



Acute and sub-acute (28-days) oral toxicity studies of *Eraippu noi chooranam*

¹Dr. Amala Hazel .A.M , ²Dr. Pattarayan. R , ³Dr.Banumathy. V

¹Lecturer, Kuzhanthai Maruthuvam Department, National Institute of Siddha, Chennai

²Former HOD, Kuzhanthai Maruthuvam Department, National Institute of Siddha, Chennai.

³Director, National Institute of Siddha, Chennai

*Corresponding Author: dramalaarulldhas@gmail.com

Abstract

Purpose: Eraippu Noi Chooranam is a herbal compound medicine mentioned in Siddha research pharmacopoeia and some modifications are introduced to increase its potentiality. It is indicated for asthma, chronic bronchitis and flatulence¹. **Aim and objective:** The objective of this study was to investigate the acute and sub-acute toxicity of ENC in Wistar rats. **Materials and Methods:** In the acute test, the limit test dose of 2000 mg/kg was administered to Wistar rats and then observed individually 1 h post-dosing, and at least once daily for 14 days. Sub-chronic toxicity was evaluated after administering daily oral doses of 300,600 and 900 mg/kg body wt., for 28 days to the rats, Biochemical and haematological assessments as well as body and relative organ weights of the rats were carried out. **Results:** The limit dose of 2000 mg/kg did not cause any mortality or signs of acute toxicity in the rats tested during the observation period. In the sub-chronic tests, the results did not show any treatment-related abnormalities in terms of haematological and biochemical parameters. The weekly body and organ weight of the rats showed no significant differences between the control and the rats treated with the ENC. **Conclusion:** Our results suggest that ENC is relatively safe when administered orally in rats.

Keywords: *Eraippu Noi Chooranam* (ENC), Acute and sub-chronic toxicity, Biochemical parameters, Haematological analysis, Wistar rats.

Introduction

Natural products, including plants, animals and minerals have been the basis of treatment of human diseases. History of medicine dates back practically to the existence of human civilization. The current accepted modern medicine or allopathy has gradually developed over the years by scientific and observational efforts of scientists. However, the basis of its development remains rooted in traditional medicine and therapies. (Patwardhan *et al.*, 2004). Whenever we administer a chemical substance to a biological system, different types of interactions can occur and a series of dose-related responses result.

In most cases these responses are desired and useful, but there are a number of other effects which are not advantageous. The types of toxicity tests which are routinely performed by pharmaceutical manufacturers in the investigation of a new drug involve acute, sub-acute and chronic toxicity. Acute toxicity is involved in estimation of LD50 the dose which has proved to be lethal (causing death) to 50% of the tested group of animals. Determination of acute oral toxicity is usually an initial screening step in the assessment and evaluation of the toxic characteristics of all compounds. (Akhila *et al.*, 2007). Acute toxicity is

produced after administration of a single dose or multiple doses in a period not exceeding 24 hours, up to a limit of 2000 mg/kg. Objective of acute toxicity studies is to identify a dose causing major adverse effects and an estimation of the minimum dose causing lethality (Robinson *et al.*, 2007). In recent times there is an increasing awareness and interest in medicinal plants and their preparations commonly known as herbal medicines (Steve *et al.*, 2009). The major hindrance to the use of traditional herbal preparations is the lack of scientific and clinical data in support of better understanding of the efficacy and safety of the drugs.

Materials and Methods

Procurement and Authentication of Raw Drugs

The ingredients of *ENC* was purchased from reputed country shop. The raw drugs were identified and authenticated by medicinal botanist of National Institute of Siddha. All the ingredients were powdered and bottled up at the ratio mentioned.

Preparation of Eraippu Noi Chooranam.

The raw drugs were purified as mentioned in Siddha literature^{3,4}. *ENC* was prepared using the procedure described in Siddha literature with slight modifications.

