



Review Article

The chemical constituents and diverse pharmacological importance of *Tinospora cordifolia*

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ABSTRACT

Tinospora cordifolia is a popular medicinal plant which is used in several traditional medicines to cure various diseases. The common names are Amrita and Guduchi and belong to the family of Menispermaceae. It is considered an essential herbal plant of Indian system of medicine (ISM) and has been used in the treatment of fever, urinary problem, dysentery, skin diseases leprosy, diabetes, and many more diseases. The plant reported containing chemical compound including Alkaloids, Terpenoids, Lignans, Steroids and others that establish the phytochemistry and pharmacological activity of *Tinospora cordifolia*. The present review highlights the pharmacological importance viz antioxidant activity, antimicrobial activity, antibacterial activity, antifungal activity, anti-diabetic activity, antistress activity, hypolipidaemic effect, hepatic disorder, anticancer anti HIV potential, antiosteoporotic effects, antitoxic effects, wound healing, anticomplementary activity, and immunomodulating activity, systemic infection and Parkinson's disease.

1. Introduction

Herbal formulations are medicinal preparation of one or more herbs present in specified quantities to give the benefits meant for cosmetic, diagnose and to mitigate diseases of human beings or animals [1]. It is also known as botanical medicine or phytomedicine. Earlier in the twentieth century, herbal medicine was the prime medication system as antibiotics or analgesics were not available. Increasing use of an allopathic system of medicine due to its fast therapeutic action and herbal medicine gradually lost their popularity among the people. For example, Curcuma is used in Traditional Chinese Medicine for more than two thousand years to treat anti-inflammatory and robust antioxidant [2, 3]. About 70–80% of people are still using herbal medicine for their primary health because of the less side effect and better compatibility with the human body [4]. Herbal medicine has gained momentum and is more effective as compared to synthetic drugs.

T. cordifolia (synonym: *Tinospora sinensis* (Lour.) Merr.) is also known as Guduchi/Amrita and its names in Latin: *Tinospora cordifolia* (Wild) Hook. f. & Thomson, English: *Tinospora* Gulancha/Indian *Tinospora*, Hindi: Giloya. It belongs to the family of Menispermaceae and is found in Myanmar, Sri Lanka, and China [5].

The plant is commonly used as traditional ayurvedic medicine and has several therapeutic properties [6, 7] such as jaundice, rheumatism, urinary disorder, skin diseases, diabetes, anemia, inflammation, allergic condition, anti-periodic, radioprotective properties, etc. [8, 9] The root of Giloya (*T. cordifolia*) is used as potent emetic and for bowel obstruction. The starch of this plant serves a beneficial household remedy for chronic fever, relieves burning sensation, increases energy and appetite.

Giloya is useful in the treatment of helminthiasis, heart diseases, leprosy, rheumatoid arthritis, support the immune system, the body's resistance to infections, supports standard white blood cell structure, function, and levels [10]. It also helps in digestive ailments such as hyperacidity, colitis, worm infestations, loss of appetite, abdominal pain, excessive thirst, and vomiting, and even liver disorders like hepatitis [11, 12]. This pharmacological activities of the plant is due to its chemical constituents like diterpenoid lactones, glycosides, steroids, sesquiterpenoid, phenolics, aliphatic compounds, essential oils, a mixture of fatty acids, and polysaccharides and is present in a different part of the plant body, including root, stem, and whole part [13].

