTION

The ancient *cittā* system, a tradition presumed to be lost is now finding its way back into the hearts and lives of people worldwide. Ancient wisdom in its nativity remains conserved by the teaching imparted by sages in primordial times, even in our postmodern world. ²Cittā, as a common noun known as "realized, perfect one", a term generally applied to a practitioner who has through his practice realized his dual goal of superhuman powers and bodily immortality.

According to *cittā*, 96 *tattuvam* influence the activities of the individual from birth to death. This system also believes body- mind- spirit is inseparable is always taken into consideration as a whole as their interconnectedness is a preestablished cosmic design of existence.

As the population increasing day by day there is an exponential increase and emergence of new diseases. The environment is getting more and more polluted, the humanity has almost come to a stage where the various available preventive measures are to be valued much well than the existing curative measures. Apart from Prevention and cure which exist in all systems, *cittā* system in addition also believes strongly in the transcendental (immortality).

Cittā medicine deals formally and systematically with each medical subject and is usually describes in a poetic verse format so that the knowledge could be passed down orally over generations from cittā physicians to their disciples³. Similar to other Indian system of medicine cittā also follows mukkuṛram theory and states that psycho-biological aspect is governs by 3 mukkuṛram vaļi, ālal and aiyam any change in body is due to the imbalance of these mukkurram⁴.

Anupava vaittiya kaļanciyam a classical cittā text classifies the diseases of childhood into 108 diseases which further classified into various sub-types based on the change in nature of disease presentation. One among the disease is māntam which is a very common entity in the pediatric population; kaṇa māntam is a classification of the disease māntam for which the sastric formulation uttāmaṇi karukku has been given.

⁵Symptoms of *kaṇa māntam* are:

"Aruntu meyyilait tirumalu maṭikkaṭic curamun Tiruntu mūkkilnīr vaṭitalum vayiruppum ciranōy Purinti raiccalum palavitam vayirupōk kuṭaṇē Varuntu kaṇṇiṇil mayakkamu māṅkaṇa māntam"

- Kulantai maruttuvam (verse 91)

- Weight loss
- Cough
- Fever
- Watery Nasal discharge
- Headache
- Diarrhea
- Bloating abdomen
- Drowsiness

Cittā literature describe a group of herbals suitable for pediatric population based on the pediatric organ developments, keeping pharmacokinetic and pharmacodynamics in mind, which indicates the in depth knowledge of cittā medicinal system in pediatric diseases. Drugs used in uttāmaņi karukku are briefly described.

The taxonomical classification, chemical constituents, *Cuvai* parameters, method of preparation of the drug, their activities has been described briefly in the upcoming topics.

Uttāmaņi karukku: (Internal medicine)

Karukku- to grind, to burn, scorch, tan, darkens by heat ⁷

Ingredients:

VacampuAcorus calamus L.10 gramUttāmaṇiPergularia daemia (Forssk.)Chiov.100 mluppuSodium chloride1 gram

Purification of *vacampu*:

The most important part of the drug preparation which makes the toxic drug into non toxic, it is one of the real identity of cittā system.

Vacampu cāmpal kuṇam:

"tērumē vacampānatil cāmpaltān

Nīrumēcalamtān atiliţa unnil

Vīrukoņţu veļivarum pētiyum

mārumenru vakuttanar nantiyē"

- *Citamparatāņu piḷḷai poruṭpaṇpu nūl part -1* Method of preparation:

⁶Vacampu has been calcinated by traditional cow dung method, *Uttāmaṇi* leaves are collected and juice is extracted, which is mixed with salt(Sodium chloride) and calcinated *vacampu* and the mixture is heated until the juice is fully absorbed and made powdery, at last again it is triturated and given for administration.

Dose: 13 to 200 mg Adjuvant: Hot water

Figure 1- Uttāmaņi karukku preparation.



Figure

tāmaṇi karukku



PROPERTIES OF THE DRUG:

Uttāmaņi karukku: Uttāmaņi:

A slender, hispid, fetid- smelling perennial climber. Leaves opposite, membranous, 3-9 cm long and wide, broadly ovate, orbicular or deeply cordate.

Figure 3- Pergularia daemia



Taxonomy

Kingdom : Plantae

: Tracheobionta Subkingdom Super division : Spermatophyta Division : Magnoliophyta Class : Magnoliopsida Subclass : Asteridae Order : Gentianales Family : Asclepiadaceae Genus : Pergularia

Species : P. daemia (Forsk) Chiv

Other names : vēliparutti, uttamamākāņi, ut-

tamakannikai

2- Ut-

Parts used : Leaves, roots and root bark

Parts used in the Trial drug : Leaves

 Cuvai
 : kaippu

 Tanmai
 : veppam

 Pirivu
 : kārppu

Action: Anti pyretic, Antispasmodic, Nervine tonic, Sedative Stomachic, Anthelmintic, Laxative, Expectorant (N. Balakrishnan et al 2009), Emetic, Anti inflammatory.