Animal Care and Husbandry:

The study protocol involving animals was reviewed and approved by Institutional Animal Ethical Committee (IAEC), KMCH College of pharmacy, Kovai estate, with the experimental protocol number IAEC NO: KMCRET/MD(S)/10/2014-15. Experiments were performed as per the instructions prescribed by the Committee for the Purpose of conduct and Supervisions of Experiments on Animals (CPCSEA), Ministry of Environment and Forest, Government of India. Male and female Wistar albino rats, (140–160 g) obtained from Sree Venkateshwara Enterprises Pvt. Ltd, Bangalore, were housed in the animal house KMCH College of pharmacy, Kovai estate. Each group of rats was separately housed in polypropylene cages in a well ventilated room under an ambient temperature of $22\pm 3^{\circ}\text{C}$ and 30-70% relative humidity, with a 12-h light/dark artificial light cycle. They were provided with food (SaiDurga Animal Feed, Bangalore) and purified water *ad libitum*. All the animals were acclimatized to the laboratory conditions at least for 7 days prior to experimentation.

Acute toxicity test

The acute oral toxicity study was performed in accordance with Organization for Economic Cooperation and Development (OECD) test guideline 423⁴. The limit test dose of 2000 mg/kg was used as stipulated in Organization for Economic Cooperation Development (OECD) guidelines. The test drug suspension was administered at a dose of 2000 mg/kg once orally to the fasted rats. In the experiments the observations like body weight, clinical signs and gross pathology were common and are as follows

Mortality or morbidity was noted

- Following test item administration weekly body weight was recorded.
- Clinical signs such as Lethality, Convulsion, Tremor, Straub tail, Sedation, Excitation, Abnormal gait (rolling), Abnormal gait (tiptoe), Jumps, Motor coordination, Loss of balance, Fore paw treading, Writhes, Piloerection, Stereotypies (chewing), Stereotypies (Head movements), Head twitches, Scratching, Respiration, Aggressiveness, Fear, Reactivity to touch, Muscle tone, Loss of righting Reflex, Ptosis, Exophthalmos, Loss of grasping, Akinesia, Catalepsy, Loss of traction, Loss of corneal reflex, Analgesia, Defecation, Salivation, Lacrimation, Others: were observed at approximately 30 mins, 1hr, 2hr and 4hr on day 1 and daily thereafter for 14 days.
- At the end of 14 days, the experimental animals were necropsied and investigated for gross pathological examination

Sub-acute toxicity test

Repeat-dose oral toxicity study was carried out according to OECD guideline 407⁵

Justification for Dose Selection

As stated in results of acute toxicity studies in wistar rats indicated that *ENC* was nontoxic up to the maximum dose level of 2000 mg/kg body weight. On the basis of these results, the doses selected for the study was 300 mg/kg, 600 mg/kg and 900 mg/kg body weight. The oral route was selected for use because oral route is considered to be a proposed therapeutic route.

Preparation and administration of dose

Repeated-dose oral toxicity study was carried out according to OECD guideline 407⁶. The animals were divided into six groups of 10 animals each (5 males and 5 females). Group 1 received distilled water and served as control. Groups II, III and IV received test drug ENC with distilled water at doses of 300,600 and 900 mg/kg body wt, respectively. Group V (Vehicle control) and Group VI ENC high dose (900 mg/kg) were included as satellite study groups to determine the delayed occurrence, or persistence of, or recovery from toxic effects. The test drug was administered daily for 28 days the same time daily and observed at least twice daily for morbidity and mortality. Body weights of the animals were evaluated weekly

The satellite groups were scheduled for follow-up observations for the next 14 days without test drug administration. The test drug suspensions (ENC with distilled water) were freshly prepared every day and administered orally (gavage) once daily for 28 consecutive days. Initial body weight of all the groups was recorded. The animals were monitored closely for signs of toxicity throughout the course of study. Appearance and behavior pattern were assessed daily and any abnormalities in food and water intake were registered.

On the 29th day, after an overnight fast, the rats were anaesthetized with ketamine and blood sample for haematological and biochemical analysis were collected into tubes with and without EDTA, respectively. Haemoglobin, Red blood cell count, White blood cell count, Mean Corpuscular Haemoglobin (MCH), Packed Cell Volume (PCV) was determined using fully automated haematology analyzer. Biochemical analysis was performed on serum obtained after centrifugation of total blood

(without anticoagulant) at 2500 rpm for 15 min. Biochemical parameters: alanine aminotransferase (ALT), aspartate aminotransferase (AST), creatinine, alkaline phosphatase, glucose, total proteins and urea also determined.