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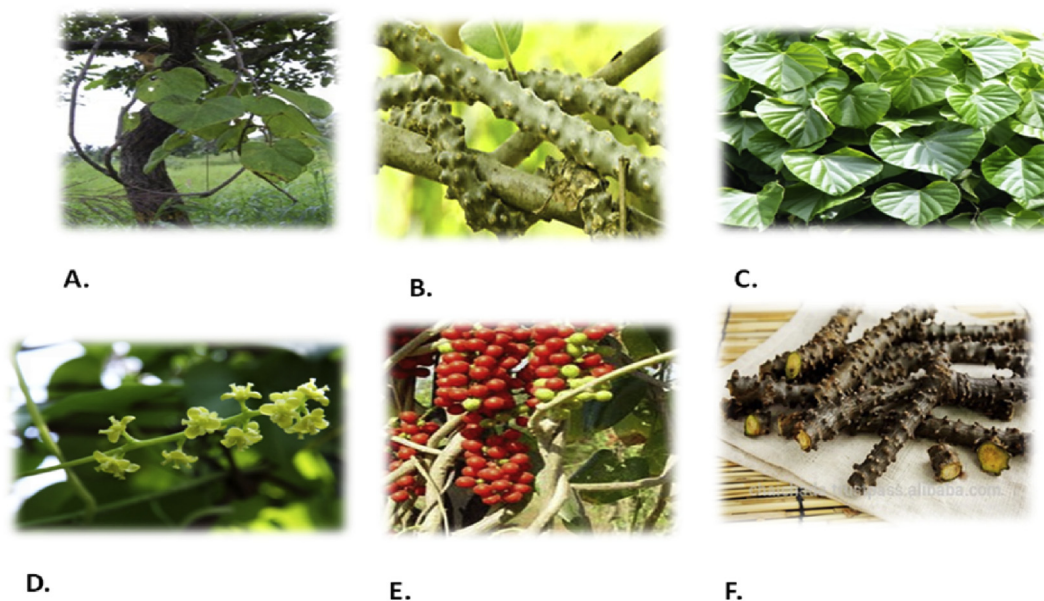


Fig. 1. Morphology of *Tinospora cordifolia* A) steam B) root C) leaves D) flower E) fruit F) seed.

Table 1
Some of the essential constituents of *T. cordifolia*.

Active Component	Compounds	References
Terpenoids	Tinosporide, Furanolactone diterpene, Furanolactone clerodane diterpene, furanoid diterpene, Tinosporaside, ecdysterone makisterone and several glucosides isolated as poly acetate, phenylpropene disaccharides cordifoliside A, B and C, cordifoliside D and E, Tinocordioside, cordioside, palmatosides C and F, Sesquiterpene glucoside tinocordifoliside, Sesquiterpene tinocordifolin.	[26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36]
Alkaloids	Tinosporine, (S), Magnoflorine, (S), Berberine, (S), Choline, (S), Jatrorrhizine, (S), 1,2-Substituted pyrrolidine(S), Alkaloids, viz. jatrorrhizine, palmatine, beberine, tembeterine, choline.	[37, 38, 39, 40, 41]
Lignans	3 (a, 4-dihydroxy-3-methoxybenzyl)-4-(4-hydroxy-3-methoxybenzyl), (S)	[42]
Steroids	Giloinsterol, (S), β -Sitosterol, (S), 20a-Hydroxy ecdysone, (S).	[43, 44, 45, 46]
Others	Giloin, Tinosporan acetate, Tinosporal acetate, Tinosporidine, Heptacosanol, Octacosanol, sinapic acid, Tinosponone, two phytoecdysones: an immunologically active arabinogalactan.	[47, 48, 49, 50, 51]

2. Main text

2.1. Pharmacognostic description

It is a large deciduous, extensively spreading climbing shrub with several coiled branches with a different type of morphology. The stem of the plant is filiform, fleshy and climbing in nature; bark is white to gray [14]. Powder of the stem is creamish brown or dark brown, characteristic odor, bitter taste and is used in dyspepsia, fever, and urinary diseases [15]. The starch obtained from the stem known as “Guduchi-satva.” It is highly nutritive and digestive. Leaves of this plant are simple, alternate, long-petioled (approximately 15 cm); round, pulvinate, heart-shaped, twisted partially and halfway around. Lamina is ovate, 10–20 cm long, seven nerved and deeply cordate at the base and membranous [16].

Flowers are unisexual, axillary position, 2–9 cm long leaflet branches and greenish-yellow in colour, male flowers are clustered, female usually solitary [17]. Its fruits are single-seeded, fruits during the winter and flowers grow during the summer [18]. The root is thread-like, aerial, squairshin, sometimes continuously lengthening touch the ground [19], aerial roots are characterized by tetra to penta arch primary structure [20]. The seeds are curved shape [21] and endocarp is variously ornamented, which provides critical taxonomic characters. morphology of the plants has been depicted in (Fig. 1).