Active constituents :

Lupeol

Beta sitosterol

Quercetin

Kaemferol

Alpha -Amyrin

Beta- Amyrin

Betaine

Isohamnetin

Uttāmaņi potukuņam:

"Icikkum valiyiraippum ettaṭippum ēkum⁶

Pacikkumati māntamum pōm pār"

- Akattiyar kuṇavākaṭam.

The verse denotes *uttāmaṇi* helps to relieve from *icivu* (vali), *iraippu* and *ati māntam*. Appetite will be increased.

Medicinal properties:

Gastro Intestinal system:

- 1. Digestive disturbance, Flatulence
- 2. Loss of appetite
- 3. Gastric ulcer
- 4. Choleric diarrhea in children
- 5. Treat infantile diarrhea and Malarial intermittent fevers.

Respiratory system:

Cough, Fever, Cold, Asthma, Whooping cough and Capillary bronchitis.

-Nadkarni's Indian material medica vol-I

Table 1. Acute toxicity of P. daemia

Treatment	Dose oral (g/kg BW)a	Quantal symp- toms	Quantal mortality
Control (vehicle)	10 mg/kg	0/10	0/10
Test extract	0.5	0/10	0/10
	1.0	0/10	0/10
	1.5	0/10	0/10
	2.0	0/10	0/10
	2.5	3/10	2/10
	3.0	2/0	0/10

A g/kg body weight/os. (S. C. Jain et al, 1998)

From Table 1, it is evident that the plant extract was well tolerated orally in mice up to a dose of 2.0 g/kg BW with no mortality or serious side effects⁹.

 $^{11} The$ methanolic extract produced zone of inhibition against S. aureus only which showed 19 mm in $75\mu L,\,20$ mm in $100\mu L$ and 21 mm in $125\mu L$ concentration. In chloroform extract active against K. pneumoniae and S. aureus which produced 17 and 12 mm in $75\mu L,\,18$ and 13 mm in $100\mu L$ and 19 and 15 mm in $125\mu L$ concentrations. In aqueous extract was the most effective against all three test pathogens, but the maximum zone of inhibition was shown at $125\mu L$ concentration which produced 15mm against E. coli and K. pneumoniae, 22 mm zone of inhibition against S. aureus.

 11 A. niger and Penicillium sp. were highly susceptible to chloroform extract (22 and 19 mm) followed by aqueous extract (17 and 20 mm) and methanolic extract (16 and 16mm) at 125μ L concentration respectively.

Acute toxicity study:

¹⁰The ethanolic and aqueous extracts did not cause any mortality up to 2000 mg/kg and were considered as safe as per OECD guidelines^{15.}

Figure 4- Acorus calamus



Table 02. Antibacterial activity of methanol, chloroform and aqueous extract of Pergularia daemia (Forsk.) Chiov. (Ramanathan R et al, 2013)

S.No	Test bacteria	Zone of inhibition (Diameter in mm)								
		Methanol extract			Chloroform extract			Aqueous extract		
		75μL	100μL	125μL	75 μL	100μL	125μL	75μL	100μL	125µL
1	Escherichia coli	-	-	-	-	-	-	-	-	15
2	Klebsiella pneumonia	-	-	-	17	18	19	11	13	15
3	Staphylococcus aureus	19	20	21	12	13	15	19	20	22

Table 02. Antifungal activity of methanol, chloroform and aqueous extract of Pergularia daemia (Forsk.) Chiov. (Ramanathan R et al, 2013)

S. No	Test fungi	Zone of inhibition (Diameter in mm)									
		Methanol extract			Chloroform extract			Aqueous extract			
		75 μL	100μL	125µL	75 μL	100μL	125µL	75 μL	100μL	125µL	
1	Aspergillus niger	14	15	16	19	20	22	15	16	17	
2	Penicillium sp.	13	15	16	17	18	19	18	19	20	

¹⁰Among the aqueous and ethanolic extracts tested, the ethanolic extract of the aerial parts of Pergularia daemia possess Hepatoprotective activity against CCl4 intoxication in rats.

