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SCREENING OF ISOLATED PHYTOSTEROL FROM LEAVES OF *HOLOPTELEA INTEGRIFOLIA* (ROXB) PLANCH FOR ANTIOXIDANT ACTIVITY

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ABSTRACT

Holoptelea integrifolia (Roxb.) Planch has been used from long time in traditional medicine. The main objective of the work was to evaluate the antioxidant activity of *Holoptelea integrifolia* isolated Phytosterol (HIIP). The antioxidant activity of HIIP isolated from petroleum ether extract of leaves was evaluated using Total Reduction Capability. Preliminary Phytochemical investigation of the petroleum ether extract (PEHI) of *Holoptelea integrifolia* leaves reveals the presence of steroids, terpenoids, alkaloids, glycosides, flavonoids, proteins, tannins, and carbohydrate. In case of Total Reduction Capability HIIP showed dose dependent increase in absorbance thereby dose dependant total Reduction capacity indicating antioxidant activity. The results confirms that HIIP from petroleum ether extract also shows antioxidant activity using Total Reduction Capability, which suggests the application of the plant as an antioxidant agent.

KEYWORDS: Holoptelea integrifoila isolated phytosterol,

Antioxidant, Total Reduction Capability.

INTRODUCTION

Oxygen is essential for the survival of all on this earth. During the process of oxygen utilization in normal physiological and metabolic processes approximately 5% of oxygen gets

univalently reduced to oxygen derived free radicals^[1,2] like superoxide, hydrogen peroxide, hydroxyl and nitric oxide radicals. All these radicals known as reactive oxygen species (ROS) exert oxidative stress towards the cells of human body rendering t each cell to face about 10000 oxidative hits per second.^[3] When generation of ROS overtakes the antioxidant defense of the cells, the free radicals start attacking the cell proteins, lipids and carbohydrates^[4-6] and this leads to a number of physiological disorders. Free radicals are involved in the development of degerative diseases.^[6] They have also been implicated in the pathogenesis of diabetes, liver damage, nephrotoxicity, inflammation, cancer, cardiovascular disorders, neurological disordes and in the process of ageing.^[7] Many plants often contain substantial amounts of antioxidants including vitamin C and E, carotenoids, flavonoids and tannins etc. and thus can be utilized to scavenge the excess free radicals from the body.^[8]

In traditional system of medicine, bark and leaves of *Holoptelea integrifoila* (HI)used as bitter, astringent, acrid, thermogenic, antiinflammatory, digestive, carminative, laxative, anthelmintic, depurative, repulsive, urinary astringent and in rheumatism.^[9,10] The plant *Holoptelea integrifolia* is used traditionally for the treatment of inflammation, gastritis, dyspepsia, colic, intestinal worms, vomiting, wound healing, leprosy, diabetes, hemorrhoids, dysmenorrhea and rheumatism.^[11] In our previous studies the antioxidant activity of petroleum ether and methanol extract of leaf of *Holoptelea integrifolia* was evaluated and it was found that petroleum ether extract (PEHI) has more significant antioxidant activity than methanolic extract (MHI)^[12] in the similar lines in this present study a phytosterol (HIIP) was isolated from petroleum ether extract and it was studied for antioxidant activity. Hence in the current dissertation the antioxidant activity of Isolated phytosterol from petroleum ether extract of leaf of *Holoptelea* for methanolic extract of leaf of *Holoptelea* for antioxidant activity. Hence in the current dissertation the antioxidant activity of Isolated phytosterol from petroleum ether extract of leaf of *Holoptelea* phytosterol from petroleum ether

MATERIALS AND METHODS

Plant Introduction

Holoptelea integrifoila belongs to the family ulmaceae commonly called as Indian Elm and frequently used in India by the tribal people for it's medicinal properties. The mucilaginous bark is boiled and the juice squeezed out and applied to rheumatic swellings.^[13] Leaves of *Holoptelea integrifoila* were collected in the Month of August from the agricultural fields of Tirunelveli district, Tamil Nadu, India. The plant was identified and leaves of *Holoptelea integrifolia* were authenticated and confirmed from Dr.V.Chelladurai, Research Officer, Botany, C.C.R.A.S. (Retired), Govt. of India by compairing morphological features (leaf and

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stem arrangement, flower /inflorescence arrangement, fruit and seed morphology etc.). The collected plant material was shade dried to retain its vital phytoconstituents and then subjected to size reduction for further extraction process.



Fig. 1: Habitat of Holoptelea integrifolia (Roxb.) Planch.

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.