2.2. Chemical constituent

The chemical constituents of *T. cordifolia* belong to different classes such as alkaloids, glycosides, steroids, phenolics, aliphatic compounds, polysaccharides, leaves are rich in protein (11.2%), calcium and phosphorus [22]. The stem contains clerodane furono diterpene glucoside (amritoside A, B, C, and D) and the structure has been established by different spectroscopic studies [23, 24, 25]. Some of the essential constituents reported in Table 1 and major constituents in Fig. 2 whereas the structure of the active chemical constituent for *Tinospora cordifolia* has been depicted in (see Fig. 3).

2.3. Pharmacological activities

Different pharmacological activities of *T. cordifolia* has been reported by the researcher, which has been described:

2.3.1. Antioxidant activity

Mehra et al., prepared the formulation and evaluated its antioxidant activity by DPPH (1-diphenyl-2-picrylhydrazyl) free radical scavenging method. They estimated the total flavonol and total phenolic content. Using the result of the formulation showed potent antioxidant activity and inhibitory concentration (IC₅₀) at 5 μ g/ml as compared to standard drug ascorbic acid [52]. George et al., reported the methanolic, ethanolic, and water extracts of *T. cordifolia* for their antioxidant activity, in which the stemic ethanol extract increased the erythrocytes membrane lipid peroxide, catalase activity and decrease the superoxide dismutase, glutathione peroxidase in alloxan-induced diabetic rats. The leaves extract of methanol, partitioned in water with ethyl acetate and butanol at 250 mg/ml, and showed their antioxidant activity, extracts of methanol phosphomolybdenum and metal chelating activity were high

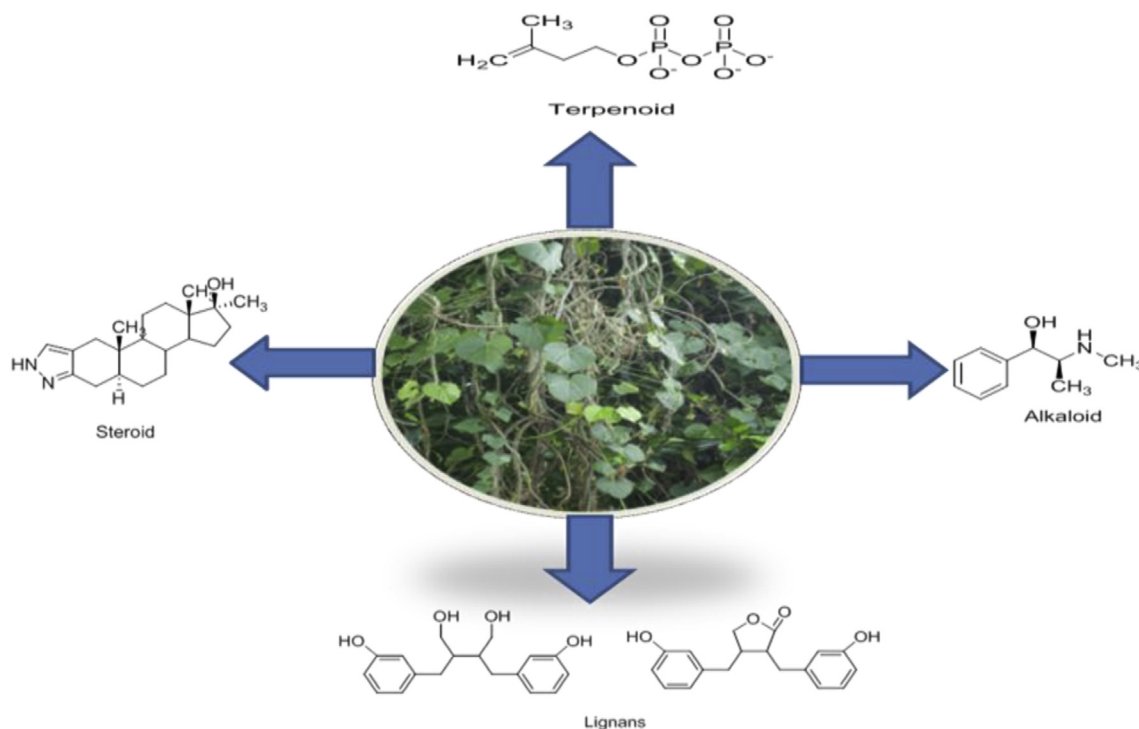


Fig. 2. Major constituent of *Tinospora cordifolia*: terpenoid, alkaloid, lignans, steroids.

