



Phytochemical Profiling and Biological Activities of *Strobilanthes urens* (B. Heyne ex Roth) J.R.I. Wood. Leaf Extract (Acanthaceae)

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Abstract: This study investigated the phytochemical composition and biological activities of *Strobilanthes urens* leaf extracts. The proximate composition, including protein, carbohydrate, lipid content, nutritive value, and total moisture, was evaluated. Qualitative phytochemical screening revealed the presence of alkaloids, flavonoids, proteins, tannins, phenols, terpenoids, saponins, glycosides, and carbohydrates in all three extracts, except for steroids in methanol, and both terpenoids and steroids in the distilled water extract. Alkaloids, tannins, saponins, glycosides, and steroids were absent in the chloroform extract. Quantitative analysis confirmed the presence of polyphenols, flavonoids, tannins, and alkaloids, while GC-MS analysis identified 22 major bioactive compounds with potential medicinal properties. Antioxidant activity assessed by the DPPH assay showed IC₅₀ values of 79.23 ± 3.78 µg/mL in methanol and 237.00 ± 12.37 µg/mL in distilled water, with no activity in chloroform. The extracts exhibited antibacterial activity but no significant antifungal effect. Cytotoxicity evaluation by MTT assay showed IC₅₀ values against MCF-7 and HepG2 cell lines ranging from 88.19 ± 8.29 to 188.10 ± 39.99 µg/mL. Overall, the findings highlight the phytochemical richness and potential pharmacological value of *S. urens* leaf extracts.

Introduction

All over the world, plants are essential to human life. Their secondary metabolites play a vital role. A large number of plants are used to treat numerous ailments. According to the WHO, eighty percent of the population in the world depends upon medicine, 80 to 85% of medicine is extracted from medicinal plant extracts (1). Herbal medicines treat and cure many diseases and health conditions through traditional systems such as Unani, Ayurveda, and Siddha. *Strobilanthes urens* (B. Heyne ex Roth) J.R.I. Wood is a genus of flowering plants in the family Acanthaceae. The native range of this species is western India. It is a subshrub and grows primarily in the seasonally dry tropical biome. Acanthaceae comprises 346 genera and 4300 species, most of which are herbs, shrubs, and vines. The plants of this family are cosmopolitan and distributed in the Old and New World. And present in Africa, Central America, Malaysia, and Indonesia, with few species extending to South Europe, Japan, the Southern coast of New Holland, and Southern to the Cape of Good Hopes. In India, Acanthaceae genera are peculiar to the Southern parts, Indian Archipelago and Malayan Peninsula but have spread from Sutlej to Sylhet and lower ranges of Himalayas (2). The plants have many therapeutic uses, such as treating inflammation, infections, wounds, skin lesions, diarrhea, scabies, leprosy, ulcers, and snake bites (3).

Plant-based medicine has become of great interest owing to its versatile applications. Medicinal plants are the source of drugs in conventional systems of medicine, including present-day medications, food supplements, nutraceuticals, folk medicines, pharmaceuticals, and synthetic drugs (4). The plant has been traditionally used to treat bone fractures, skin conditions, urinary diseases, and allergies (5). *Strobilanthes hamiltoniana* exhibits potent pharmacological activities, including antibacterial and anthelmintic activities (6). The present research aims to study the potent phytochemicals (Qualitative and Quantitative), their antimicrobial, antioxidant, and anticancer activities.

Materials and Methods

Collection of Plant Material

The collection of *S. urens* (B. Heyne ex Roth) J.R.I. Wood is from Kalgi village, Kalburgi District, Karnataka, India, identified by A N Sringswara, curator, Mahatma Gandhi Botanical Garden, University of Agricultural Sciences, KGVK Bangalore, collection Number 009, Accession number UASB 5720. The plant and its part identified can be seen in **Figure 1**.

