

# Evaluation of Synergistic Anthelmintic Activity of Polyherbal Formulation on *Pheritima Posthuma*

Abirami AV, Rajamanickam V, Venakateshan N, Sivakumar V

Department of Pharmaceutical Chemistry, Arulmigu Kalasalingam College of Pharmacy, Krishnankoil, India

**Abstract:** - The prevalence of worm infestation is high in underdeveloped and developing countries because of poor sanitation and lack of health education. Our study was aimed to search out the individual and synergistic result of *Ocimum sanctum* (Family: Lamiaceae), *Coriandrum sativum* (Family: Apiaceae), *rosid dicot genus pinnata* (Family: Fabaceae), *Epipremnum aureum* (Family: Araceae) and *curcuma longa* (Family: Lamiaceae) extract as a result of they need been used historically for the treatment of worm infestation. additional exploring the phytoconstituents gift within the herbs of the polyherbal formulation can aid us within the bioactivity gift.

Various concentration of ethanolic extracts of varied herbs and therefore the polyherbal formulation were subjected for assessment of anthelmintic activity in *Pheritima posthuma*. Time of dysfunction associated time of death was used as an analysis parameter. helminthic change state (10 mg/ml) was used as a customary drug. Phytochemical take a look at discovered the presence of alkaloids, flavanoids, glycosides, carbohydrates, phenoplast compounds and tannins. Concentration dependent anthelmintic result was discovered with the extract. once one hundred mg/ml concentration of ethanolic extracts of the polyherbal formulation was more it showed dysfunction of take a look at worm (earthworm) at eighteen minutes and death at thirty-nine minutes. The polyherbal formulation has shown substantial anthelmintic activity exploitation in vitro model on earthworms which can be attributed to the polyphenols gift within the extract.

## I. INTRODUCTION

India is one of the world's leading biodiversity with the presence of over 45000 different plant species. India is perhaps the largest producer of medicinal herbs & is rightly called the botanical garden of the world. Medicinal herbs have been for thousands of years is one form or another under the indigenous systems of medicine like Ayurveda, siddha, unani since independence in 1947. India had made tremendous progress in agro technology standardization, quality control & research development etc.

The world have received greater attention in recent time because of its diversity of curing diseases, safety & well tolerated remedies compared to the conventional medicines. The herbs natural combination of constituents as a whole or naturally occurring remedies which has proved to be more effective & safe than conventional medicines.

The ability of herbal medicine to affect the body systems depends on the chemical constituents that it contains.

Research on isolated plant constituents are of greater importance.

The study of diseases & their treatment must also have been contemporaneous with the dawn of the human intellect.

It is well known plant generally own their virtues as medical agents to certain characteristic alkaloid & principles present in them because a complete & full chemical analysis of the medicinal plants of India have not yet been performed.

## II. MATERIALS AND STRATEGIES

### Extraction Method:

The leaves powdery material was extracted by continuous hot extraction methodology exploitation soxhlet equipment. Powdery leaves were taken in a very white textile and through the flask containing 500ml of boiling volatile solvent fermentation alcohol and maintain temperature at 70°C. The vapour arising within the flask passes by the aspect tube into the condenser. The soluble matter dissolved from the drug within the extractor remaining within the flask. Finally the soluble matter was distillates then evaporate to xerotes in china dish exploitation heating mantle at temperature 70°C.

The preliminary photochemical analysis of leaves of *Ocimum sanctum*, *Coriandrum sativum*, *rosid dicot genus pinnata*, *epipremnum aureum* and *curcuma longa* showed the presence of alkaloids, flavanoids, glycosides, carbohydrates, phenoplast compounds and tannins.