followed by ethyl acetate, butanol, and water extract [53]. It also decrease level of free radical species of diabetic rat and up-regulate the anti-oxidant enzyme [54, 55, 56], scavenging activity for free radical of methanol extract was high compared with phenol extract [57]. This plant modifies the different enzymatic system which controls the production of these reactive species and maintains the oxidative load by regulating the lipid peroxidation process and glutathione level [58, 59]. Premnath *et al.*, dried the leave of *T. cordifolia* and powdered and extracted with chloroform, methanol, ethanol hexane, and water. Antioxidant assay by different *in-vitro* models, lipid peroxidation inhibitory activity, DPPH radical scavenged, and superoxide radical scavenging activity. Other solvent extracts showed weak antioxidant activity, whereas ethanol extract had high antioxidant activity. The results suggested that the antioxidant compound are better in ethanol extract, and there is a direct correlation between the total polyphenols extracted and its anti-oxidant activity [60].

2.3.2. Antimicrobial activity

Antimicrobial activity of the *T. cordifolia* with different solvents on different micro-organism, showed good antifungal and antibacterial activity [61]. Jeyachandran *et al.*, reported the antimicrobial activity of stem extracts by *in-vitro* analysis against both gram-positive and gram-negative bacteria and showed good therapeutic activity on the infectious disease. It has taken a methanolic extract of *T. cordifolia* against both bacteria group [62]. Narayanan *et al.*, have reported antibacterial activity of plants to extract against *Escherichia coli*, *Proteus vulgaris*, *Salmonella typhi*, *Salmonella paratyphi*, *Salmonella typhimurium*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa*, *Enterobacter aerogene*, *Shigella flexneri*, *Staphylococcus aureus* and *Serratia marcescens* (Gram-positive bacteria) [63]. The aqueous, ethanol and acetone extract of *T. cordifolia* inhibited the activity on clinical isolates of urinary pathogens *Klebsiella pneumoniae* and *Pseudomonas aeruginosa*. [64] Singh *et al.*, has reported silver nanoparticles from the stem of *T. cordifolia*, which possess antibacterial activity against the different strains of bacterias [65]. Allemailem *et al.*, have reported the antifungal activity of *T. cordifolia*, which was determined using the agar well plate diffusion method. The aqueous extract of *T. cordifolia* showed potent activity against *A. fumigatus*,

Aspergillus flavus, and *Aspergillus niger* (fungus) in the study [66]. Agarwal *et al.*, studied *in-vitro* extract of *T. cordifolia* was obtained using 100% ethanol by maceration process. They prepared etholic extract seven different concentrations and tested against *S. mutans* in brain–heart infusion agar medium. Plates were incubated aerobically at 37 °C for 48 h, using Vernier caliper and measured the zone of inhibition. 0.2% chlorhexidine and dimethylformamide were used as positive and negative controls, respectively. This experiment data were analysed by descriptive-analytic tests. Which showed the maximum antibacterial activity of *T. cordifolia* a volume of 40 µl at 2% concentration with a zone of inhibition of 19 mm. A 30 µl volume of 0.2% chlorhexidine showed a zone of inhibition of 28 mm, and dimethylformamide showed no zone of inhibition [67]. Khan *et al.*, reported the antifungal activity TCAE (*Tinospora cordifolia* aqueous extract) was tested for *in-vitro* against the isolates of different *Aspergillus* species. To evaluate *in vivo* activity, different doses (10, 25, and 50 mg/kg) of TCAE were orally administered in *A. fumigatus*-infected mice for seven days for the evaluation of *in vivo* activity. The effectiveness of aqueous extract on the basic survival rate and assessing the fungal burden in the kidney of the treated mice [68]. Prasad *et al.*, studied the anti-oxidant and antimicrobial properties phenolic extract of *T. cordifolia* stem and root. Total reducing power, hydrogen peroxide scavenging activity assay, and hydroxyl radical scavenging activity were checked using different *in-vitro* assays. The ethanolic extract showed maximum 87.2% and 91.0% free radical scavenging activity concerning H₂O₂ scavenging and hydroxyl free radical scavenging assay [69].

