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EVALUATION OF ANTIDEPRESSANT ACTIVITY OF ISOLATED PHYTOSTEROL FROM LEAVES OF *HOLOPTELEA INTEGRIFOLIA* (ROXB.) PLANCH IN EXPERIMENTAL ANIMALS

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ABSTRACT

Background: Majority of scientific documentation suggested promine nt role of Phytosterols towards Antidepressant activity. The main objective of the work was to evaluate Antidepressant activity of Holoptelea integrifolia isolated Phytosterol (HIIP) from petr oleum ether extract (PEHI) of leaves of Holoptelea integrifolia (Roxb) Planch. **Methods:** The Antidepressant activity of different doses of HIIP (10 a nd 30 mg/kg-p.o.) was evaluated using Tail Suspension Test (TST) in mice. **Results:** HIIP-30 mg/kg was more potent than HIIP-10 mg/kg for sho wing antidepressant activity. **Conclusions:** The results indicate that HIIP shows antidepressant activity which was dose dependent.

KEYWORDS: Tail Suspension Test, Holoptelea integrifoila Isolated Phytosterol, Depression, Fluoxetine.

1. INTRODUCTION

In traditional system of medicine, bark and leaves of *Holoptelea integrifoila* (HI) used as bitter, astringent, acrid, thermogenic, anti inflammatory, digestive, carminative, laxative, anthelmintic, depurative, repulsive, urinary astringent and in rheumatism.[1,2] In our previous studies the Antidepressant activity of petroleum ether and methanol extract of leaf of *Holoptelea integrifolia* in experimental animals was evaluated and it was found that petroleum ether extract (PEHI) has shown comparable effects with the standard drug and more significant antidepressant activity than methanolic extract (MHI).[3] On the similar lines

in this present study a phytosterol (HIIP) was isolated from petroleum ether extract and it was studied for Antidepressant activity.

2. MATERIALS AND METHODS

2.1 Chemicals and Drugs

Standard drug *fluoxetine* was obtained from Crescent Therapeutics limited, Himachal Pradesh. The test drug HIIP was prepared individually as suspension in distilled water with tragacanth (1%w/v) as a suspending agent. A gastric catheter was used for oral drug administration. All the solvents used for the extraction were of AR grade.

2.2 High Performance Thin Layer Chromatography (HPTLC) Instrumentation

HPTLC system of CAMAG, Muttenz, Switzerland, Anchrom Enterprises (I) Pvt. Ltd, Mumbai, consisting of sample applicator (Linomat 5), Twin trough chamber with lid (20×10 cm, CAMAG, Muttenz, Switzerland), UV cabinet (Aetron, Mumbai) with dual wavelength (254/366 nm) and the HPTLC photo documentation (Aetron, Mumbai) were used for study. UV spectra was recorded using CAMAG TLC Scanner – IV.

LC/MS was recorded using SHIMADZU- LC/MS 2020, IR spectra was recorded using SHIMADZU-IR PRESTIGE-21, NMR spectra was recorded using Mercury plus 300 MHZ NMR Spectrometer.

2.3 Plant Material, Extraction and Isolation of *Holoptelea integrifolia* phytosterol (HIIP) from petroleum ether extract by preparative TLC

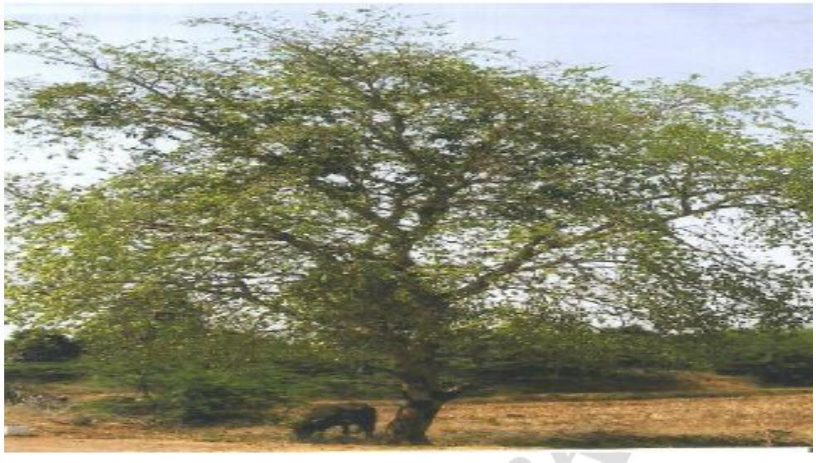


Fig. No. 1: *Holoptelea integrifolia* (Roxb.) Planch tree.

The dried and powdered leaves (1kg) of *Holoptelea integrifolia* was extracted with petroleum ether (b.p. 60-80ºC) for three times. After evaporation of the solvent under reduced pressure, the yield obtained was 4.8%w/w.