Preparation of Leaf Extract

The fresh plant leaves were washed with ionized water to

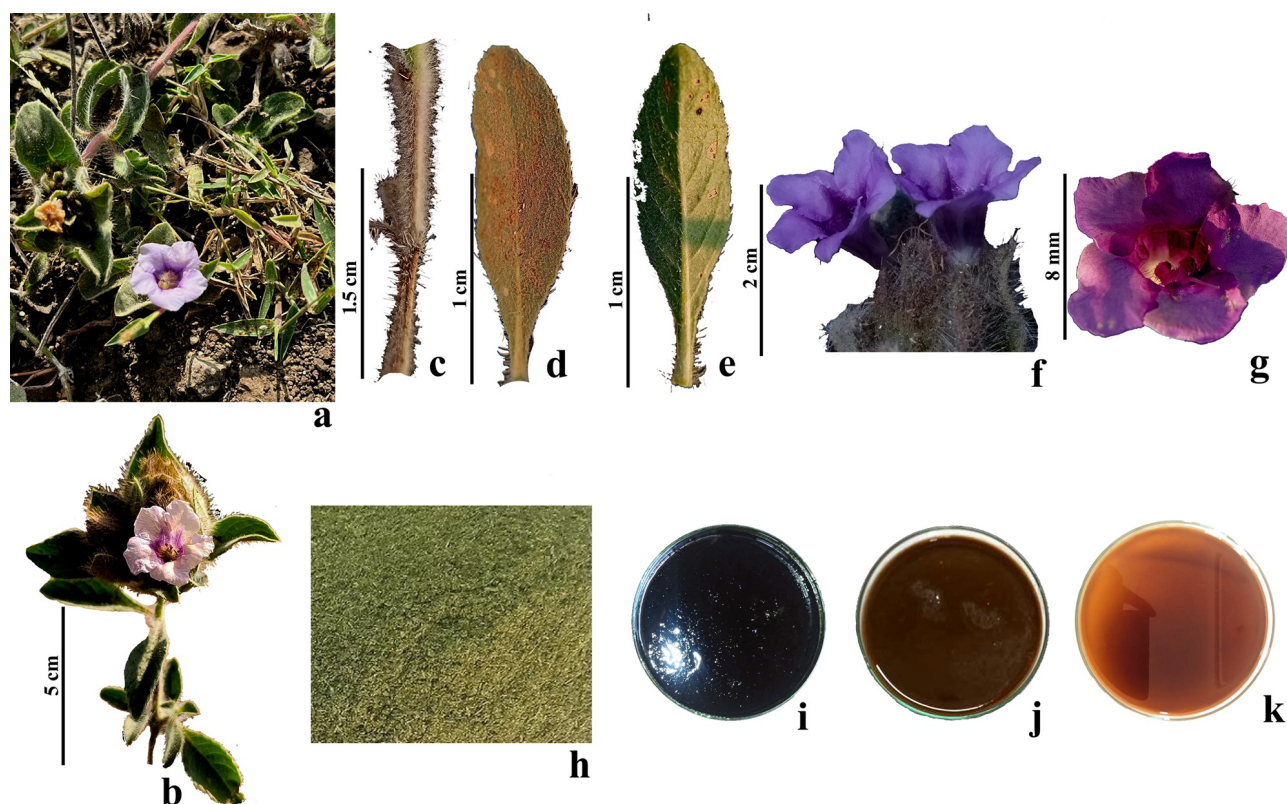


Figure 1. (A) Habit, (B) flowering twig, (C) stem, (D) adaxial leaf, (E) abaxial leaf, (F) inflorescence, (G) flower, (H) dry powder, (I) methanol extract, (J) water extract, and (K) chloroform extract of *Strobilanthes urens*.

remove surface impurities and unwanted debris, then air-dried in a shaded area to ensure proper moisture removal. The dried material was ground into a fine powder using a blender. Approximately 25 g of the leaf powder were packed into filter paper and subjected to Soxhlet extraction using 250 mL each of distilled water, methanol, and chloroform.

Proximate Compositions

The dried plant powder was used to analyze moisture content, which was determined by drying the biomass for 10 h at 100 °C. Lipid content was estimated using the phenol-sulfuric acid method, carbohydrate content by the Kjeldahl method, and protein content according to AOAC procedures. Ash content was measured in a muffle furnace at 550 °C to determine the nutritive value (7).

Total Phenolic Contents

The total phenolic content (TPC) was estimated using the Folin-Ciocalteu (FC) reagent, with gallic acid as the standard antioxidant. A 100 µL aliquot of appropriately diluted extract was mixed with 0.5 mL of FC reagent and incubated at room temperature for 10 min. Then, 2 mL of 7% Na₂CO₃ solution was added. The mixture was boiled for 1 min, after which the absorbance was measured at 750 nm using a spectrophotometer (Shimadzu UV-1800, Kyoto, Japan). The results were expressed as micrograms of gallic acid equivalents (µg GAE) (8).

Total Flavonoid Estimation

The total flavonoid content was estimated using 200 µL of the sample mixed with 5 mL of chromogen reagent, 25 mL of methanol, 25 mL of concentrated HCl, and 0.1%

cinnamaldehyde solution in a cooled mixture totaling 75 mL. After incubation for 10 min, the absorbance was measured at 640 nm. The flavonoid content was expressed as micrograms of catechin equivalents (µg CE) per milligram of extract (9).

Total Alkaloid Estimation

A 200 µg portion of the sample was dissolved in 1 mL of 2N HCl, then filtered and washed with 10 mL of chloroform. Similarly, atropine standard solutions (20, 40, 60, 80, and 100 µg/mL; Sigma Chemical, USA) were prepared. To each solution, 5 mL of bromocresol green (BCG) reagent (prepared by dissolving 69.8 mg of BCG in 3 mL of 2N NaOH and diluting with distilled water) and phosphate buffer (pH 4.7) were added, and the total volume was made up to 10 mL. The absorbance was measured at 470 nm using a spectrophotometer, with a blank prepared under the same conditions but without atropine or sample. The alkaloid content was expressed as micrograms of atropine equivalents (µg AE) per milligram of extract.

Total Tannins Estimations

The tannin content was estimated using 0.05 mL of the sample, and the volume was adjusted to 0.5 mL with distilled water. Then, 0.25 mL of 1N Folin-Ciocalteu reagent was added, followed by 1.25 mL of 20% sodium carbonate solution. The mixture was vortexed and incubated for 40 min at room temperature. Absorbance was measured at 725 nm using a spectrophotometer. The tannin content was expressed as micrograms of tannic acid equivalents (µg TAE) per milligram of extract (10).

Qualitative Phytochemical Analysis

A preliminary qualitative phytochemical analysis was conducted for different secondary metabolites (Phenols, Alkaloids, Flavonoids, Tannins, Terpenoids, Saponins, Glycosides, Steroids, Proteins, Carbohydrates) using specific chemical reagents (11).

