

LC-MS BASED TARGETED METABOLITE PROFILING, AAS BASED MINERAL PROFILING IN METHANOL LEAF EXTRACT OF *Terminalia bellerica* (Gaertn.)

Comment [T11]: Write AAS in full

ABSTRACT

AIM: Natural products in medicinal plants are one of the important sources for modern drug discovery. Plant-based formulations play an important role in management of numerous diseases. To discover the medical potential of natural products, it is important to understand the bioactive compounds present in medicinal plants. Hence in this study, it is planned to assay the volatile compounds and minerals in the leaves of *Terminalia bellerica*

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STUDY DESIGN: In the present study, metabolites in leaves of *Terminalia bellerica* have been studied by AAS and LC-MS techniques.

Hence, the aim of this study, it is to identify the volatile compounds and minerals present in *Terminalia bellerica* leave.

RESULTS: AAS assay recorded the leaves of *T.bellerica* are rich in iron content (260mg/kg) when compared with Zn, Mn and copper content. LC-MS assay revealed chebulinic acid, chebulagic acid, punicafolin, myricetin rutine, galloyl punicalagin are in highest amount.

Comment [T13]: This study investigates the metabolites present in the leaves of *Terminalia bellerica* using AAS and LC-MS techniques.

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CONCLUSION: This plant could provide reservoir of iron that have greater therapeutic potential with fewer side effects than to synthetic drugs. Due to rich phyto nutrients in leaves of *T.bellerica*, it may be used in the development of health supplements. To develop new drugs in pharmaceutical industry, compounds in leaves of *T.bellerica* lead a pathway. Further isolation of novel compounds leads to the development of pharmacologically important drugs of specific function.

Comment [T15]: The AAS assay showed that the leaves of *T. bellerica* are particularly rich in iron (260 mg/kg) compared to their zinc, manganese, and copper content. Meanwhile, the LC-MS analysis identified chebulinic acid, chebulagic acid, punicafolin, myricetin, rutin, and galloyl punicalagin as being present in the highest concentrations

Key words: Medicinal herbs, *Terminalia bellerica* leaves, Phytoceutics, AAS assay, GC-MS assay.

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1. Introduction:

Plants have served as an essential source of drugs and remedies against diseases and health disorders since ancient times. Health benefits may be credited to the presence of the various phytochemicals like polyphenols, terpenes, anthocyanins, flavonoids, alkaloids and glycosides. Plants of antimicrobial activity can be related to their essential oils (1) or other isolated compounds such as alkaloids (2), flavonoids, tannins, phenolic acids (3), among

other chemical classes including naturally occurring peptides (4,5). These compounds possess their antimicrobial activity through various mechanisms such as depressing the nuclear or ribosomal enzyme(s) synthesis, altering the membrane structure and the electron flow, or affecting the metabolic activity of the microbial cell, as well as inhibiting the secretion of their toxins (6,7,8).

Comment [T17]: Downregulating

Natural products from medicinal plants are one of the important sources for modern drug discovery. Triphala is an ayurvedic drug prepared by mixing dried fruit powder of *Terminalia chebula*, *Terminalia bellerica* and *Phyllanthus emblica* in the ratio of 1:1:1 and also its individual constituent's fruit powder exhibit immuno-modulatory activity. The important constituents of the Triphala are chebulagic acid (CA), chebulinic acid (CI), gallic acid. These compounds inhibit TNF α induced pro-angiogenic and pro-inflammatory activities in retinal capillary endothelial cells by inhibiting p38, ERK and NF κ B phosphorylation. Polyphenol containing plants, of *Terminalia* species, have reportedly shown various health benefits and applications in pharma industry. *Terminalia* have a wide spectrum of pharmacological activities including antioxidant, anti-inflammatory, anti-cancer, hepatoprotective, and antimicrobial activities. Polyphenols have been also proven to lower the risk of cardiovascular diseases, enhance liver regeneration, and increase life expectancy.

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Traditional medicinal plants practices are used for the treatment of infectious diseases and they have very low level/low-level risk compared to synthetic drugs. Natural compounds in medicinal plants are essential sources for drug discovery (9) and are reported that natural products take up to 35% of the global medicine market (10). To discover the medical values of natural products, it is important to understand the ethno pharmacological uses of various medicinal plants (11). Although the modern medicines are quickly growing, large population still preferring herbal medicines due to the effectiveness, medical effectiveness, enhancing cost of modern medicines and cultural preferences (12,13,14). Based on WHO reports, 80% of the global population depends on traditional medicine, and 60% of the Indian rural population use herbal medicines (14). These natural medicines are generally safe, cost-effective, and efficient (14). Apart from the medical values, plants are also widely used as food, health care products (15), veterinary medicine (16), have extensive impacts on daily life. Herbal medicine is still the mainstay of health care in several developing countries. The World Health Organization has estimated that more than 80 % of the world's population in developing countries depends primarily on herbal medicine for basic healthcare needs.

