

Quantitative Screening of Phytochemicals of Different Parts of *Ficus benghalensis* Linn.

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Abstract

The present study was carried out to investigate the quantitative and phytochemical screening of different part of *Ficus benghalensis* Linn using standard technique. The Methanolic extract of different part of *ficus benghalensis* were analyzed quantitatively by spectrophotometric method for the phytochemicals: flavonoids, total phenol and flavonols. Phytochemical screening of the plant showed the presence of flavonoid, cardiac glycosides. The quantitative screening of phytoconstituents contained in the different plant extract of the *ficus benghalensis* revealed that flavonol are high 363 µg/ml, flavonoid is relatively moderate 129.408 µg/ml and phenol content is 5.359 µg/ml. This study vindicates the curative usage of *ficus benghalensis* in traditional medicine.

Keyword: *Ficus Benghalensis* Linn., Antioxidant, Phytochemical, ascorbic acid

1. Introduction

Plants are utilized extensively as raw drugs for many formulations in traditional systems of medicine^{1,2}. Medicinal plants have curative properties due to the presence of various complex chemical substances of different composition, which are found as secondary plant metabolites in one or more parts of these plants^{2,3}. The plants have the ability to produce large variety of secondary metabolites such as terpenoids,

alkaloids, flavonoids and phenyl propanoids, which together account for 200,000 compounds⁴. Phytochemical are bioactive compound found in plant that work with nutrient and dietary fibers which protect against various diseases⁵.

The most widely occurring group of Phytochemical is flavonoids and phenolic compound present in plant. Biological activity of these groups of Phytochemical was reported in several studies⁶. Phenolic compounds have gained much attention, due to their antioxidant activities and free radical-scavenging abilities, which potentially have beneficial implications for human health. Flavonoids are regarded as one of the most widespread groups of natural constituents found in plants. Polyphenols and flavonoids are used for the cure and prevention of various diseases⁷.

Ficus benghalensis also known as Indian banyan tree⁸ and its belong to the family Moraceae. The imposing banyan tree of poetry and legend is a store-house of invaluable remedies for some of the deadliest diseases⁹. *Ficus benghalensis* which is an indigenous plant possessing reputed medicinal properties, have been listed in Ayurvedic literature^{2,10}. Various scientific studies have been carried out on *Ficus benghalensis* and various pharmacological activities have been reported. It has been reported to possess immunomodulatory, hypoglycemic, antioxidant, antistress and antiallergic, anthelmintic activities¹¹. The main

focus is given to the study of phytochemical screening and the quantitative activity of these plants.

2. Materials And Methods

PLANT MATERIALS AND EXTRACTION:

The different plant parts (leaves, bark, fruit, and stem) of *ficus benghalensis* were collected from University campus of MGCGV, Chitrakoot, Satna (M.P. India) washed with water and then washed with methanol and allowed to dry in shade at room temperature. Dried parts of plants grind and passed through 120 no sieve. Take 20gm of sieved powder of *F. benghalensis* then added 100ml (80%) methanol and using cold maceration method for extraction. Extract was filtered through the whatman No.1 filter paper. The filtrate was concentrated under room temperature for dryness.

2.1 Preliminary Phytochemicals Screening

Phytochemical screening performed were using by given standard procedure.

Flavonoids

In 1 ml methanolic extract added 1ml dilute ammonia in a test tube, yellow color appeared then added few drop of conc. H_2SO_4 , yellow color was disappear. Indicate presence of Flavonoids.

Tannins

1 ml of the methanolic extract and 2 ml of water was added in a test tube. Then 2 to 3 drops of diluted ferric chloride solution was added and observed for green to blue-green (catechic tannins) or a blue-black (gallic tannins) coloration.

Carbohydrate

In 2ml of methanolic extract and 1ml molish reagent was added in a test tube. Then added 2 to 3 drop of conc. H_2SO_4 at the side wall of test tube and observed purple and violet color ring at the junction of two liquid.

Reducing Sugar

The methanol extract (1 ml) and 2 drops of boiling Fehling's solution (A and B) was added in a test tube. The formation of a precipitate red-brick in the bottom of the tube indicates the presence of reducing sugars.