Antimicrobial Activity

Antimicrobial activity was evaluated using the agar diffusion method (12). The samples were tested against gram-positive *Staphylococcus aureus* (MTCC-7443) and gram-negative *Escherichia coli* (MTCC-7410), with incubation at 37 °C for bacterial strains. The inoculum was adjusted to approximately 5×10^5 CFU/mL using sterile saline solution. Samples were dissolved in DMSO at a concentration of 20 mg/mL to prepare the stock solution and loaded into wells at concentrations ranging from 200 µg to 800 µg. Mueller-Hinton agar was used for bacterial cultures, and Czapek-Dox agar was used for fungal species. The fungi, *Aspergillus flavus* (MTCC-9606) and *Pichia anomala* (MTCC-237), were incubated at 28 °C for 72 h. After incubation, the diameters of the inhibition zones (mm) were measured.

GC-MS Analysis

Gas chromatography-mass spectrometry (GC-MS) analysis was performed to determine the phytochemical constituents. The procedure was carried out using a Shimadzu QP2010S system operating in electrospray ionization (ESI) mode, equipped with an ELITE-5MS capillary column (film thickness: 0.25 µm; length: 30 m; internal diameter: 0.25 mm). The GC oven temperature was initially set at 80 °C and programmed to increase to 450 °C at a rate of 20 °C/min to achieve effective analyte separation. Sample injection was conducted using a 2 mm direct injection technique. Compound identification was based on the comparison of relative retention times and mass spectral data with reference spectra from the National Institute of Standards and Technology (NIST) library. The analytical procedure followed established protocols to ensure accurate compound characterization, as described by Chakraborty et al. (2022) (13).

FTIR Analysis

Fourier Transform Infrared Spectroscopy (FTIR) was employed to analyze the bioactive constituents. The analysis was performed using a spectrophotometer (Thermo Fisher Scientific, Waltham, MA, USA). The dried powder of biosynthesized nanoparticles was mixed with potassium bromide (KBr) and compressed into pellets. The spectra were recorded within the range of 400–4000 cm⁻¹ to identify characteristic infrared absorption bands. The presence of functional biomolecules in the sample was confirmed through spectral analysis, facilitating comprehensive characterization of the nanoparticles' chemical composition (14).

DPPH Assay

The radical scavenging activity of the samples was evaluated using the stable DPPH radical, as previously described (15). Different concentrations of the samples (0–100 µg/mL) were mixed with 2 mL of DPPH solution (100 µM) and 3 mL of methanol. The mixture was incubated at room temperature in the dark for 45 min. After incubation, the absorbance was measured at 517 nm using a spectrophotometer (Shimadzu

UV-1800, Kyoto, Japan) against a blank (without sample or standard). The free radical scavenging capacity of the samples was calculated and expressed as IC₅₀ values, relative to vitamin C as the standard.

Cytotoxicity Property

The IC₅₀ value was determined using the MTT assay. Cultured cells (1×10^5) were seeded into 96-well plates and incubated for 48 h at 37 °C in a 5% CO₂ incubator. After incubation, the monolayer was washed with fresh medium, and 100 µL of different test concentrations of the samples was added to each well. The cells were further incubated under the same conditions. Subsequently, the medium was removed, and 100 µL of MTT solution was added to each well, followed by incubation at 37 °C for 4 h. After the supernatant was discarded, 100 µL of DMSO was added to each well and incubated for 10 min to solubilize the formazan crystals. The optical density was measured at 590 nm. The percentage of cell growth inhibition was calculated, and the IC₅₀ values were determined from the dose-response curve (16).

Results and Discussion

Proximate Composition

The proximate analysis of *S. urens* leaf extract revealed the following composition: total protein, $28.62 \pm 2.57\%$; total carbohydrates, $17.85 \pm 1.24\%$; total lipids, $12 \pm 0.75\%$; total ash, $14.26 \pm 0.59\%$; moisture content, $2.35 \pm 0.24\%$; and nutritive value, 233.88 kcal.

In comparison, *Strobilanthes crispus* leaves were reported to have an ash content of 21.6%, with mineral concentrations of calcium (5185 mg/100 g), potassium (10,900 mg/100 g), sodium (2953 mg/100 g), iron (255 mg/100 g), and phosphorus (201 mg/100 g) in notably high amounts (17). *Strobilanthes crispa* Blume was reported to contain relatively low levels of protein, carbohydrates, fat, moisture, ash, and minerals (18). Similarly, *Strobilanthes auriculata* Nees was found to contain protein (8.40 mg/100 g), amino acids (2.25 mg/100 g), tannins (4.80 mg/100 g), crude lipids (0.99 mg/100 g), and crude fiber (0.14 mg/100 g) (19). Furthermore, *S. crispus* was reported to have moisture ($82.43 \pm 0.94\%$), ash ($6.97 \pm 0.34\%$), carbohydrates ($2.94 \pm 0.92\%$), protein ($4.56 \pm 0.12\%$), and fat ($3.05 \pm 0.41\%$) (20).

Table 1. Qualitative phytochemical analysis of *Strobilanthes urens* leaf extract.

No.	Phytochemical tests	SULF M	SULF DW	SULF CHL
1	Phenols	+	+	+
2	Alkaloids	+	+	-
3	Flavonoids	+	+	+
4	Tannins	+	+	-
5	Terpenoids	+	-	+
6	Saponins	+	+	-
7	Glycosides	+	+	-
8	Steroids	-	-	-
9	Proteins	+	+	+
10	Carbohydrates	+	+	+

Note: (+) = present and (-) = absent.