Histopathology

Necropsy was done in all animals on day 29 except the satellite groups for which it was done on day 42. After blood collection, all the animals were euthanized for gross pathological examinations of all major internal organs. The organs such as brain, heart, liver, spleen, kidneys, were weighed and relative organ weights were calculated. The organs were fixed in 10% neutral buffered formalin, trimmed and a 5µ thickness of tissue sections were stained with hematoxylin and eosin for histopathological investigation.

Statistical analysis

All the values are expressed as mean ± S.E.M. The data were statistically analyzed by one-way ANOVA followed by Dunnet-t test. P values < 0.05 were considered significant.

Results

Acute toxicity studies

The limit dose of 2 g/kg did not cause mortality or any sign of acute toxicity in the three rats dosed for a short period (48 h) and long period (14 days).

No behavioural changes and death were observed at the end of the treatment. Similarly, no significant differences in bodyweight were observed between control and treated groups during this period (see Table1).

Table 1: Dose finding experiment and its behavioral Signs of Toxicity

No	Dose mg/kg	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20
1	Control	+	-	-	+	+	+	-	-	-	-	-	-	-	-	-	-	-	-	-	-
2	2000mg/kg bw	+	-	-	+	+	+	-	-	-	-	-	-	-	-	-	+	+	-	+	+

1.Response 7. Decreased Motor Activity 8.Tremors 9.Convulsions 10. Muscle Spasm 11. Catatonia 12. Muscle relaxant 13. Hypnosis 14.Analgesia 15.Lacrimation 16.Exophthalmos 17.Diarrhoea 18.Writhing 19.Respiration 20. Mortality

Sub acute toxicity study

There were no treatment-related toxicity signs and mortality observed in both sexes of rats treated at 300, 600 and 900 mg/kg orally for a period of 28 days and

in the satellite group of rats. No significant difference in body weight gain was observed between control and treated groups during the study (Table 2). Feed and water consumption of ENC treated groups were found to be insignificant in both the sexes when compared to

control. Hematological parameters such as haemoglobin, red blood cells, white blood cells, platelet count, mean corpuscular haemoglobin, were found to be well within the clinical range of rats⁶ in experimental groups (Table3). There were no significant difference in plasma biochemical profile such as glucose, total cholesterol, triglycerides, total protein, alkaline phosphatase, bilirubin, creatinine, blood urea (Table 4) observed between control and treated groups. The levels of liver marker enzymes like SGOT and SGPT were found to be well within the clinical range of rats⁶ in ENC treated groups (Table4). There were no significant differences in organ weight

of brain, heart, liver, spleen, lungs, kidneys and sex organs recorded between the control and ENC groups (Table5). In our study, hisopathological examinations in control and high dose group revealed no abnormalities. There were no hematological, biochemical and histopathological alterations observed with ENC administration even at 900 mg/kg/day in rats for a period of 28 days compared to control. The No-Observed Adverse Effect Level (NOAEL) of ENC was estimated to be greater than 900 mg/kg/day in rats. Hence, it can be concluded that ENC is safe for oral administration.

Table 2: Effect of Eraippu Noi Chooranam on Body weight of experimental Wistar rats in repeated oral toxicity study

Treatment	1 st day	7 th day	14 th day	21 st day	28 th day	35 th day	42 nd
Control	147.90±1.27	151.30±1.33	154.80±1.19	158.20±1.03	161.50±1.09	-	-
ENC300mg /kg /day	147.50±1.12	150.60±1.48	154.50±1.45	157.60±1.38	160.80±1.35	-	-
ENC600mg /kg /day	147.60±1.61	151.10±1.54	154.70±1.51	158.30±1.48	161.20±1.67	-	-
ENC900mg /kg /day	146.60±1.48	150.00±1.52	153.40±1.63	156.70±1.61	159.90±1.52	-	-
Satellite group							
Control	148.50±1.08	152.20±0.99	155.70±0.90	159.50±0.82	163.00±0.75	166.40±0.65	169.60±0.62
ENC900mg /kg /day	147.40±1.42	150.90±1.50	154.30±1.54	157.70±1.46	161.10±1.44	164.40±1.45	167.70±1.40

Comparison was made between vehicle control and test groups using the one-way ANOVA test followed by Dunnetts test. Values are mean ± SEM for 10 rats in each group (*P* < 0.05).