### PRELIMINARY PHYTOCHEMICAL STUDIES

S.no	Extracts	Alkaloids	Carbohydrate	Tannins	Volatile oils	Glycoside	Phytosterols	Triterpene	Flavonoids	Phenolic Compounds
1.	MIXTURE (OCEPL)	+	+	+	+	+	+	+	+	+
2.	OS	-	+	+	+	-	-	-	+	+
3.	CS	-	+	+	+	-	+	-	+	+
4.	CL	-	+	+	+	+	-	+	+	+
5.	PP	+	+	+	-	-	-	-	+	+
6.	EA	-	+	+	+	-	-	+	+	+

OS: *Ocimum sanctum*, CS: *Coriandrum Sativum*, PP: *rosid dicot genus pinnata*, EA: *epipremnum aureum*, CL: *Curcuma longa*

**Anthelmintic activity**Worm: *Pheritima posthuma*

Control: solution

Cluster I: OCEPL extract      Cluster II: OS extract  
 Cluster III: CS extract      Cluster IV: CL extract  
 Cluster V: PP extract      Cluster VI: EA extract

**Procedure:**

The methodology delineated by Dashetal was used for evaluating anthelmintic activity. *Pheritima posthuma* (obtained from gardening department, Madurai, Tamilnadu, India) of roughly equal size (15 cm) was divided into nineteen teams. Every cluster consists of six earth worms of same kind and treated with any of the subsequent.

Fifty ml of take a look at resolution containing twenty, fifty and one hundred mg /ml of take a look at extracts and helminthic change state (10mg/kg).

The mean solar time of dysfunction and death was recorded in minutes. The dysfunction time was recorded once no movement of any kind might be discovered except once the worms were agitated smartly. Time for death of worms was recorded, once worms were neither moved whereas agitated smartly, nor once lordotic in heat water (50°C).

Anthelmintic activity of extracts

Category	Dose	Time of dysfunction Mean $\pm$ S.E.M (min)	Time of death Mean $\pm$ S.E.M (min)
Piperazinecitrate	10 mg/ml	26 $\pm$ 0.04*	47 $\pm$ 0.29*
MIXTURE (OCEPL)	20 mg/ml	22 $\pm$ 0.91	44 $\pm$ 0.37
	fifty mg/ml	20 $\pm$ 0.93*	41 $\pm$ 0.36*
	one hundred mg/ml	18 $\pm$ 0.75*	39 $\pm$ 0.90*
OS	20 mg/ml	64 $\pm$ 1.79	72 $\pm$ 1.75
	50 mg/ml	54 $\pm$ 1.70*	63 $\pm$ 1.63
	100 mg/ml	45 $\pm$ 0.76*	56 $\pm$ 0.52*
CS	20 mg/ml	54 $\pm$ 1.71	62 $\pm$ 0.65
	50 mg/ml	49 $\pm$ 1.63	56 $\pm$ 0.54*
	100 mg/ml	42 $\pm$ 0.53*	49 $\pm$ 0.32*
CL	20 mg/ml	54 $\pm$ 1.36	69 $\pm$ 0.75
	50 mg/ml	46 $\pm$ 1.72	65 $\pm$ 0.61*
	100 mg/ml	40 $\pm$ 0.92*	59 $\pm$ 0.72*
PP	20 mg/ml	48 $\pm$ 1.31	72 $\pm$ 0.55
	50 mg/ml	42 $\pm$ 1.53	66 $\pm$ 0.74*
	100 mg/ml	36 $\pm$ 0.58*	59 $\pm$ 0.42*
EA	20 mg/ml	48 $\pm$ 1.75	67 $\pm$ 0.35
	50 mg/ml	45 $\pm$ 1.53	61 $\pm$ 0.64*
	100 mg/ml	41 $\pm$ 0.64*	55 $\pm$ 0.72*

Student "t" test take a look at, \*P< 0.001 (Compared to standard) was thought of important.

**III. RESULTS & DISCUSSION**

Phytochemical take a look at discovered the presence of alkaloids, flavanoids, glycosides, carbohydrates, phenoplast compounds and tannins. Concentration dependent anthelmintic result was discovered with the extract.

The OCEPL extracts showed important anthelmintic activity. Once one hundred mg/ml concentration of ethanolic extracts of the polyherbal formulation was more it showed dysfunction of take a look at worm (earthworm) at eighteen minutes and death at thirty-nine minutes. The anthelmintic activity of extracts might be because of the presence of phenoplast compounds.