Table 2. Quantitative phytochemical estimation of *Strobilanthes urens* leaf extract.

No.	Samples	Polyphenols (µg/mg GAE)	Flavonoids (µg/mg QE)	Tannins (µg/mg TAE)	Alkaloids (µg/mg AE)
1	SULF DW	235.50 ± 5.44	135.57 ± 8.88	28.88 ± 2.89	105.71 ± 5.20
2	SULF M	270.50 ± 3.30	260.57 ± 3.93	130.03 ± 7.02	211.79 ± 1.99
3	SULF ChI	199.25 ± 2.50	150.15 ± 5.63	2.45 ± 3.51	53.06 ± 5.85

Table 3. Bioactive compounds found in *Strobilanthes urens* using GC-MS.

Peak#	Retention Time	Area	Area%	Similarity	Base m/z	Compound Name
1	3.842	1646387	2.15	98	96.00	Furfural
2	5.687	1058709	1.38	98	110.05	Furfural <5-methyl->
3	6.844	442838	0.58	96	79.05	Benzyl alcohol
4	6.981	648737	0.85	75	113.00	Thiofuran <2-methyl-, 3-methyl->
5	7.614	395493	0.52	83	43.00	Pineapple ketone
6	9.670	2762250	3.60	94	97.00	Hydroxymethyl furfural
7	11.987	438285	0.57	66	136.05	Copaen-4-alfa-ol <beta->
8	13.821	3136242	4.09	65	73.05	Decamethylcyclopentasiloxane
9	14.530	1534616	2.00	73	73.05	Tridecanol <n->
10	14.621	1261145	1.65	53	73.05	Tridecane <n->
11	17.451	348926	0.46	96	68.05	Neophytadiene
12	17.521	355595	0.46	92	43.05	Phytone
13	18.793	4311986	5.63	46	73.05	Decamethylcyclopentasiloxane
14	19.358	1395407	1.82	96	73.00	Hexadecanoic acid <n->
15	21.223	2949334	3.85	54	73.05	Decamethylcyclopentasiloxane
16	21.894	1446147	1.89	92	82.05	Neophytadiene
17	22.214	455586	0.59	95	67.05	Linoleic acid
18	22.319	1033923	1.35	89	79.05	Linolenate <methyl->
19	28.165	1204901	1.57	69	277.10	Phytol
20	38.235	1249610	1.63	67	81.10	Torulosol
21	38.584	4600200	6.00	65	83.10	Larixol
22	39.440	5567702	7.26	66	145.10	Torulosol

Preliminary Qualitative Phytochemical Analysis

The preliminary qualitative phytochemical analysis (PQPA) of *S. urens* leaf extract revealed the presence of alkaloids, phenols, tannins, flavonoids, terpenoids, saponins, glycosides, steroids, and proteins. These compounds were detected across all three solvent extracts (methanol, water, and chloroform), except steroids in methanol; terpenoids and steroids in distilled water; and alkaloids, tannins, saponins, glycosides, and steroids in chloroform (**Table 1**).

Similarly, *S. alternata* leaves have been reported to contain alkaloids, saponins, flavonoids, tannins, and hydrogen cyanide, with additional evaluation of their anti-nutritional components (21). *Strobilanthes integrifolius* leaves were shown to contain alkaloids, carbohydrates, glycosides, saponins, proteins, phytosterols, fixed oils, phenolics, and flavonoids, while its stems exhibited the presence of alkaloids, carbohydrates, phytosterols, terpenoids, phenolics, and flavonoids. Likewise, *Strobilanthes blume* leaves were reported to contain carbohydrates, phytosterols, terpenoids, fixed oils, and phenolics, and its

stems to contain alkaloids, carbohydrates, phytosterols, terpenoids, phenolics, and flavonoids (22).

Quantitative Phytochemical Estimation

The quantitative phytochemical estimation (QPE) of *S. urens* leaf extract revealed that the total polyphenol content was 235.50 ± 5.44 µg/mg in distilled water, 270.50 ± 3.30 µg/mg in methanol, and 199.25 ± 2.50 µg/mg in chloroform. The total flavonoid content was 135.57 ± 8.88 µg/mg in distilled water, 260.57 ± 3.93 µg/mg in methanol, and 150.15 ± 5.63 µg/mg in chloroform. The total tannin content was 28.88 ± 2.89 µg/mg in distilled water, 130.03 ± 7.02 µg/mg in methanol, and 2.45 ± 3.51 µg/mg in chloroform. The alkaloid content was 105.71 ± 5.20 µg/mg in distilled water, 211.79 ± 1.99 µg/mg in methanol, and 53.06 ± 5.85 µg/mg in chloroform (**Table 2**).