Starch

The aqueous extract 5ml was treated with the reagent of the starch (iodine) blue violet color obtain indicates the presence of starch.

Protein

2 ml Methanolic extract of was added 1drop 1% NaOH and 2 drop 1% $CuSO_4$ in a test tube. Blue or purple color is obtained indicates the presence of protein.

Saponins

1 ml of methanolic extract was added and few volume of distilled water in a test tube. The solution was shaken vigorously and observed for a

stable persistent froth for 20 min, indicates the presence of saponin.

Terpenoids

In 2 ml of methanolic extract was added 1 ml chloroform shake properly and added H_2SO_4 . Raddish brown ring in obtained. Indicate the presence of terpenoids.

Steroids

2 ml of methanolic extract few amount of chloroform was added and shake properly and added 3 to 4 drop of acetic anhydride and 1drop H_2SO_4 in a test tube. Formation of purple color change in to green. Indicate the presence of steroids.

Glycosides

2 ml of methanolic extract, 0.5 ml glacial acetic acid shake properly and added 3 to 4 drop of $FeCl_3$ in a test tube. Formation of brown color ring at the interface of test tube. Indicate the presence of cardiac glycosides.

Phenol

2 ml of methanolic extract was added 2 ml 5% aqueous $FeCl_3$ in a test tube. Formation of blue color obtained. Indicate the presence phenol.

Alkaloids

2 ml of methanolic extract was added 2 ml Mayer reagent in a test tube. Form white ppt. Indicate the presence alkaloids.

2.2 Quantitative Analysis

Total polyphenolic content

Total polyphenolic content of different part of plant extract was measured by using Folin - Ciocalteu reagent. The 25 μ l of plant extract diluted with 125 μ l water followed by addition of 150 μ l of Folin- ciocalteu (1N) & 25 μ l of Na_2CO_3 (20% W/V) and incubated at 45 $^{\circ}C$ for 60 min then absorbance was measured by spectrophotometrically at 765nm (Bio Tek Synergy H4 multimode micro plate reader Bio Tek instrument, Inc Winooski, VT, USA). Absorbance was recorded triplicates. Quantification was performed with respect to the standard curve of quercetin (equation). Result was expressed as milligram of quercetin equivalent per ml of extract¹⁴.

Total flavonoid content

Total flavonoid in the plant extracts, in brief, 100 μ l of sample, followed by 100 μ l of $AlCl_3 \cdot 6H_2O$ in ethanol and 150 μ l Sodium acetate, solution added. The absorbance at 430nm was taken (BioTeksynergyH4 multi-mode microplate reader, Bio Tek Instruments, Inc Winooski, VT, USA), after 2.5 h of incubation at 200 C. Total flavonoid contents were calculated with respect to the standard curve of the flavonoid quercetin dehydrate (equation). Results were expressed as micrograms of quercetin dehydrate equivalents (QE) per ml of the extract¹⁴.

Flavonol

Flavonol content in the sample (100 times diluted with methanol) was measured by mixing equal volume of plant extract with 2% $AlCl_3 \cdot 6H_2O$ in a 96 well plate. Absorbance was recorded at 420 nm spectrophotometrically (Bio Tek synergyH4 multimode microplate reader, BioTek Instruments, Inc Winooski, VT, USA). Flavonol contents in the extracts were determined with respect to the standard curve of the flavonoid quercetin (equation). Results were expressed as micrograms of quercetin equivalents (QE) per ml of the extract.

3. Results And Discussion

The result of screening of Phytochemical from different part of *Ficus bhenghalesi* Linn showed the presence of medicinal active constituent like Flavonoid, Tannin, saponin, terpenoids, steroids and glycosides were given in **Table-1**.

Phytochemical analysis conducted on the plant extracts revealed the presence of constituent which are known to exhibit medicinal as well as physiological activity. Analysis of plant extract revealed the presence of Phytochemical such as alkaloid, flavonoid, tannin, saponin, glycosides, terpenoids, steroids have been associated with medicinal properties. Phytochemical are the basic source for the establishment for several Pharmaceutical industries. The constituent present in the plants play a significance role in the identification of crude drugs. Phytochemical screening is very important in identifying the new

source of therapeutically and industrial important compounds like alkaloids, flavonoids, terpenoids, phenolic compound, saponin, steroids etc.