2.3.3. Anti-toxic effects

Gupta *et al.*, reported the extract to scavenge free radicals generated during aflatoxicosis. It showed protective effects of *T. cordifolia* on thiobarbituric acid reactive substances (TBARS) levels and increase the level of GSH, ascorbic acid, protein, and the activities of anti-oxidant enzymes viz., Superoxide Dismutase (SOD), Catalase (CAT), GPx enzyme, Glutathione S-transferase (GST) and glutathione reductase (GR) in kidney. The alkaloids such as choline, tinosporin, isocolumbin, palmatine, tetrahydropalmatine, and magnoflorine present in the plant of *T. cordifolia* showed protection against aflatoxin-induced nephrotoxicity.[htt](#)

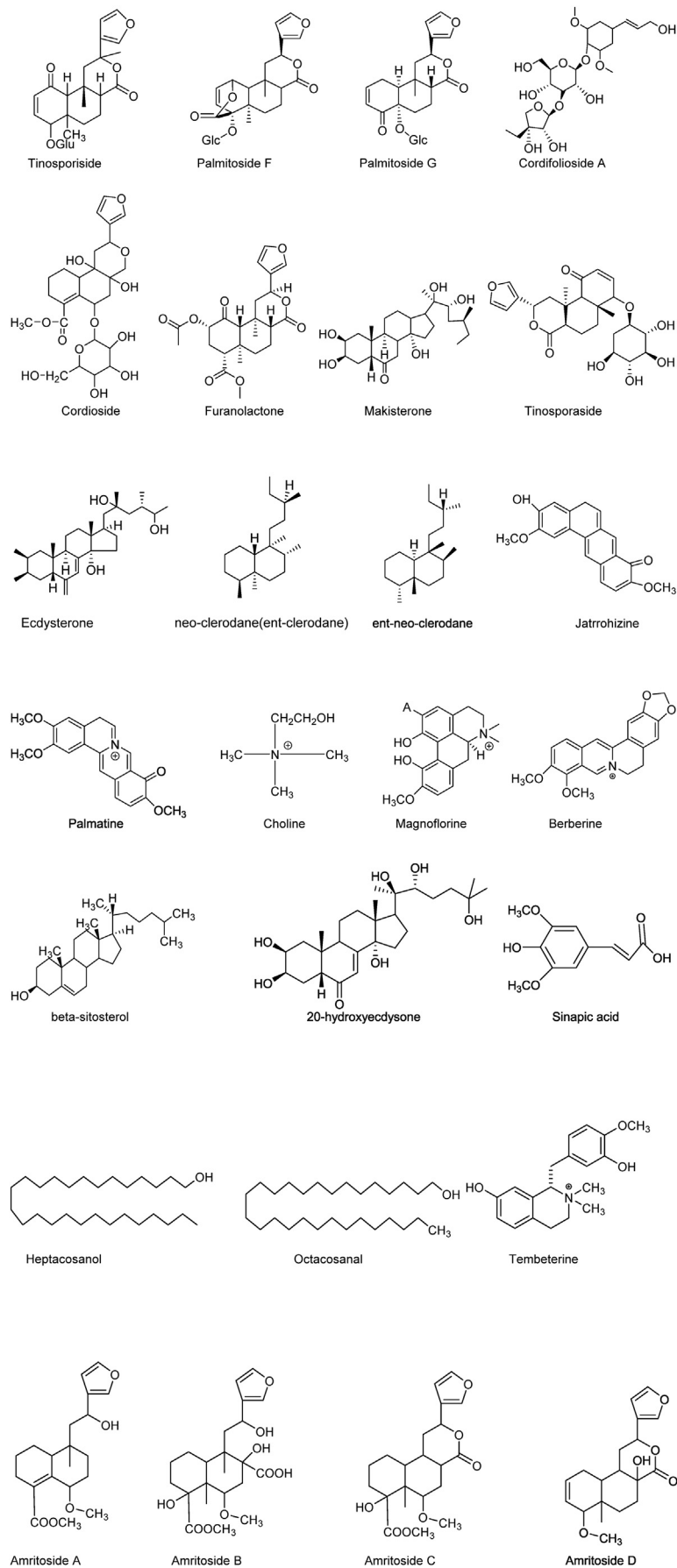


Fig. 3. Structure of the chemical constituent of *T. cordifolia*.