Although altogether experimental concentrations there's variation within the time to paralyze the worms, however once it gets paralytic, it took terribly short time for the parasites to die. this might be prompt that combination of plant extracts possesses vermifugal activity in nature and will exert a reversible action on the contractor system of the worms and therefore the inactiveness caused would lead the parasite to be swept back out of the host's body

Dose-dependent effectiveness was additionally discovered with exposure to numerous concentrations of every combination, as a rise in concentration, shortens the dysfunction amount.

**IV. CONCLUSION**

Thus, these plants besides having vermifugal/vermicidal activity additionally showed a synergistic result once treated together.

The polyherbal formulation has shown substantial anthelmintic activity exploitation in vitro model on earthworms which can be attributed to the polyphenols gift within the extract.

**REFERENCES**

- [1]. Gazi Shaikh International journal of ayurvedic & herbal medicine 2(3) June. (455463) 456 year (2012).
- [2]. American Family Physician www.aafp.org/afp Volume 87, Number 9 May 1, 2013
- [3]. Menter A, Gottlieb A, Feldman SR, et al.Guidelines of care for the management of psoriasis and psoriatic arthritis: Section 1. Overview of psoriasis and guidelines of care for the treatment of psoriasis with biologics. J Am Acad Dermatol.,58(5):826-850.4. Dung NX, Truong PX, Ky PT, Leclercq PA. Constituents of the leaf oil of *Curcuma domestica* L. from Vietnam. J. Essential oil Res; 1997. p. 677 (2008)
- [4]. Mathan C Nisha, Sevanan Rajeshkumar. (Survey of crude drugs from Coimbatore city).Indian Journal of Natural Products and Resources, 2010; 1(3): 376-383.5.
- [5]. Nambiar, V., Seshadri, S.: Retention of total and beta carotene from fresh radish leaves in steamed, sautéed and baked products of Western India. J Food Science Techn. 38 (5): 458-461(2001).
- [6]. Nambiar, V., Bhadalkar, K., Daxini, M.: Drumstick leaves in the ICDS-SFP. Ind J Peadiat.70 (5):11-15 (2003).Richmond R, Pombo-Villar E Chromatogr. A. Gas Chromatography-Mass Spectrometry coupled with Pseudo-Sadtler retention indices for the identification of components in the Essential Oil of *Curcuma*