The quantitative analysis of *Ficus bhenghalesis* Linn. total phenolic content, total flavonoid and flavonol were carried out using multimode micro plate reader. The total phenolic content in plant extract is expressed in term of Catechol equivalent **Fig. 1** (The equation of standard curve is $Y = 0.0139 X + 0.085$ $R^2 = 0.9859$). The concentration of total phenolic content was estimated 5.395 in leaf, 0.359 in stem, 3.093 in bark and 2.589 $\mu g/ml$ in fruit were given in **Table-2**. Phenolic compound occur ubiquitously in plants and a variety of biological activities have been attributed to them.

The total flavonoid content in plant extract is expressed in term of quercetin equivalent **Fig. 2** (The equation of standard curve is $Y = 0.0049 X + 0.0949$ $R^2 = 0.9969$). The concentration of total flavonoid content in plant extract was estimated 55.326 in leaf, 129.408 in stem, 134.91 in bark, and 99.6122 in fruit were given in **Table-3**.

The flavonol content in plant extract is expressed in term of quercetin equivalent **Fig. 3** (The equation of standard curve is $Y = 0.0024 + 0.0635 X$ $R^2 = 0.9855$). The concentration of flavonol content in plant extract was estimated 123.5417 in leaf, 363.125 in stem, 268.9583 in bark and 111.875 in fruit were given in **Table-4**.

Table 1: Phytochemical screening of different part of ficus benghalensis

Phytochemicals	Test	Leaf	Stem	Bark	Fruit
Flavonoid	-	+	+	+	+
Tannin	-	+	+	+	+
Carbohydrate	Molish	+	+	+	+
	Fehling	+	+	+	+
Protein	Biuret	+	-	-	+
Saponin	Foam	+	+	+	+
Terpenoids	-	+	+	+	+
Steroids	-	+	+	+	+
Glycoside	-	+	+	+	+
Phenol	-	+	+	+	+
Alkaloids	Meyer	-	-	-	-
Starch	Iodine	+	+	+	+

Table 2: Total Polyphenol content of different part of ficus benghalensis

S.NO.	Plant name	Botanical name	Total Polyphenol
1.	Bargad leaf	<i>Ficus benghalensis</i>	5.395 µg/ml
2.	Bargad bark	<i>Ficus benghalensis</i>	3.093 µg/ml
3.	Bargad stem	<i>Ficus benghalensis</i>	0.359 µg/ml
4.	Bargad Fruit	<i>Ficus benghalensis</i>	2.589 µg/ml

Table 3: Flavonoid content of different part of ficus benghalensis

S.NO.	Plant name	Botanical name	Total Flavonoid
1.	Bargad leaf	<i>Ficus benghalensis</i>	55.326 µg/ml
2.	Bargad bark	<i>Ficus benghalensis</i>	139.91 µg/ml
3.	Bargad stem	<i>Ficus benghalensis</i>	129.408 µg/ml
4.	Bargad Fruit	<i>Ficus benghalensis</i>	99.6122 µg/ml

Table 4: Flavanol content of different part of ficus benghalensis

S.NO.	Plant name	Botanical name	Total Flavanol
1.	Bargad leaf	<i>Ficus benghalensis</i>	123.54 µg/ml
2.	Bargad bark	<i>Ficus benghalensis</i>	268.95 µg/ml
3.	Bargad stem	<i>Ficus benghalensis</i>	363.12 µg/ml
4.	Bargad Fruit	<i>Ficus benghalensis</i>	111.87 µg/ml