[p://www.ancientscienceoflife.org/article.asp?issn=0257-7941;year=2012;volume=31;issue=4;page=151;epage=159;aulast=Saha](http://www.ancientscienceoflife.org/article.asp?issn=0257-7941;year=2012;volume=31;issue=4;page=151;epage=159;aulast=Saha) [70] Sharma *et al.*, studied the stem and leaves extract of the plant has shown hepatoprotective effect in Swiss albino male mice against lead nitrate induced toxicity [71]. Oral administration of plant extracts prevented the occurrence of lead nitrate induced liver damage [72]. They also checked the decreased level of SOD, CAT and increased level of aspartate aminotransferase (AST), alanine aminotransferase (ALT), alkaline phosphate (ALP), and acid phosphates (ACP). <http://www.ancientscienceoflife.org/article.asp?issn=0257-7941;year=2012;volume=31;issue=4;page=151;epage=159;aulast=Saha> [72] Synergistic administration of aqueous extract of stem and leaf increased the activities of SOD and CAT and decreased the levels of AST, ALT, ALP, and ACP enzymes. <http://www.ancientscienceoflife.org/article.asp?issn=0257-7941;year=2012;volume=31;issue=4;page=151;epage=159;aulast=Saha> [72] Protective role of aqueous extract of the plant stem and leaves overcoming the toxic effects was also reported by them. <http://www.ancientscienceoflife.org/article.asp?issn=0257-7941;year=2012;volume=31;issue=4;page=151;epage=159;aulast=Saha> [71].

2.3.4. Antidiabetic activity

The anti-diabetic activities is due to alkaloids (Magnoflorine, Palmatine, Jatrorrhizine), tannins, cardiac glycosides, flavonoids, saponins, etc. [73] The crude extract of the stem in ethyl acetate, dichloromethane (CDM), chloroform and hexane was studied for inhibition of the alpha-glucosidase enzyme. The activity of the enzyme inhibited hypoglycemic action in diabetic animal and normal animals. The aqueous extract was studied in the rats, without the addition of *Tinospora cordifolia* extract increase in glucose by 21.3%, insulin by 51.5%, triglycerides by 54.12%, and glucose-insulin index by 59.8 when plant containing extract was given. The fructose-induced abnormalities in the liver involving lipid peroxidation, protein carbonyl groups, GSH levels, and enzymatic antioxidants decreased [74]. Methew *et al.*, have reported *in vivo* studies of different extracts of the plants on a diabetic patient. Sedimental extract of *Tinospora* on the subject was studied at 30 d ay. Different doses (200 and 400 mg/kg b. w) of Ethanolic extract of *T. cordifolia* leaves were prepared. The doses were administered orally for ten days and 30 days in streptozotocin-diabetic albino rats. *T. cordifolia* showed the antidiabetic activity in diabetic animals an efficacy an 50%–70% compared to insulin [75].

From Guduchi Prasant *et al.*, isolated alkaloids, cardiac glycosides, saponins, flavonoids, tannins, and steroids that contains anti-diabetic property. Alkaloids from this plants showed insulin-mediated actions due to insulin hormone [76]. Gestational diabetes can increase the GSH content and other reactive species that can act as a threat to the mother as well as the fetus. The study based upon the pregnant rat using *T. cordifolia* was incorporated in the daily diet to a diabetic-pregnant rat (streptozotocin-induced diabetes), which showed a protective effect by reducing the oxidative load thereby preventing the relative incidence of diseases and any birth defect [77]. In a diabetic rat model, *T. cordifolia* root extracts of Guduchi attenuated the brain mediated lipid level and down-regulated the blood glucose and urinary glucose level emphasizing its anti-diabetic and lipid-lowering activity [78].