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

The dried and powdered leaves (1kg) of *Holoptelea integrifolia* was extracted with petroleum ether (b.p. 60-80°C) 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 μ m thickness with CAMAG Linomat V (Switzerland) was used. The plates were washed by methanol and

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activated at 120° 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 composition 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 110 °C. 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.^[14-22]

Drugs

Phosphate buffer, Hydrogen peroxide and Vitamine- C, All the chemicals used in the study were of analytical grade and procured from Merck India Pvt. Ltd.

The Petroleum ether extracts of *Holoptelea integrifolia* leaves were subjected to the following investigations,

- 1. Preliminary photochemical screening
- 2. Antioxidant activity

Preliminary phytochemical screening of petroleum ether extract

The extract was subjected to follow chemical tests to detect the phytochemical constituents present in it. 0.5 gm of extract was dissolved in 5 ml of distilled water and filtered. The filtrate was used to determine the presence of various phytoconstituents.^[23]

Assessment of Antioxidant Activity

Total reduction capability

Total reduction capability of HIIP ($20\mu g/ml$, $40\mu g/ml$, $60\mu g/ml$, $80\mu g/ml$ and $100\mu g/ml$) and Vit C ($20\mu g/ml$, $40\mu g/ml$, $60\mu g/ml$, $80\mu g/ml$ and $100\mu g/ml$) were estimated by using the method of Oyaizu. Different concentrations of HI extracts and Vit C were mixed with phosphate buffer (2.5 ml, 0.2 M, pH 6.6) and potassium ferricyanide (1%, 2.5 ml). The mixture was incubated at 50° C for 20 min. A portion of (2.5 ml) trichloroacetic acid (10 %) was added to the mixture. Then it was centrifuged for 10 min at 1000 g. The upper layer of the solution (2.5 ml) was mixed with distilled water (2.5 ml) and ferric chloride (0.5 ml, 0.1%) and the absorbance was measured at 700 nm by using spectrophotometer. Higher absorbance of the reaction mixture indicated greater reducing power.^[24,25]

RESULTS

Total Reduction capability of HIIP

HIIP has shown dose dependent increase in absorbance thereby dose dependant total reduction capacity indicating antioxidant activity. The observations are given in table no.1

Concentration	HIIP	Vitamin-C
20 µg/ml	0.5654	0.7143
40 µg/ml	0.6412	0.7710
60 µg/ml	0.7512	0.8440
80 μg/ml	0.9214	0.9122
100 µg/ml	1.0433	1.1023

Table No. 1: Total Reduction capability of HIIP At Different Concentrations.

Where HIIP: *Holoptelea integrifolia* Isolated Phytosterol, Vitamin- C was used as the positive control.

DISCUSSION

Free radicals are chemical entities that can exist separately with one or more unpaired electrons. The propagation of free radicals can bring about thousands of reactions and thus may cause extensive tissue damage. Lipids, proteins, and DNA are all susceptible to attack by free radicals.^[4,5] Antioxidants may offer resistance against oxidative stress by scavenging the free radicals, inhibiting lipid peroxidation etc.

In the present investigation, preliminary phytochemical analysis and the earlier scientific studies have shown that petroleum ether extract of *Holoptelea integrifolia* leaves showed the prominent presence of steroids, triterpenoids, glycosides, saponins, flavonoids, proteins,

tannins and carbohydrates. The previous scientific studies have shown that these secondary plant metabolites are mainly responsible for the pharmacological actions and thus thereby it supported the traditional uses.^[26,27] Which may also be responsible for the various actions of *Holoptelea integrifolia*. This gives a green signal towards further exploration of this plant for the validation of traditional claims for various complaints for which there is either no or very limited satisfactory pharmacotherapy.

As the plant has shown its potential effectiveness in treating various disorders for which most common mechanism may be through its antioxidant potential. Also, *Holoptelea integrifolia* has been traditionally claimed to possess antioxidant properties. So in order to assess its efficacy as a potent antioxidant agent the plant was investigated using in vitro model namely Total Reduction Capability.

Total reducing ability is considered as the ability of Fe3+–Fe2+ transformation. The reducing capacity of a compound may serve as a significant indicator of its potential antioxidant activity. The HI extracts showed dose dependent total reduction capacity by dose dependent increase in absorbance indicating total reducing ability in turn potential antioxidant activity. These results indicate its usefulness in various disorders associated with oxidative stress.^[28,29]

CONCLUSION

Majority of scientific documentation suggested prominent role of phytosterols towards anti oxidant activity. Hence, HIIP from petroleum ether extract of leaves of HI have naturally exhibited significant antioxidant activity using Total Reduction Capability, which support the ethnomedicinal application of the plant as an antioxidantagent, also our studies confirm the application of *Holoptelea integrifolia* leaves as an antioxidant.

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