S. alternata has been reported to contain phytates, tannins, oxalates, and cyanogenic glycosides (23). *Strobilanthes kunthiana* was reported to possess tannins, phenolics, and flavonoids (24). Similarly, *Strobilanthes heyneana* root extract exhibited a total phenolic content of

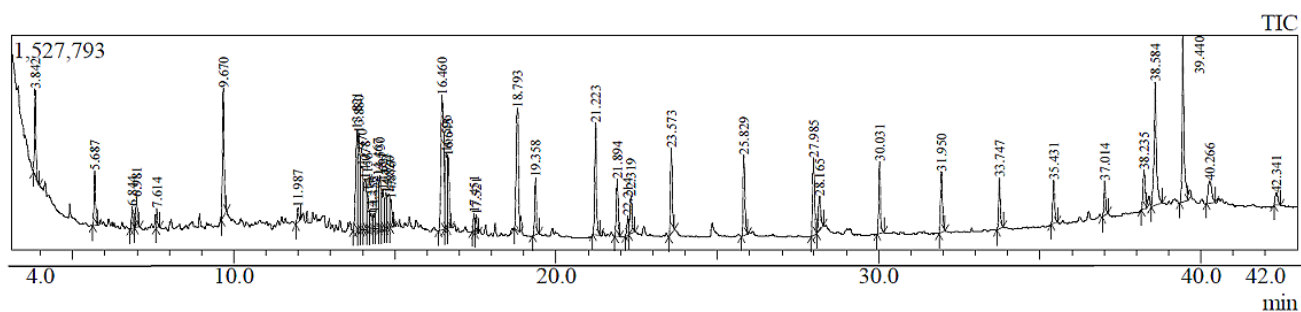


Figure 2. Chromatogram of *Strobilanthes urens* methanol leaf extract.

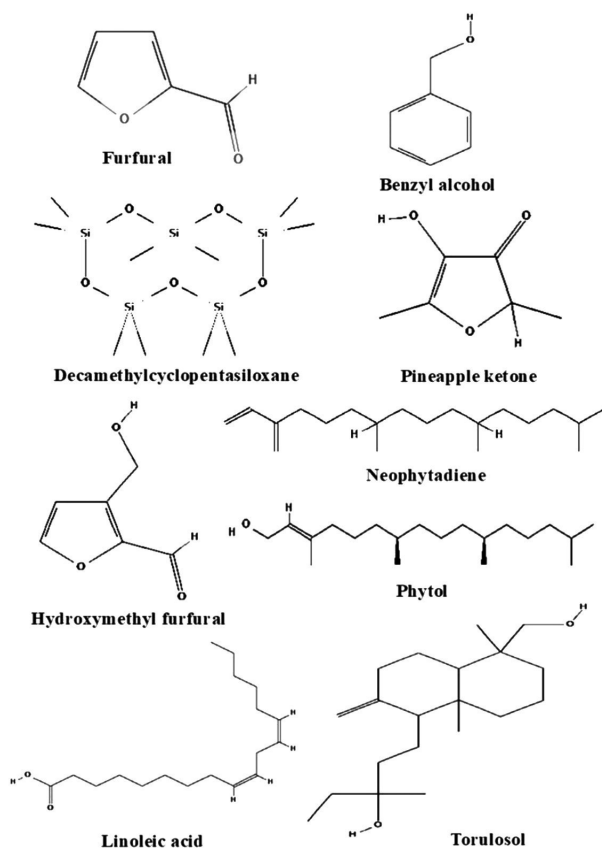


Figure 3. Structures of compounds identified in the methanolic leaf extract of *Strobilanthes urens*.

125.53 ± 2.29 $\mu\text{g}/\text{mg}$ and a flavonoid content of 32.79 ± 0.62 $\mu\text{g}/\text{mg}$ (25).

GC-MS Analysis

The GC-MS analysis of the methanolic extract of *S. urens* revealed a total of 45 bioactive compounds with distinct retention times and area percentages (Table 3 and Figure 2). The major compounds identified include Furfural (RT 3.842), Pineapple ketone (RT 7.614), 5-Methylfurfural (RT 5.687), Benzyl alcohol (RT 6.844), Decamethylcyclopentasiloxane (RT 13.821), Neophytadiene (RT 17.451), Phytone (RT 17.452), n-Hexadecanoic acid (RT 19.358), Linoleic acid (RT 22.214), Phytol (RT 28.165), Torulosol (RT 38.235), and Larixol (RT 38.584). The corresponding chemical structures of these compounds are shown in Figure 3.

Similarly, GC-MS analysis of *S. crispus* extracts reported

18 compounds in the leaves, 23 compounds in the stem, and 12 compounds in the roots, including hexadecanoic acid, 9-octadecanamide, eicosane, squalene, cholesterol, vitamin E, campesterol, stigmasterol, γ -sitosterol, lupeol, and betulin (26). *Strobilanthes glutinosus* chloroform and n-hexane extracts revealed 81 compounds, such as methyl esters, n-hexadecanoic acid, lupeol, linoleic acid, stigmasterol, γ -sitosterol, and linoelaidic acid, many of which have been reported to exhibit antioxidant, antidiabetic, anti-inflammatory, antimicrobial, antiprotozoal, and anticancer activities (27).

FTIR Analysis

FTIR analysis was performed to identify the functional groups of bioactive components based on peak values, allowing classification of compound types (Figure 4). The detected peaks ranged from 4000 cm^{-1} to 600 cm^{-1} . The major absorption bands corresponded to N-O-H, N-H, C-H, N=C=S, C=C, O-H bending, and C-F stretching, which indicate the presence of intermolecular vibrations, secondary amines, alkanes, isothiocyanates, conjugated alkenes, nitro compounds, carboxylic acids, and fluoro compounds.

Additional peaks such as O-H, C-H, O=C=O, S-C $\equiv$ N, C=C, S=O, and CO-O-CO were associated with alcohols, aldehydes, carboxylic acids, thiocyanates, cyclic alkanes, sulfonyl chlorides, and anhydrides. The O-H, N-H, O=C=O, C=C, C-H, S=O, C-N, and C-O stretching bands corresponded to alcohols, secondary amines, carbon dioxide, conjugated alkenes, alkanes, sulfonyl chlorides, aromatic amines, and primary alcohols, respectively, in methanol, water, and chloroform extracts.