The root extract of Guduchi showed an antihyperglycemic effect in the alloxan-induced diabetic model by decreasing its excess glucose level in urine as well as in normal [79]. Certain herbal preparation, including Guduchi like Ilogen-Excel, Hyponidd, and Dihar have been tested in diabetic rat models, the anti-diabetic activity of *T. cordifolia* was observed. The effects by Ilogen Excel down the level of excess glucose in the blood and enhance the insulin efficiency by increasing its amount in the systemic circulation. Hyponidd is reported, and it maintained the oxidative load by decreasing reactive species and reduced the glucose-mediated hemoglobin count. when the tested of 'Dihar' for one and a half month in streptozotocin-induced diabetic model decreased the urea as well as creatinine amount in the blood with an increase in enzyme activities [54, 80, 81, 82].

2.3.5. Antistress activity

Sarma *et al.*, reported ethanolic extract of *T. cordifolia* at the dose of 100 mg/kg gives significant anti-stress activity in all parameters compared with standard drug diazepam (dose of 2.5 mg/kg) [83]. The plant extract gives a moderate degree of behavior disorders and mental deficit response. The clinical research showed the improved I. Q level of patients. In Ayurveda, it acts as Medhya Rasayana or brain tonic by increasing mind power like memory and recollection [84].

2.3.6. Hypolipidemic effect

Stanely *et al.*, studied the hypolipidemic effect of an aqueous extract of the root on the rats weighing 2.5 and 5.0 g/kg body weight on sixth weeks, that resulted in decrease tissue cholesterol, reduction in serum, phospholipids, and free fatty acid in alloxan diabetic rats. The dose of root extract 5.0 g/kg body weight showed the highest hypolipidaemic effect. When the level of serum lipids in diabetes increased, they represented coronary heart disease, lower the serum lipids level decreased the risk of vascular disease. The ability of *T. cordifolia* root extract to reduce the level of serum or tissue lipids in diabetics animals have never been studied before till then [85].

2.3.7. Hepatic disorder

Protective Effects of *Tinospora cordifolia* water extract (TCE) on Hepatic and Gastrointestinal Toxicity was reported by Sharma *et al.*, a significant increase in the levels of gamma-glutamyl transferase, aspartate transaminase, alanine transaminase, Triglyceride, Cholesterol, HDL and LDL ($P < 0.05$) in alcoholic sample whereas their level get down-regulated after TCE intervention, patients showed the normalized liver function of *T. cordifolia* stand to relieve the symptoms [86].

2.3.8. Anticancer activity

Ali *et al.*, studied the anticancer activity of *T. cordifolia* palmatine extract in animal models, alkaloid using response surface methodology (RSM). The extract indicates the anticancer potential in 7,12-dimethylbenz(a)anthracene DMBA induced skin cancer model in mice [87]. Rahul *et al.*, prepared the extract of 200, 400, 600 mg/kg dry weight in a dose depend upon manners. 50% methanolic extract of *cordifolia* to C57 BI mice for 30 days at a dose of 750 mg/kg body weight the tumor size reduced life span [88]. Mishra *et al.*, showed the anti-brain cancer potential, 50% ethanolic extract of *T. cordifolia* (TCE) using C6 glioma cells significantly induced differentiation in C6 glioma cells, and reduced cell proliferation [89].

2.3.9. Anti-HIV potential

Kalika *et al.*, showed that the root extract of *T. cordifolia* affects the immune system of HIV positive patient. The stem extract of *Tinospora cordifolia* reduces the ability of eosinophil count, stimulation of B lymphocytes, macrophages, level of hemoglobin, and polymorphonuclear leucocytes [90, 91].

2.3.10. Wound healing

Shanbhag T *et al.*, The present study was aimed at evaluating the wound healing profile of alcoholic extract of *T. cordifolia* and its effect on dexamethasone suppressed healing. Incision, excision, and dead space of the wound models were employed to investigate the wound healing potential of the plant increased tensile strength extract of *T. cordifolia* may be attributed to the promotion of collagen synthesis. The extract of *T. cordifolia* did not reverse dexamethasone suppressed wound healing [92].

2.3.11. Anti-osteoporotic effects

Abiramasundari *et al.*, reported *T. cordifolia* affect the proliferation, differentiation, and mineralization of bone-like matrix on osteoblast model systems *in-vitro* and hence finds potential application as an anti-osteoporotic agent. Alcoholic extract of *T. cordifolia* has been shown to stimulate the growth of osteoblasts, increasing the differentiation of cells

Table 2
Biological activities of *T. Cardifolia* concerning different parts of the plant.