Table 3: Effect of Eraippu Noi Chooranam on Hematological Parameters of experimental Wistar rats in repeated oral toxicity study

Treatment	Hematological parameters							
	WBC (10 ³ /uL)	RBC(10 ⁶ /uL)	Hb g/dl	PCV %	MCH pg	Lymphocyte %	Monocyte %	Platelets
Control	8.70 ±0.20	7.02 ± 0.18	15.36 ±0.47	42.95 ±1.36	22.14 ±0.53	82.40 ±1.23	2.70 ± 0.26	827.70 ±23.92
ENC300mg /kg /day	8.71 ± 0.26	6.94 ± 0.19	14.62 ± 0.63	41.18 ± 1.07	22.21 ± 0.67	81.60 ± 1.71	2.50 ± 0.34	803.80± 25.11
ENC600mg /kg /day	8.57 ± 0.28	6.96 ±0.17	15.43 ±0.52	39.27 ± 0.74	20.65 ± 0.48	84.30 ±1.46	2.60 ± 0.31	844.40±27.69
ENC900mg /kg /day	8.70 ± 0.27	6.95 ± 0.26	15.54 ± 0.59	39.96 ± 1.33	21.35 ± 0.70	84.40 ± 1.38	2.70 ±0.26	806.10±30.02
Satellite group								
Control	8.61 ± 0.28	6.94 ±0.21	15.69 ±0.51	41.64 ± 0.06	21.81 ±0.70	83.90 ±1.30	2.30 ±0.26	802.70±28.79
ENC900mg /kg /day	8.51 ± 0.31	7.10 ± 0.28	15.59 ±0,36	40.04 ± 1.29	20.76 ± 0.80	83.60 ± 1.61	2.50±0.34	843 ±29.36

Comparison was made between control and test groups using the one-way ANOVA test followed by Dunnetts test. Values are mean ± SEM for 10 rats in each group (*P* < 0.05).

RBC: red blood corpuscles; HB: hemoglobin; PCV: packed cell volume; MCH: mean corpuscular hemoglobin; WBC: white blood cells.

Table 4: Effect of Eraippu Noi Chooranam on biochemical parameters of experimental Wistar rats in repeated oral toxicity study

Treatment	Biochemical parameters								
	Glucose mg/dl	TG Mg/dl	Cholesterol mg/dl	SGPT U/I	ALP U/I	SGOT U/I	T. Protein g/dl	Creatinine mg/dl	Urea Mg/dl
Control	89.50 ±1.65	74.35 ±2.80	100.10 ±2.45	65.02 ± 2.32	296.93 ±10.75	144.77 ± 2.42	7.08 ± 0.27	0.93 ± 0.04	30.05 ±1.59
ENC300mg /kg /day	93.00 ±1.86	76.27 ±1.78	101.00 ±3.79	71.97 ±1.65	291.85 ±14.25	153.89 ± 2.06	7.48 ±0.32	0.71 ±0.06	35.01 ±2.96
ENC600mg /kg /day	88.80 ±1.97	78.49 ±2.60	97.37 ±3.11	67.19 ±4.37	293.26 ±9.81	151.17 ±3.77	7.36 ±0.41	0.84 ±0.05	37.00 ±1.13
ENC900mg /kg /day	88.80 ±1.81	75.58 ± 1.85	95.02 ± 2.71	62.55 ±2.02	296.03 ±12.23	151.33 ±3.61	7.54 ±0.25	0.74 ±0.06	34.06 ± 2.24
Satellite group									
Control	94.20 ±1.84	74.11 ± 2.51	102.40 ± 2.59	66.47 ± 2.76	279.35 ± 9.42	147.15 ± 3.31	8.17 ± 0.31	0.87 ± 0.04	32.81 ±2.22
ENC900mg /kg /day	94.00 ± 2.25	72.00 ± 1.51	96.35 ±1.75	66.0 ± 2.73	294.58 ± 9.92	162.64 ± 5.06	7.30 ± 0.43	0.66 ± 0.05	32.18 ± 2.28