S. crispus leaves were previously reported to exhibit functional groups such as C-Cl (halo compounds), C-N (amines), C=C (alkenes), N-H (aliphatic primary amines), M-O (metal-oxygen bonds), C-O (tertiary alcohols), and O-H stretching (alcohol groups) (28). Similarly, *Strobilanthes ciliatus* Nees showed characteristic peaks for S=O (sulfates), C-H bending (alkanes), C-O (alkanes), N-H (amines), C-H (carboxylic acids), C-H (aryl ketones), N-H (amides), C=O (amides), =C-H (aldehydes), and C=O (ketones) (29).

Antioxidant Activity by DPPH Assay

The antioxidant activity of *S. urens* extracts revealed IC₅₀ values of 79.23 ± 3.7 $\mu\text{g}/\text{mL}$ and 237.00 ± 12.37 $\mu\text{g}/\text{mL}$ for the methanol and distilled water extracts, respectively, while the chloroform extract showed no detectable activity. Vitamin C was used as a standard (Table 4). The strong antioxidant potential observed in the methanolic extract may be attributed to its high flavonoid content, as flavonoids are known to exhibit antioxidant activity primarily through metal chelation mechanisms (35).

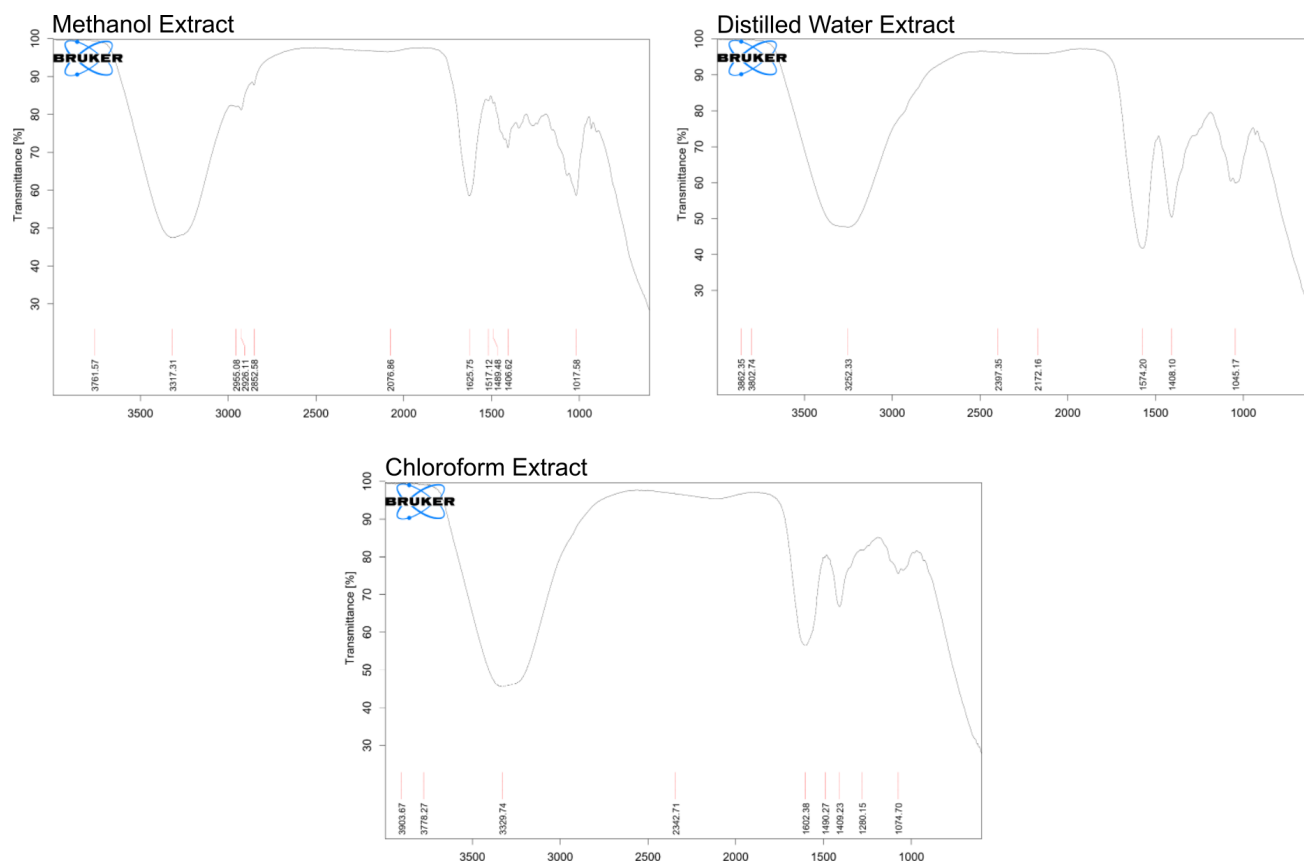


Figure 4. Analysis of functional groups in methanolic, distilled water, and chloroform extracts of *Strobilanthes urens* by FTIR.

Table 4. Antioxidant activity of *Strobilanthes urens* leaf extract.