A. calamus is a perennial plant with creeping and extensively branched, aromatic rhizome, cylindrical, up to 2.5 cm thick, purplish-brown to light brown externally and white internally. At the rhizome forming, perennial that can grow to 2 meters resembling an iris 12

TAXONOMY

Kingdom : Plantae

Division : Magnoliophyta
Class : Liliopsida
Order : Acorales
Family : Acoraceae
Genus : Acorus

Species : calamus/ A. aromatics / A.

calamus var. americanus

Other species : Acorus gramineus

Other Tamil names

ukkiram, vacam, vacai, vēṇi, cuṭuvāṇ, uraippāṇ, pēr collā maruntu, piḷḷai maruntu⁶, koṭi kevuri, vaṅkirācam, kimaṇattippi, viṣaram, ōmacam, tiripaṅkucāti, ācuvētayam, vacuntakirumiyari.

Parts used : Dried Rhizome
Parts used in the trial drug : Ash of the Rhizome.

Cuvai : kārppu, kaippu
tanmai : Veppam

Pirivu : kārppu

Action

Stomachic, Anti periodic, Stimulant, Disinfectant, Germicide, Carminative, Expectorant and Antispasmodic.

Constituents:

- Alpha asarone- 1.17%
- Beta- Asarone- 92.68%
- Delta- Asarone- 2.27%
- 1-8 cineole, Terpionolene, sugar 8.342%
- Nitrogen 2.029%, Protein 10.318%
- A volatile essential oil- acronic.
- A bitter principle- acoretin(choline calamine)
- The essential oil of Acorus calamus is yellowish brown and is found to be composed of as aryl aldehyde, esters of palmitic acids and a small quantity of phenol, eugenol, methyl eugenol, calameneol and calameone.

- Indian Materia Medica (P. No: 35)

Active constituents:

The rhizomes of A. calamus Linn. has mixed fatty acids, as indicated by gas chromatography of the corresponding methyl esters were myristic (1.3%), palmitic (18.2%), palmitoleic (16.4%), stearic (7.3%), oleic (29.1%), linoleic (24.5%) and arachidic (3.2%). The nature of the sugars was defined by paper chromatography and confirmed by direct comparison with authentic samples. Composition of the sugars, as indicated by densitometer, was maltose (0.2%), glucose (20.7%) and fructose (79.1%). The content of the oil in dried sweet flag rhizomes was 1.20+/-0.12%. Acorenone was dominant in the rhizomes (20.86%) followed by isocalamendiol (12.75%). Besides Monoterpene hydrocarbons, sequestrine ketones, (Trans- or Alpha) Asarone (2, 4, 5trimethoxy-1- propenylbenzene), and Beta-asarone (cis- isomer) and eugenol were also identified. Some other compound identified in A. calamus are (-)-4-Terpineol, 2-Allyl-5-ethoxy-4methoxyphenol, Epieudesmin, Lysidine, (-)-Spathulenol, Borneol, Furyl ethyl ketone, Nonanoic Acid, 2,2,5,5-Tetramethyl-3hexanol, Bornyl acetate, Galgravin, Retusin, (9E,12E,15E)-9,12,15- Octadecatrien-1-ol, Butyl Butanoate, Geranylacetate, Sakuranin, Acetic Acid, Camphor, Isoelemicin, á-Ursolic acid, Acetophenone, Dehydroabietic acid, Isoeugenol Methylether, Apigenin 4',7-dimethyl ether, Dehydrodiisoeugenol, Linalool, Elemicin, Linoleic acid¹³.

The biochemical constituent in Vacampu is asarone. This compound has analgesic, sedative and neuro depressive activity which can produce sedation and hence reduce the irritable cry among neonates. In neonates with diarrhea, the spasmolytic and antisecretory effect of the extract can reduce the frequency of loose stools¹⁴.

¹²Heavy Metals Analysis The content of heavy metals such as lead was present within the permissible limit, cadmium; mercury and arsenic were not found in the drug Acorus calamus (table.4).

Table. 4. Heavy Metals Analysis (Ushakanthan et al., 2017)

S. N o.	Name of the Element	Results	Permissible Limit	
1	Lead	0.0914 ppm	10 ppm (WHO)	
2	Cadmium	Not detect- ed	o.3 ppm (WHO)	
3	Arsenic	Not detect- ed	3 ppm (API)	The eth-

anolic extract of *Acorus calamus* significantly protects against liver injuries resulting in improved serum biochemical parameters such as SGOT, SGPT and SALP¹⁶.

Fatty acids in *Acorus calamus* have palmitic acid and its ester which possess significant antifungal and antibacterial activity¹⁷. Various essential oil components present in essential oils like linalool, 1,8-cineol, caryophyllene, α humulene, and asarone have been reported to possess antioxidant activity.

Uppu:

Sodium chloride

Other names: kariyuppu, $c\bar{o}rruppu$, kataluppu, $v\bar{t}tuppu$, ilavanam, $camuttira lavanam^{18}$.