Comparison was made between vehicle control and test groups using the one-way ANOVA test followed by Dunnett's test. Values are mean ± SD for 10 rats in each group ($P < 0.05$).

TGL: triglycerides, SGOT: serum glutamic oxaloacetic transaminase, SGPT: serum glutamic pyruvic transaminase, ALP:alkaline phosphatase.

Table 5: Effect of Eraippu Noi Chooranam on Organ weight of experimental Wistar rats in repeated oral toxicity study

Treatment	Relative organ weight (g %)							
	Brain	Lungs	Heart	Liver	Kidney	Spleen	Sex organs	
							Testis (n=5)	Ovaries (n=5)
Control	0.83±0.06	0.85±0.02	0.57±0.01	3.72±0.02	0.99±0.03	0.29±0.01	1.07±0.02	0.05±0.00
ENC300mg /kg /day	0.82±0.06	0.87±0.02	0.57±0.02	3.74±0.10	0.95±0.03	0.30±0.01	1.12±0.06	0.05±0.00
ENC600mg /kg /day	0.83±0.06	0.84±0.02	0.58±0.01	3.66±0.10	0.97±0.02	0.29±0.01	1.12±0.04	0.06±0.01
ENC900mg /kg /day	0.84±0.06	0.87±0.02	0.59±0.01	4.23±0.10	1.01±0.04	0.30±0.01	1.08±0.01	0.05±0.01
Satellite group								
Control	1.20±0.01	0.90±0.03	0.55±0.01	3.79±0.10	0.96±0.02	0.28±0.02	1.03±0.04	0.06±0.00
ENC900mg /kg /day	1.19±0.02	0.90±0.05	0.55±0.01	4.02±0.12	0.99±0.03	0.27±0.01	1.09±0.03	0.05±0.01

Comparison was made between control and test groups using the one-way ANOVA test followed by Dunnett's test. Values are mean ± SEM for 10 rats in each group ($P < 0.05$).

Discussion

Since ENC is in clinical use for treatment of respiratory diseases particularly for bronchial asthma for more than two decades, a limit test was performed in acute oral toxicity study. According to the OECD test guideline 423 when there is information in support of low or non-toxicity and immortality nature of the test material, then the limit test at the highest starting dose level (2000 mg/kg body weight) was conducted. There were no mortality and toxicity signs observed at 2000mg/kg. ENC can be classified under category-5 and LD₅₀ value was greater than 2000mg/kg in accordance with Globally Harmonised System of Classification and Labeling of chemicals and this provides us a direct relevance for protecting human and animal health. Therefore, it can be concluded that ENC when administered at single dose is non-toxic and can be used safely in oral formulations.

A 28-day repeated oral toxicity study was performed followed OECD test guideline 407 in both male and female wistar albino rats. Since examination of clinical signs plays major role in toxicological testing⁷ mortality and morbidity were recorded twice a day throughout the study. ENC did not produce any alterations in feed and water consumption and this reveals that it did not adversely affect the basic metabolic processes of the experimental animals. The haemopoietic system serves as important target for toxic chemicals and is a sensitive index for pathological conditions both in humans and animals⁸. In the present study, treatment with ENC did not produce any alteration in haematological parameters (i.e. RBC, WBC, haemoglobin, etc.), which indicate that ENC did not affect blood cells nor their production. Clinical biochemistry and hematological data holds significant role in determining the toxicity induced by drugs⁶ Transaminases (SGOT and SGPT) are good indicators of liver function and biomarkers to predict the possible toxicity of drugs⁹ Any elevation pertaining to these enzymes indicate their outflow into the blood stream due to damage in liver parenchymal cells.