No.	Samples	DPPH (IC ₅₀ µg/ml)
1	SULF M	79.23 ± 3.78
2	SULF DW	237.00 ± 12.37
3	SULF CHL	ND
4	Std (Vit-C)	26.21 ± 2.84

Note: Not detected (ND) indicates there is no activity found in the tested concentration.

S. kunthiana has been reported to show significant antioxidant activity, with IC₅₀ values of 19.97 ± 1.30 µg/mL and 20.97 ± 1.74 µg/mL in the DPPH assay, and 4.36 ± 0.38 µg/mL in the ABTS assay (27). Similarly, *S. crispus* leaves demonstrated potent radical-scavenging activity, with IC₅₀ values of 44.1, 58.2, and 78.3 µg/mL as determined by the DPPH assay (30).

Antimicrobial Activity

The antimicrobial activity of *S. urens* extracts demonstrated clear zones of inhibition at the tested concentrations across all three solvent extracts. The bacterial strains used for antibacterial testing were *S. aureus* and *E. coli*, while the fungal strains tested were *A. flavus* and *P. anomala*. No antifungal activity was observed at the tested concentrations. Kanamycin was used as the standard control (see **Figure 5** and **Supplementary Table 1**).

Previous studies on *S. crispus* and *Clianthus nutans*

reported activity against a total of eight Gram-positive and eight Gram-negative bacteria using the agar diffusion method with ethanol, acetone, and chloroform extracts. Among these, *Pseudomonas aeruginosa* exhibited the largest zone of inhibition, indicating strong susceptibility. This suggests that certain phytochemicals present in the extracts may specifically target *P. aeruginosa*, contributing to its pronounced antibacterial effect (31).

Cytotoxicity by MTT Assay

The cytotoxicity assay results revealed that the percentage of cytotoxicity against the MCF-7 cell line showed IC₅₀ values of 188.10 ± 39.99 µg/mL and 88.18 ± 8.29 µg/mL, while the HepG2 cell line exhibited IC₅₀ values of 129.34 ± 3.94 µg/mL and 156.38 ± 5.64 µg/mL. Doxorubicin was used as the standard control (**Figures 6, 7, and Supplementary Table 2**).

The methanolic extract of *S. crispus* was previously reported to show limited cytotoxicity against HepG2 cells at concentrations ranging from 31.25 µg/mL to 250 µg/mL, indicating no significant cell death (32). In contrast, *S. crista* leaf extracts in ethyl acetate and chloroform demonstrated potent cytotoxic activity against the HepG2 cell line, with IC₅₀ values of 38.8 µg/mL (33). Furthermore, *S. crispus* stem and leaf extracts exhibited cytotoxic activity against the MCF-7 cell line, with IC₅₀ values of 23 µg/mL and 38 µg/mL for the leaf extract, while the stem extract showed higher IC₅₀ values of 74–86 µg/mL (34). Lower IC₅₀ values indicate higher potency due to stronger interactions with cellular targets, whereas higher IC₅₀ values reflect weaker cytotoxic potential (35).

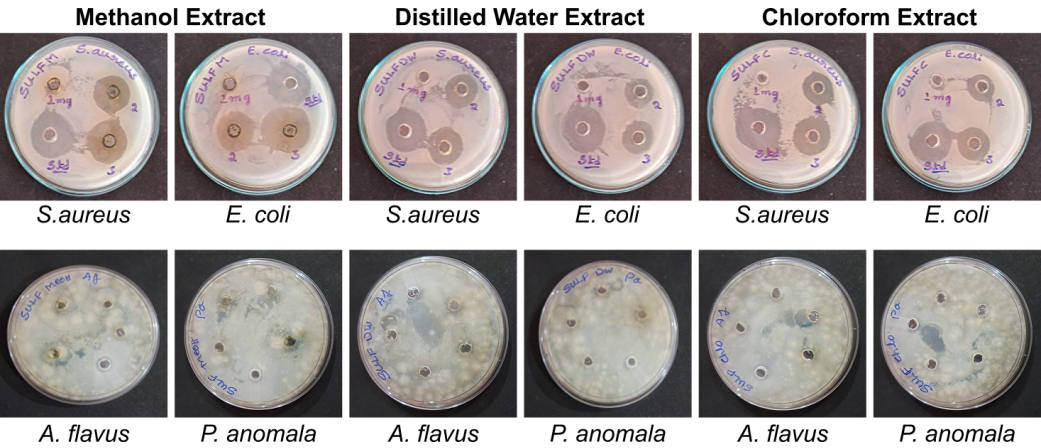


Figure 5. Antimicrobial activity of methanolic, distilled water, and chloroform leaf extracts of *Strobilanthes urens*.