Kariyuppin potukunam:

"Māntam porumalarum vāyuvumpōmtīpaṇamām tontitta aiyan toṭarumō- cantatamum akkiṇiyiṇ puṣṭi aṭaruṅ kariyuppāl cikkukiṇra nīriraṅkuñ ceppu"

Kariyuppāl māntam, vayirrup porumal, vāyu, kapam nīnkum. Nīraṭaippu tīrum. Paciyum camākkiniyum atikappaṭum.

CONCLUSION

Cittā medicine has its own unique identity in its formulation of drugs. Despite of, various climatic change and geographical nature changes, the composition of drug most of its activities are similar mainly uttāmaṇi karukku, the herbo mineral combination has its own composition which has Hepatoprotective and Anti oxidant activity. Uttāmaṇi karukku will be more prioritized by further scientific validation.

FINANCIAL SUPPORTS

Nil

CONFLICTS OF INTEREST

None declared.

REFERENCES

- 1. WHO, Expert Committee on diabetes mellitus, World Health Organization, 1980, Geneva, pp.12-15.
- 2. David Gordon white, The Alchemical Body: Siddha Traditions in Medieval India, Edition 2007.
- 3. Thambyayah M & Amuthan A. Infantile seborrheic dermatitis: A pediatric cittā medicine treatise. Clinics in Dermatology 2014, vol-33, 356-357.
- 4. Natarajan, K. Principles of diagnosis in Siddha, Chennai: Department of Indian Medicine and Homeopathy, Chennai, 2009: pp-106, 154-159.
- 5. Pon G: Balavagadam, Department of Indian Medicine and Homeopathy, Chennai, Edition 6th, 2016, pp-100, 160.
- 6. K.S.Murugesa Mudhaliar, Gunapadam mooligai vaguppu, edition 9, 2013, pg.no.135,787
- 7. Tamil Lexicon, Vol II, Part I, University of Madras, 1982, pg. 750
- 8. Sambasivam pillai, T.V. Tamil English dictionary IV part II. Chennai: Department of Indian Medicine and Homeopathy, 1998, 2nd Ed.
- 9. Jain, S. C., Jain, R., Mascolo, N., Capasso, F., Vijayvergia, R., Sharma, R. A., & Mittal, C. (1998), Ethno pharmacological evaluation of Pergularia daemia (Forsk.) Chiov. Phytotherapy Research, 12(5), 378–380.
- 10. Sureshkumar, S. V., & Mishra, S. H. (2006), Hepatoprotective effect of extracts from Pergularia daemia Forsk. Journal of Ethno pharmacology, 107(2), 164–168.
- 11. R Ramanathan, R Baby, Antimicrobial activity of Canthium parviflorum Lam. And Pergularia daemia (Forsk.) Chiov, International Journal of Comprehensive Pharmacy, 2013, 09 (4).
- 12. Ushakanthan et al. bio chemical and physiochemical evaluation of "vacampu choornam" (Acorus calamus rhizome powder), World Journal of Pharmaceutical Research, Volume 6, Issue 8, 2017, 1451-1459.
- 13. R. Balakumbahan, K. Rajamani and K. Kumanan, Acorus calamus: An overview (2010), Journal of Medicinal Plants Research Vol. 4(25), pp. 2740-2745
- 14. Tanigasalam, V., Vishnu Bhat, B. ., Adhisivam, B., Plakkal, N., & Harichandra Kumar, K. T. (2017). *Vacampu (Acorus calamus) Administration: A Harmful Infant Rearing Practice in South India. The Indian Journal of Pediatrics*, 84(10), 802–803.
- 15. OECD, 1996. OECD Guidelines for the Testing of Chemicals Test no. 423: Acute Oral Toxicity—Acute Toxic Class Method.
- 16. S. Palani, S. Raja, Therapeutic efficacy of antihepatotoxic and antioxidant activities of *Acorus calamus* on acetaminophen- induced toxicity in rat, International journal of integrative biology, 2009, Vol; 7, No.1.
- 17. Archana Parki, Pinky Chaubey, Seasonal variation in essential oil compositions and anti oxidant properties of *Acorus calamus* L. Accessions, Medicines open access journal, 2017, Vol 4, issue 4.
- 18. Dr. R. Thiyagarajan, Gunapadam thathu seeva vaguppu, Department of Indian Medicine and Homeopathy, Chennai, 8th edition, 2013, 384-386.
- 19. V.H. Bhaskar and N. Balakrishnan, Vēliparutti (Pergularia daemia (Forsk.) Chiov.) As a phytomedicine: A review, 2009, International Journal of PharmTech Research CODEN (USA): IJPRIF, Vol.1, No.4, pp 1305-1313