The petroleum ether extract was prepared in petroleum ether as a sample solution applied on Precoated silica gel aluminium plates 60F254, 20 cm x 10 cm with 250 µm thickness with CAMAG Linomat V (Switzerland) was used. The plates were washed by methanol and activated at 1200 C for 20 min before the start of chromatography. The sample solution was applied by using CAMAG microlitre syringe on the plates. The distance between the 2 bands was 5 mm with constant application rate of 1.0 µl/s was applied.

The compostion of mobile phase used for isolation of phytosterol was Chloroform: Ethyl acetate, in the volume ratio of 4:6 (v/v) and 20 ml of mobile phase was used per chromatography.

The plates were developed in 20 cm x 10 cm twin trough glass chamber saturated with filter paper Whatmann No.1 in mobile phase for 20 min at room temperature, and length of chromatogram run was 8.0 cm.

TLC plates were dried with the help of air dryer. Later on, densitometric scanning was performed with CAMAG TLC Scanner IV at 540 nm. The TLC Plate was dipped in Anisaldehyde Sulphuric acid reagent and then dried in oven at 110oC. Concentration of the compound was then determined.

The yield of HIIP obtained was 6 mg for a total of 40 preparative TLC Plates. In order to get sufficient quantity of HIIP, TLC plate of 1mm thickness was used. 20 gm of PEHI has given 228 mg, HIIP yield by using this method.[4-11]

2.4 Preparation of test samples

The test drug HIIP was prepared individually as suspension in distilled water with tragacanth (1%w/v) as a suspending agent. For the all pharmacological studies freshly prepared suspensions were used.

2.5 Animals

Albino wistar mice of either sex weighing between 20-30g were procured from Central Animal House, Rajah Muthiah Medical College & Hospital, Faculty of Medicine, Annamalai

University, Annamalai Nagar 608002, TamilNadu, India for experimental purpose. The animals were acclimatized to laboratory conditions for 7 days. The animals were supplied with commercially available standard diet. They were maintained at 25 ± 2° C and relative humidity of 45 to 55% and under standard environmental conditions (12 hour. light 12 hour. dark cycle). Water was allowed ad libitum under hygienic conditions. All animal studies were performed in accordance guideline of CPCSEA and Institutional Animal Ethical Committee (IAEC) of Central Animal House, Rajah Muthiah Medical College & Hospital, Annamalai University, Tamil Nadu, India(CPCSEA registration number-160/1999 /IAEC/CPCSEA, Proposal no:1029). All experiments were carried out between 12:00- 16:00 h.

2.6 Acute toxicity study

Healthy adult female Wistar albino mice were subjected for acute toxicity studies as per OECD-425 guidelines for isolated compound, HIIP. The test substances were administered orally in a single dose by gavages using a stomach tube. Mice were fasted prior to dosing (food was withdrawn overnight and water was withdrawn 3-4 h before drug administration). Following the period of fasting, the mice were weighed and the test substance, HIIP was administered. After the administration of the substances, food was withheld for1-2 h in mice. Mice were observed for its onset and duration of behavioural changes, toxicity and mortality upto 24h and observations were done for a period of 14 days after acute toxicity. For determining LD50 value, HIIP was administered in mice as per OECD-425 guidelines, the isolated compound HIIP was given as 100,200,300,400 and 500 mg/kg/p.o/b.wt. If the first animal survived; the second animal received a higher dose. If the first animal died or appeared moribund, the second animal received a lower dose.

2.7 Evaluation of antidepressant activity of HIIP using Tail Suspension Test (TST) model in mice

On the 14th day immediately after administration of last dose, each mice individually was suspended on the edge of the table 50 cm above the floor by adhesive tape placed 01 cm from the tip of the tail for the period of 05 minutes using stop watch and immobility duration was recorded. Mice are considered immobile when they hanged passively and completely motionless. In this test, suspending mice suspended upside down leads to characteristic behavior immobility which resembles to human depression. The decrease in immobility duration was considered as behavioral profile that indicated antidepressant like action.

Group 1- Control group of mice treated with vehicle 10- ml/kg/p.o/b.wt for 14 days.

Group 2- Test group of mice treated with low dose of HIIP- 10 mg/kg/p.o/b.wt for 14 days.

Group 3- Test group of mice treated with high dose of HIIP- 30 mg/kg/p.o/b.wt for 14 days.

Group 4-Test group of mice treated with (std.) Fluoxetine-10 mg/kg/p.o/b.wt for 14 days.

STATISTICAL ANALYSIS

Comparison was made against the vehicle treated control group. All the data was presented as Mean ± SEM. The data was analyzed by one way analysis of variance (ANOVA) followed by Dunnett's 't' test ,*P<0.05, **P<0.01,***P<0.001.

3. RESULTS AND DISCUSSION

3.1 Acute toxicity study

Table. 1: LD50 Values of isolated compound HIIP.