- longa Chromatogr. A;1997. p. 303.
- [7]. Colombo AL, Nucci M, Park BJ, Nouér SA, Arthington-Skaggs B, et al. ( Epidemiology of candidemia in Brazil: a nationwide sentinel surveillance of candidemia in eleven medical centers. J Clin Microbiol 44: 2816–2823. doi: 10.1128/jcm.00773-06(2006)
- [8]. Pfaller M, Neofytos D, Diekema D, Azie N, Meier-Kriesche H, et al. Epidemiology and outcomes of candidemia in 3648 patients: data from the Prospective Antifungal Therapy (PATH Alliance) registry, 2004–2008. Diag Microbiol Infect Dis 74: 323–331. (2012)
- [9]. Messer SA, Moet GJ, Kirby JT, Jones RN Activity of contemporary antifungal agents, including the novel echinocandin Anidulafungin, tested against *Candida* spp., *Cryptococcus* spp., and *Aspergillus* spp.: Report from the SENTRY Antimicrobial Surveillance Program (2006 to 2007). Clin Microbiol J 47: 1942–1946. doi: 10.1128/jcm.02434-08 (2009)
- [10]. Garg SN, Bansal RP, Gupta MM, Kumar S. Variation in the Rhizome Essential Oil and Curcumin contents and oil quality in the land races of turmeric, *Curcuma longa* of North Indian Plains. Flavour Fragr. J; 1999. p.315-8.
- [11]. Martins AP, Salgueiro L, Gonçalves MJ, da Cunha AP, Vila R, Cañigueral S, Mazzoni V, Tomi F, Casanova J. Essential oil composition and antimicrobial activity of three Zingiberaceae from S. Tomé e Príncipe. Planta Med 2001;67:580-4.
- [12]. Behura S, Srivastava VK. Essential Oils of Leaves of *Curcuma* Species; 2004. p. 109-10.
- [13]. Balcerek M, Matławska I. Preventive role of curcumin in lung cancer. Przegl Lek 2005;62:1180-1.
- [14]. Farid et al. Effect of *Ocimum basilicum* on glucose and lipids metabolism. 2009 187-199.
- [15]. Gupta S K, Prakash J, Srivastava S V. Validation of claim of *Tulsi*, *Ocimum sanctum* Linn as a medicinal plant. Indian J Experimental Biology 2002, 40(7):765–773( 2002)
- [16]. Hannan J M A, , Marenah L, Ali L, Rokeya B, Flatt P R and Abdel-Wahab YHA *Ocimum sanctum* leaf extracts stimulate insulin secretion from perfused pancreas, isolated islets and clonal pancreatic b cells. Journal of Endocrinology 2006, 189: 127–136
- [17]. Joglekar GV, Chaudhary NY & Aiman R. Effect of indigenous plant extracts on glucose-absorption in mice. Indian Journal of Physiology and Pharmacology 1959, 3: 76
- [18]. Jyoti S, Satendra S, Sushma S, Anjana T, Shashi S. Antistressor activity of *Ocimum sanctum* (*Tulsi*) Against experimentally induced oxidative stress in Rabbits. Methods Find Exp Clin Pharmacol. 2007, 29(6):411-6.
- [19]. Karthikeyan, P. Gunasekaran, N. Ramamurthy and S. Govindasamy Anticancer Activity of *Ocimum Sanctum*. Summary Pharmaceutical Biology 1999, Vol.37, No. 4, Pages 285-290 .
- [20]. Kath R K & Gupta R K. Antioxidant activity of hydroalcoholic leaf extract of *Ocimum sanctum* in animal models of peptic ulcer. Indian J Physiol Pharmacol 2006, 50 (4): 391–396.
- [21]. Anonymous, The Wealth of India; Raw Material, CSIR; New Delhi; 1950.p. 402.
- [22]. Nigam MC, Ahmed A. *Curcuma longa*: Terpenoid composition of its Essential Oils. Indian Perfumer; 1991. p. 355.
- [23]. Zachariah TJ, Baby KN. Effect of Storage of fresh turmeric Rhizome on Oleoresin and Curcumin contents. J. Spice Arom. Crops; 1992. p. 55-58.
- [24]. Shah NC. Traditional use of Turmeric (*Curcuma longa*) in India J Med Arom Plant Sci; 1997. p. 948-95.
- [25]. Masuda T, Isobe J, Jitoe A, Nakatani N. Antioxidative Curcumions from Rhizomes of *Curcuma Xanthorrhiza*. Phytochemistry; 1992. p. 3645-47. . Singh S, Kher A Biological effects of curcumin and its role in cancer chemoprevention and therapy. Anticancer Agents Med Chem 2006;6:259-70.
- [26]. Ammon HP, Wahl MA. Pharmacology of *Curcuma longa*. Planta Med 1991;57:1-7.
- [27]. Kumar V. The Pharma Review Clinical Trials in India: Balancing Economic Opportunity with the Public Health Context. Kongposh Publication Pvt. Ltd. New Delhi.
- [28]. Jennings W, Shibamoto T. Qualitative Analysis of Flavour and Fragrance volatiles by Glass Capillary Column Gas Chromatography. Academic Press: New York. 1980.
- [29]. Swinggarm AA, Silverstein RM. Monoterpenes, Aldrich Chemicals Co. Inc.: Milwaukee, WI, 1987.
- [30]. Adams RP. Identification of Essential Oils by Ion Trap Mass Spectroscopy. Academic Press: San Diego CA, 1989.
- [31]. Dwivedi S., Dwivedi A., Kapadia R. and Kaul S.. Anthelmintic activity of alcoholic and aqueous extract of fruits of *Terminalia chebula* Retz., Ethno. Leaflets, 12:741-743. (2008)
- [32]. Kailashraj R. and Kurup A., Ind. J. Phar., 1962, 74.
- [33]. Rios, J.L., M.C., Recio and A., Villar, "Screening methods for natural products with antimicrobial activity: Journal of Ethnopharmacol 1988; 23, p. 127-149