Comment [T111]: "Make up" will be better

Comment [T112]: "are progressively increasing"

Comment [T113]: high cost

Comment [T114]: I'm uncertain whether natural medicines are generally safe unless their phytochemical constituents have been identified and they have received approval from the relevant regulatory bodies. Please ensure this information is included or provide appropriate references.

Terminalia bellirica Roxb. is a deciduous tree that is widely distributed in the tropical regions. The fruit extract of *T. bellirica* has shown to be hepatoprotective(17) and the main phyto constituents reported are tannins such as chebulic acid, chebulinic acid, chebulagic acid, gallic acid, corilagin, ellagic acid and flavonoids, sterols, amino acids, fructose, resin, fixed oils etc. *T. bellirica* contains different chemical constituents in different parts such as stem bark contains arjungenin and its glycosides, belleric acid, belleriosides. Ripe fruits are used as astringent in

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combination with chebulic myrobalan (*Terminalia chebula*) and *Phyllanthus emblica* as the famous Triphala drug of Ayurveda are also useful in eye problems like cataract, glaucoma, progressive myopia, and conjunctivitis.

Comment [T116]: "are used for eye diseases such as"

Secondary plants metabolites possess various compounds differ widely in terms of structure, biological properties and wide mechanisms of actions (10). Polyphenols have been also proven to lower the risk of cardiovascular diseases, enhance liver regeneration, and increase life expectancy. These secondary metabolites include flavonoids, phenols, phenolic glycosides, saponins, and glycosides (18). *Terminalia* species exhibit nutraceutical value with numerous health benefits in the treatment of some diseases (19). For example, fruits of *T. bellirica* (Gaertn.) Roxb usually form Triphala, the well-known polyherbal formulation in Ayurvedic has pharmacological applications as a laxative, detoxifying, and rejuvenating effects (20). Thus, our present study aims to analyze the phytochemical composition of *T. bellirica* leaves and their antimicrobial activity. In this study, methanol extract of leaves of *T. bellirica* is analyzed by LC-MS techniques for the phytochemicals profile and antibacterial assay is also carried.

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2. Methods and materials:

2.1. Preparation of extract:

Fresh mature plant leaves of *T. bellirica* were collected from trees growing in Orathanadu taluk, Thanjavur District, Tamilnadu. The leaves of *T. bellirica* (250 g) were air-dried, ground, and extracted with 95% methanol at room temperature for 8 hours (6 × 500 ml). The methanol extracts were filtered, and reduced under vacuum at 40 °C.

2.2. LC-MS assay:

Leave extract was analyzed by shimadzu LCMS2010 / LCMS-QP8000α and the solvents used were 0.5% (v/v) acetic acid (A) and 100% methanol (B). The isocratic elution was as follows: (i) 55% of solvent A, from 0 to 10 min, (ii) 65%, from 11 to 20 min (iii) 35%, at 21-30 min of total run time. The PDA detector (UPLC LG 500 nm) was monitored at 340 nm and the column temperature was maintained at 30 °C. The mass spectrometer (MS) was operated in the positive ionization mode with the mass range of 150 m/z to 1000 m/z, the capillary voltage of 3.50 kV, cone voltage of 30 V, extractor voltage of 3V, the gas flow of 30 L/Hr and collision gas flow of 0.18 mL/Min. The mass spectrometry (MS) was determined for phytochemicals using a shimadzu LCMS-2010 / LCMS- QP8000α plus instrument by direct injection. The experiment was carried out in NIFTEM, Thanjavur, Tamilnadu. Detection of the ions was performed in electrospray ionization (ESI) and quadrupole ion trap mass analyzer (21).

2.3. AAS Analysis:

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Leaves extract was taken in a precleaned and constantly weighed silica crucible and heated in a muffle furnace at 400C till there was no evolution of smoke. The crucible was cooled at room temperature in a desiccator and carbon-free ash was moistened with concentrated sulphuric acid and heated on a heating mantle till

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fumes of sulphuric acid ceased to evolve. The crucible with sulphated ash was then heated in a muffle furnace at 600°C till the weight of the content was constant (~2–3 h). One gram of sulphated ash obtained above was dissolved in 100 mL of 5% HCl for Atomic Absorption Spectroscopy (AAS). Standard solution of each element was prepared and calibration curves were drawn for each element using AAS (22).