Active compound	Biological activity
Terpenoids	Stem: Respiratory track infection [98], skin disease [5], Anti-hyperglycemic property [99].
Alkaloids	Stem and root of plant: Anti-cancer property [100], Antioxidant activity [101].
Lignans	The root of plant: Anti-neoplastic property [102], Antioxidant activity [101].
Steroids	Arial part of stem: Anti-stress activity [83].
Other	The whole part of Plant: Antidote to snakebite and scorpion sting, Analgesic and Neuropharmacological activities, Diabetes, Rheumatoid arthritis, Gout, Cancer, high cholesterol content, antipyretic, antileprotic, radioprotective [8,103, 104, 105, 106, 107].

N.B. Biological activities of different parts of *T. Cardifolia*. The number denoted on the disease are the references cited.

into the osteoblastic lineage and also increasing the mineralization of bone-like matrix [93]. Ecdysteroids isolated from the plant have been reported of protein anabolic and anti-osteoporotic effects in mammals. Beta-Ecdysone (Ecd) from *T. cordifolia* extracts have been reported to induce a significant increase in the thickness of joint cartilage, induce the osteogenic differentiation in mouse mesenchymal stem cells [94] and to relieve osteoporosis in osteoporotic animal models [62, 94]. Further 20-OH- β -Ecd isolated from *T. cordifolia* has been reported for its anti-osteoporotic effects [93], thus highlighting the role of *Tinospora cordifolia* in the treatment of osteoporosis and osteoarthritis [95].

2.3.12. Anticomplement activity and immunomodulating activity

Kapil et al., Studied the syringin (TC-4) and cordiol (TC-7) isolate from *T. cordifolia* inhibited the *in-vitro* immune hemolysis of antibody-coated sheep erythrocytes by guinea pig serum. Immune hemolysis was reduced due to inhibition of the C3-convertase of the classical complement pathway. The compounds of *T. cordifolia* rise to significant increases in IgG antibodies in guinea pig serum. Cordioside (TC-2), cordiofolioside A (TC-5) and cordiol (TC-7) activated macrophase with increasing incubation times [96]. Sharma et al., isolated and characterised different classes of active compounds reported their immunomodulatory activity [7].

2.3.13. Parkinson's disease

Birla et al., reported *T. cordifolia* extract is highly attractive against the Parkinsonism. They observed the anti-inflammatory activity of aqueous extract in 1-methyl-4-phenyl-1,2,3,6-tetra hydroxyridine (MPTP)-intoxicated Parkinsonian mouse model. The extract reversed the behavior of the target MPTP-intoxicated mice and its suggest that *T. cordifolia* protected dopaminergic neurons by suppressing neuroinflammation in MPTP-induced Parkinsonian mouse model [97].

The plant exhibited multiple biological activities due to diverse chemical constituents present in it. The biologically active chemical molecules are present in different parts of the *T. Cordifolia*. That is the explanation for curing various ailments in human being using a different part of the miraculous plant from the ancient ages (Table 2).

3. Conclusions

T. cordifolia is a medicinal plant having various type of compounds. The different bioactive compounds, including alkaloids, steroids, glycosides, sesquiterpenoids, etc have been discussed. Present review spotlights the artistic antifungal activity, antioxidant activity, antimicrobial activity, antibacterial activity, hypolipidaemic effect, hepatic disorder, anticancer, Anti HIV potential, Antiosteoporotic effects, Antitoxic effects, Wound healing, anticomplementary activity, immunomodulating activity, systemic Infection and Parkinson's disease of *T. Cordifolia*. It has been used successfully in Ayurvedic medicine from the ancient era, and its

products are used for their better economic and therapeutic utilization. In this regard, further studies need to be carried out to explore *T. cordifolia* for its potential in preventing and treating diseases. This review can be used for further research investigations as well as clinical purpose in the development of novel drugs.

Declarations

Author contribution statement

All authors listed have significantly contributed to the development and All authors listed have significantly contributed to the development and the writing of this article. the writing of this article.

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The authors declare no conflict of interest.

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