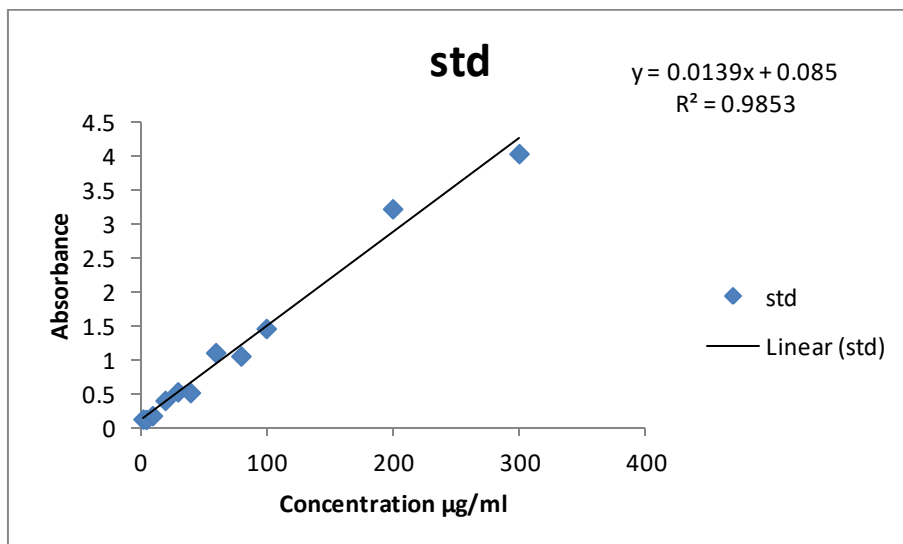


Fig. 1: Standard graph of catechol for total phenol

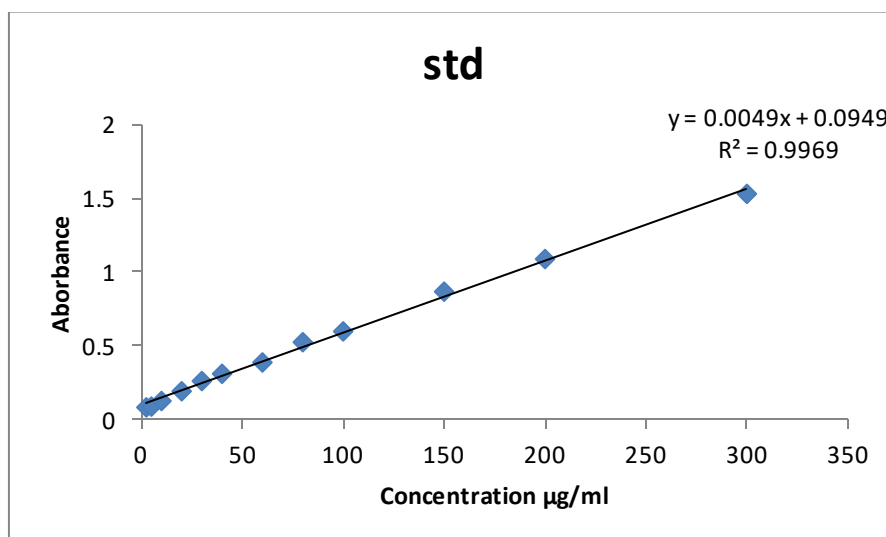


Fig. 2: Standard graph of quercetin for flavonoid

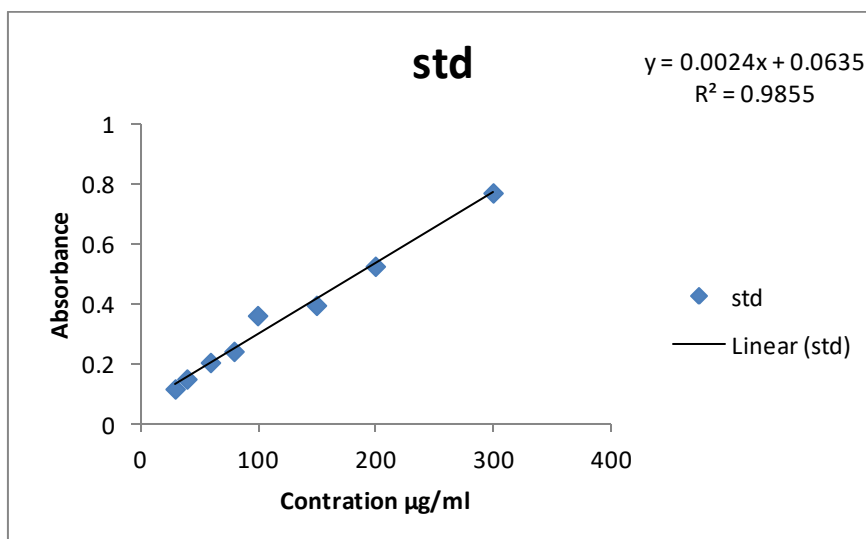


Fig. 3: Standarded graph of quercetin for Flavonol

4. Conclusion

The present research work concludes that *Ficus benghalensis* is important medicinal plants and contains various active phyto constituents. The overall result obtain by present study we observed that the total flavonoid content was highest in *Ficus benghalensis* leaf and lowest in stem. Highest flavonol content was found in *F. benghalensis* stem, lowest in fruit and total phenolic content was highest in leaf and lowest in stem.