Isolated Compound	Up and down doses in mg/kg/p.o/b.wt				
HIIP	100	200	300	400	500
	5/5	4/5	3/5	0/5	-

The LD50 of HIIP, was performed at the dose level of 100 to 300 mg/kg/p.o/b.wt. At 300 mg /kg/p.o/b.wt two mice were dead. And hence, the LD50 dose of HIIP, was fixed at 300 mg/kg /p.o/b.wt.

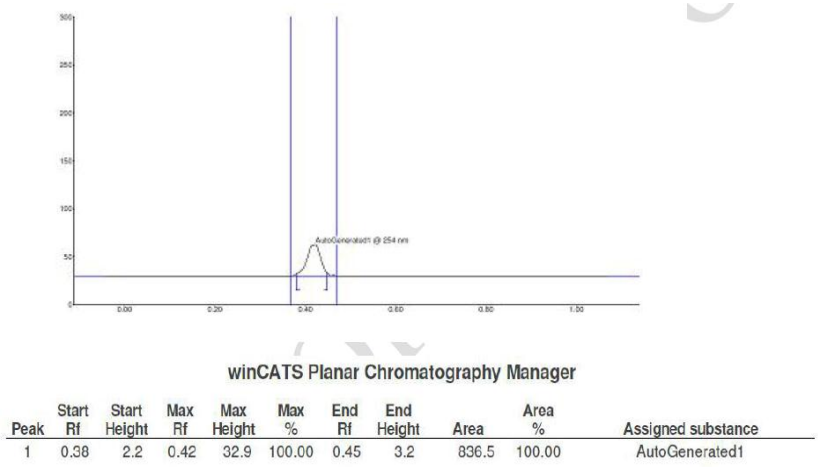


Fig.No.2: HPTLC chromatogram of a new phytosterol (HIIP) from petroleum ether extract of leaves of *Holoptelea integrifolia* by preparative TLC (Rf : 0.42).

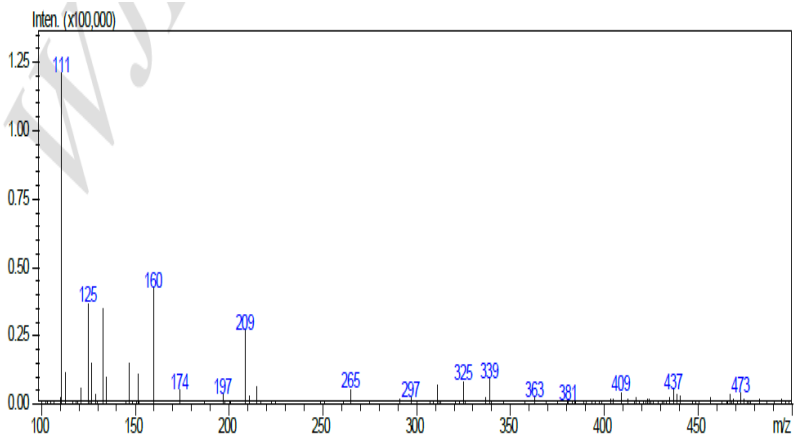
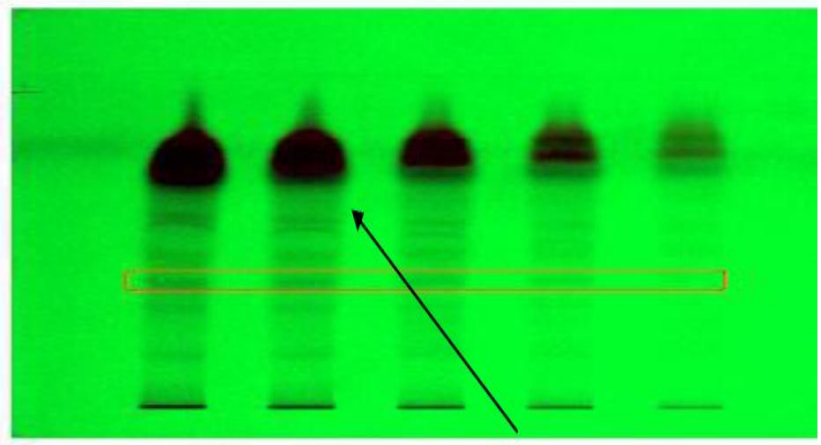
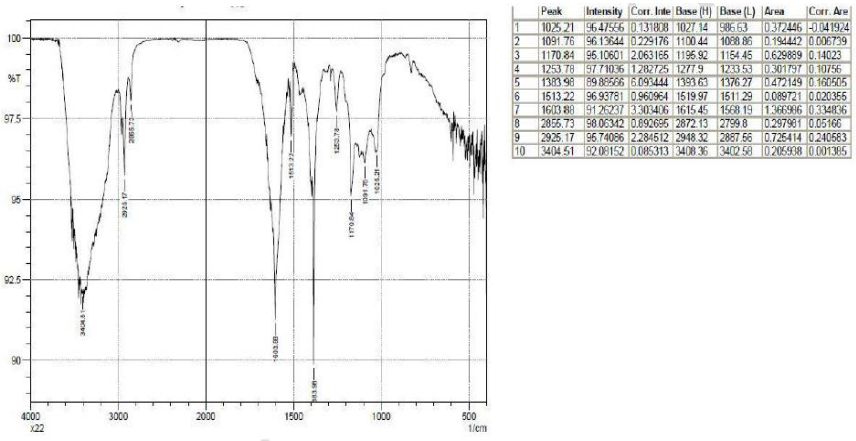