There were no changes in the SGPT and SGOT levels which reveal that ENC did not affect liver function/or metabolism. In the present study, there were no treatment related abnormalities in renal function and other biochemical parameters suggesting that ENC is non-toxic. Histopathological studies provide supportive evidence for biochemical and haematological observations¹⁰ The organ weights were found to be non-significant between the control and ENC treated rats. No abnormality was recorded with respect to gross or histopathological examinations of all organs examined. Since there were no signs of

toxicity with respect to hematology, clinical chemistry, organ weight, gross and histopathological examinations noted in ENC satellite group, it can be inferred that ENC will not produce delayed onset of toxicity. Based on these results, the No Observed Adverse Effect Level (NOAEL) of "ENC" is greater than 900 mg/kg/day.

Conclusion

In accordance with Globally Harmonised System of Classification and Labeling of chemicals, ENC can be classified as Category 5. Based on 28 day repeated dose toxicity study, NOAEL of ENC is greater than 900 mg/kg/day. The present investigation substantiates, at least in part, the safety of ENC, which was found to be in line with the long history of its use in Siddha system of medicine.

Acknowledgments

The authors wish to acknowledge Mr.G. Ariharasivakumar, Associate Professor, Dept of Pharmacology, KMCH College of Pharmacy, Kovai Estate, Kalapatty Road, Coimbatore, 641048, for his guidance in Toxicological studies.

References

- 1).The Pharmacopoeia of siddha medicines- Dr.M.Shanmugavelu & Dr.G.D.Naidu.Published by Dr.G.D.Naidu Industrial labour welfare association ltd, Coimbatore.
- 2) R. Thiagarajan, Gunapaadam-Thathu Jeeva Vaguppu, Department of Indian Medicine and Homeopathy Publishers, B.I.M., 4th edition, 2004.
- 3) Sarakusuthi seimuraiyagal
- 4) OECD, OECD Guideline for Testing of Chemicals, Acute Oral Toxicity-Acute Toxic Class Method: TG 423-Adopted, OECD, Paris, France, 2001.
- 5) OECD, OECD Guideline for Testing of Chemicals, Repeated Dose 28-day Oral Toxicity Study in Rodents: TG 407-Adopted, OECD, Paris, France, 1995.
- 6) Petterino C, Argentino-Storino A: Clinical chemistry and haematology historical data in control Sprague-Dawley rats from pre-clinical toxicity studies. *Exp Toxicol Pathol.* 2006, 57: 213-219. 10.1016/j.etp.2005.10.002.
- 7) Stevens KR, Mylecraine L: Issues in chronic toxicology. *Principles and Methods of Toxicology.* Edited by: Hayes AW. 1994, New York: Raven Press, 673-third

- 8) Adeneyea AA, Ajagbonnab OP, Adeleke TI, Bellod SO: Preliminary toxicity and phytochemical studies of the stem bark aqueous extract of *Musanga cecropioides* in rats. *J Ethnopharmacol.* 2006, 105: 374-379. 10.1016/j.jep.2005.11.027.
- 9) Hilaly JE, Israili ZH, Lyouss B: Acute and chronic toxicological studies of *Ajuva Iva* in experimental animals. *J Ethnopharmacol.* 2004, 91: 43-50. 10.1016/j.jep.2003.11.009.
- 10) Matsuzawa T, Nomura M, Yonezawa H, Unno T: Selection of appropriate parameters, use of a quality control concept, and suitable statistical analyses for clinical pathology examination of animals in toxicity studies: results of a current survey by the Japanese Pharmaceutical Manufacturers Association. *Comp Haematol Int.* 1995, 5: 196-200. 10.1007/BF00368044.

Access this Article in Online	
	Website: www.ijarbs.com
	Subject: Siddha Medicine
Quick Response Code	

How to cite this article:

Amala Hazel .A.M , Pattarayan. R , Banumathy. V. (2016). Acute and sub-acute (28-days) oral toxicity studies of *Eraippu noi chooranam*. *Int. J. Adv. Res. Biol. Sci.* 3(6): 106-112.