Reference

1. Babu Koilpillai, Shanker Sabesan Gokul, Rai Sadananda, Comparative Pharmacognostic Studies on the barks of four *Ficus* species, Turk J Boat, 34, 215-224, 2012.
2. Pandeya Krishna Bihari, Tripathi Indra Prasad, Mishra Mahendra Kumar, Dwivedi Neelesh, Pardhi Yogesh, Kamal Arti, Gupta Priyanka, Dwivedi Nupa, Mishra Chinmayi, A Critical Review on Traditional Herbal Drugs: An Emerging Alternative Drug for Diabetes, International Journal of Organic Chemistry, 3, 1-22, 2013.
3. Singh Shailja, Jaiswal Shalini, Therapeutic Properties of *Ficus Religiosa* International Journal of Engineering Research and General Science, 2, 2014
4. Agrawal Supriya, Katare Charu, Prasad Gbks, Antioxidant activity, total Phenolic compound and Flavonoid content of vaccum dried extract of *L. Siceraria*, Global Journal of pharmaceutical research, 4, 302-308, 2015.
5. Taskeen Abida, Naeem Ismat, Mubeen Hifsa and Mehmood Talib, Reverse Phase High Performance Liquid Chromatographic analysis of flavonoids in two *Ficus* species, New York Science Journal, 2, 32-35, 2009.
6. Shaikh Abusufyan, Ibrahim Mohammed, Mohib Khan, Comparative in vitro Antidiabetic and Antioxidant Activity of Various Extracts of Ficus Species, A Multifaceted Journal in the field of Natural Products and Pharmacognosy, 10, 349-354, 2018.
7. Chandrasekar S. B., Bhanumathy M., Pawar A. T., 1 and Somasundaram T., Phytopharmacology of *Ficus religiosa*, Pharmacogn Rev.4, 195-199, 2010.
8. Mandal SG, Shete RV, Kore KJ, Otari KV, Kale BN and Manna AK, Review: Indian national tree (*Ficus bengalensis*), International Journal of Pharmacy and Life sciences, 1, 268-273, 2010.
9. Deraniyagala, S.A., Wijesundera, R.L.C. and Weerasena, O.V.D.S.J. Antifungal activity of *Ficus racemosa* leaf extract and isolation of active compound, Journal of the National Science Foundation of Sri Lanka, volume 26, 19-26, 1998.
10. Deshmukh V.K., Shrotri D.S. and Aiman R., Isolation of a Hypoglycemic principle from the bark of *Ficus bengalensis*, Ind., J. Physiol. and Pharmacol. 4, 182-185, 1960.

11. Garg Vipin Kumar, Paliwal Sarvesh Kumar, Wound-healing activity of ethanolic and aqueous extracts of *Ficus benghalens*, J Adv Pharm Technol Res.. 6, 110-114, 2011.
12. Zohar S. F., Meriem B., Samira S., A. M. M. S., Phytochemical Screening and identification of some compounds from Mallow, Journal of Natural Product and Plant Resources, 2, 512-516, 2012
13. Ayoola G.A., Coker H., Adesegun S A, Adepoju-Bello AA, Obaweya K, Ezennia EC, Atangbalyia TO, Phytochemical Screening and Antioxidant Activities of Some Selected Medicinal Plants used for Malaria Therapy in Southwestern Nigeria Journal of Pharmaceutical Research, 7, 1019-1024, 2008.
14. I.P. Tripathi, M. K. Mishra, C. Mishra, R. Tripathi, A. Kamal, P. Tripathi. V.P. Shukla, R. Gangele and K.B. Pandey, Assessment of Antioxidant and Total Polyphenolic content of some plants of Euphorbiaceae Family, Indian Journal of Applied Research, volume 3, 1-4, 2013.