Fig. No. 3: Liquid Chromatography/ Mass Spectrometry (LC/MS) of isolated new phytosterol (HIIP) (Rf: 0.42).



New Phytosterol Rf 0.42

Fig. No. 4: UV Spectra of a new phytosterol (HIIP) at 254 nm, isolated from petroleum ether extract of leaves of *Holoptelea integrifolia* using preparative TLC.



	Peak	Intensity	Corr. Inte	Base (H)	Base (L)	Area	Corr. Are
1	1025.21	96.47556	0.131808	1027.14	986.53	0.372446	-0.041924
2	1091.76	96.13644	0.229176	1100.44	1068.86	0.194442	0.006739
3	1170.84	95.10601	2.063166	1195.92	1154.45	0.629889	0.14023
4	1253.78	97.71036	1.282725	1277.9	1233.53	0.301797	0.10756
5	1383.98	89.89596	6.093444	1393.63	1376.27	0.472149	0.160505
6	1513.22	96.93781	0.960964	1519.97	1511.29	0.089721	0.020355
7	1603.88	91.26237	3.303406	1615.45	1568.15	1.366986	0.334836
8	2855.73	98.06342	0.892695	2872.13	2799.8	0.297981	0.05166
9	2925.17	95.74086	2.284512	2948.32	2887.66	0.726414	0.240663
10	3404.51	92.06152	0.085313	3408.36	3402.58	0.205938	0.001365

Fig. No. 5: IR Spectrum of isolated new phytosterol (HIIP) (Rf : 0.42).

3.2 Assessment of Antidepressant Activity of HIIP

Effect of HIIP on TST induced depression in mice: The above mentioned doses were administered as mentioned earlier. It was observed that both low and high doses of HIIP exhibited a significant antidepressant effect by showing significant delay in latency to rearing with forelimb clonus when compared to control group. HIIP-30 mg/kg was more potent than HIIP-10 mg/kg for showing antidepressant activity. It was observed that the antidepressant activity of HIIP was dose dependent. The standard drug *fluoxetine* (10 mg/kg-i.p.) exhibited a significant antidepressant activity. The observations are given in Table 2.

Table. 3: Effect of HIIP on TST induced depression in mice.

Treatment	Immobility (second)
Vehicle control-10 ml/kg/p.o/b.wt for 14 days	83.56 ± 2.38
HIIP-10 mg/kg/p.o/b.wt for 14 days	75.39 ±1.66*
HIIP-30 mg/kg/p.o/b.wt for 14 days	68.41±1.744**
Fluoxetine-10 mg/kg/p.o/b.wt for 14 days	71.04 ± 1.961**

Values are mean ± SEM; n=6; One way analysis of variance (ANOVA) followed by Dunnetts test. Where, *P<0.05, **P<0.01,***P<0.001 HIIP: *Holoptelea integrifolia* isolated phytosterol The tail suspension test has been described by Steru et al.[12] As a facile means of evaluating potential anti depressants. The immobility displayed by rodents when subjected to an unavoidable and inescapable stress has been hypothesized to reflect behavioral despair, which in turn may reflect depressive disorders in humans. Clinically effective antidepressants reduce the immobility that mice display after active and unsuccessful attempts to escape when suspended by tail. Also recent studies have shown that the dopaminergic activation is also involved in struggling (climbing) behaviour.[13,14,15] On observation and reference to reported data from Phytochemical tests, it was clear that, HIIP, isolated from Petroleum ether extract of *Holoptelea integrifolia* (Roxb) Planch leaves showed the presence of flavonoids, steroids, triterpenoids. Flavonoids, sterols and terpenoids in preliminary phytochemical screening, these phytochemicals have been implicated in various pharmacological actions on central nervous system including antidepressant and anxiolytic activity.[14,15]

4. CONCLUSION

Majority of scientific documentation suggested prominent role of phytosterols towards Antidepressant activity.[14,15] From the above data it is concluded that the isolated phytosterol compound, HIIP. possesses significant antidepressant activity against Tail Suspension Test (TST) induced depression in mice.

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