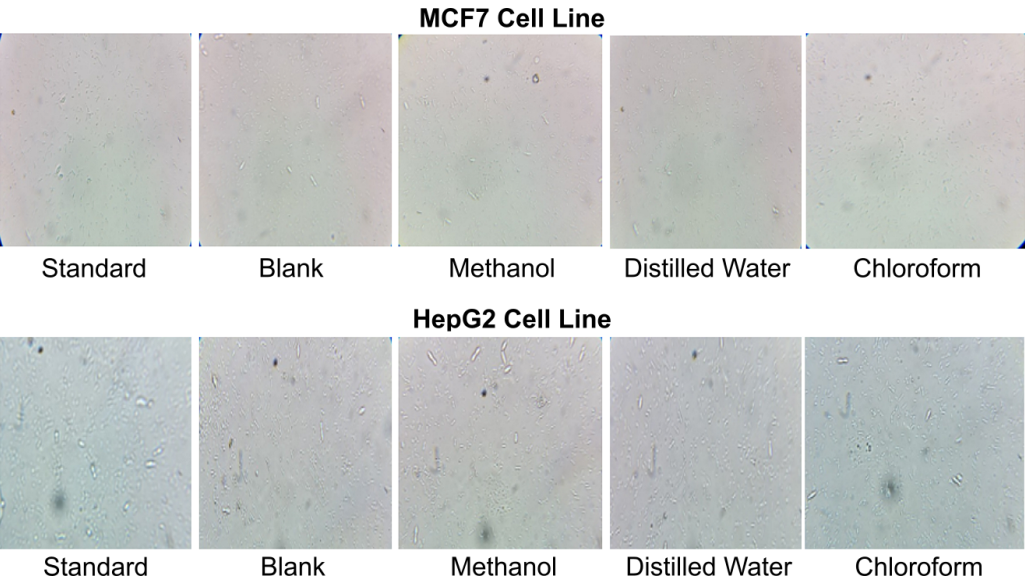


Figure 6. Microscopic images showing the cytotoxic effects (% cell viability) of methanolic, distilled water, and chloroform leaf extracts of *Strobilanthes urens* on MCF7 and HepG2 cell lines at a concentration of 160 µg/mL.

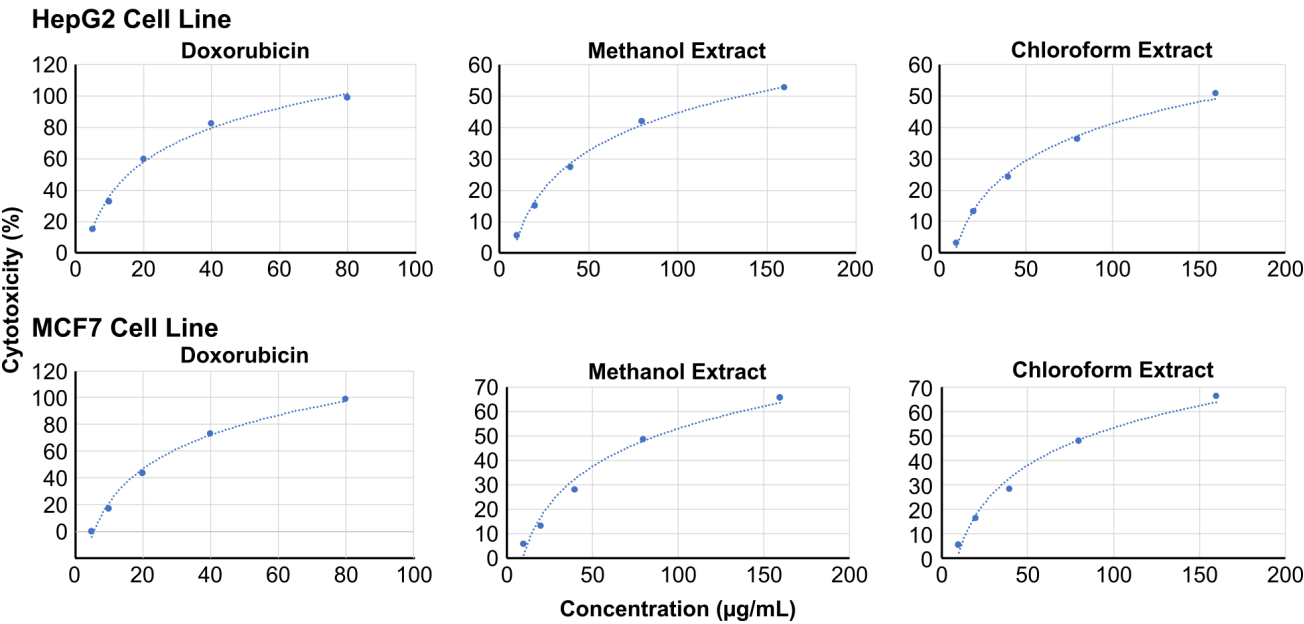


Figure 7. Cytotoxicity assay showing different concentrations of *Strobilanthes urens* leaf extracts against HepG2 and MCF7 cell lines.

Conclusion

The present study reveals that the various phytochemicals present in *S. urens* (B. Heyne ex Roth) J.R.I. Wood. These are linked to various pharmacological and biological activities. The various compounds identified might be a good source of novel drugs as the GC-MS analysis shows a good anti-cancer potential from MCF-7 and HepG2 show good therapeutic values in crude extracts against various ailments. From the present research, the Torulosol bioactive compound shows potent activity. It is concluded that the methanol crude extract shows a promising tool in the pharmaceutical and therapeutic applications. Further study is needed to isolate, characterize, and determine the pharmacological activities of compounds reported in our study.

Abbreviations

SULF M = *Strobilanthes urens* methanol extract; SULF DW = *Strobilanthes urens* distilled water extract; SULF CHL = *Strobilanthes urens* chloroform extract; SULF = *Strobilanthes urens* leaf; GC-MS = Gas chromatography mass spectroscopy.

Declarations

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Conflict of Interest

The authors declare no conflicting interest.

Data Availability

The unpublished data is available upon request to the corresponding author.

Ethics Statement

Not applicable.

Funding Information

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Supplemental Material

The supplementary data can be downloaded at the link: <https://etflin.com/file/document/202510310200431243264911.docx>

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Additional Information

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