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3. Results and discussion

T. bellirica has a diverse pharmacological spectrum and has been used in Ayurveda, Siddha, Chinese medicine etc, because of having important phytoconstituents (23,24,25). LC-MS analysis of *T. bellirica* leaves recorded 21 compounds and out of these chebulinic acid, chebulagic acid, quercetin and bellericoside were found to be higher in the leaves (Fig 1,2,3). Chebulinic acid is known to possess numerous biological activities including anti tumor activity, anti antherogenic, anti fibrotic, anti inflammatory, antiulcer, antioxidant, hepatoprotective, antidiabetic and antiviral etc. Chebulagic acid, ellagitannins and proantho cyanidins are the major chemical constituents present in the leaves extract of *T. bellirica*.

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Chebulagic acid is benzopyran tannin and an antioxidant that has many potential uses in medicine. It has been found to be immunosuppressive (26,27), hepatoprotective (28) and a potent alpha-glucosidase inhibitor (29,30), a human gut enzyme useful in diabetic studies has been shown to be active against *Staphylococcus aureus* and *Candida albicans* (31). It is formed from geraniin through a glutathione-mediated conversion (32). Gallo-tannic acid, bellericanin, ellagic acid, gallic acid, termilignan, thanni lignan, flavone and anolignan B, tannins, ellagic acid, ethyl gallate, galloyl glucose and chebulaginic acid, phenyllembin, sitosterol, mannitol, glucose, fructose and rhamnose were also found present in it. Therefore, because of the presence of these compounds, the leave extract showed possible many of the broad pharmacological activities such as antisecretory, analgesic, antihypertensive, antidiarrhoeal activity, antimicrobial activity, antidiabetic, antioxidant (33,34) antiulcer, antipyretic, hepatoprotective, anticancer, angiogenesis, antidepressant-like activity (35). It is useful in the treatment of gastric ulcer, constipation, general debility, piles etc. still, and it has actually not been explored comprehensively. These compounds are traditionally used in relief in a cough, asthma, indigestion, dental problems, sore throat and wounds (36).

Glucosides, tannins, gallic acid, ellagic acid, ethyl gallate, galloyl glucose, chebulanic acid are mainly responsible for its wide therapeutic actions. It has anti-HIV1, antimalarial and antifungal activity. It is used as antioxidant, antimicrobial, antidiarrheal, anticancer, antidiabetic, antihypertensive and hepatoprotective agent (37,38). It also possesses analgesic, antipyretic and anti-ulcerogenic effect and antimicrobial activity (39,40). Many phytoconstituents have been isolated from the leaves, chebulagic acid, gallic acid have antimicrobial activity of

8. REFERENCES:

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Table 1: Compounds identified in LC-MS assay

S.No	Tentatively identified compounds	m/z
1.	7- Hydroxy3,4(methylene dioxyflavan)	270
2.	Luteolin	286
3.	Beta sitosterol	414
4.	Termilignan	296
5.	Anolignan	266
6.	Gallic acid	170
7.	Ellagic acid	302
8.	Methyl Gallate	184
9.	Phyllembin	198
10.	Chebulinic acid	956
11.	Chebulagic acid	955
12.	3-0- alpha glucopyranosylL-aminopyranose	326
13.	Hexahydroxydiphenic acid	338
14.	Arjungenin	505
15.	Bellericoside	666
16.	Quercetine	302
17.	Kempherol	286
18.	Rutin	610
19.	Quercetin – 3-07-0-alpha rhamnopyranoside	756
20.	Quercetin-3-beta-glucoside	464
21.	Dihydroclerodin	436
22.	Quinic acid	191
23.	Mailc acid	133
24.	Galloyl hexoside	933
25.	puigluconin	801
26.	Grahatin	951
27.	corilagin	633
28.	Galloyl punicalagin	1235
29.	Trigaloxyl hexoside	635
30.	Elgatannin	967
31.	Punicafolin	937
32.	Quercetin coumaroyl glucoside	609
33.	Vitexin	431
34.	Chebolic acid	355
35.	Cuarictin	935
36.	Myricetin rutine	935
37.	Myricetin glucoside	479

Chart 1: Minerals Assay by AAS in leaves of *T.bellerica*

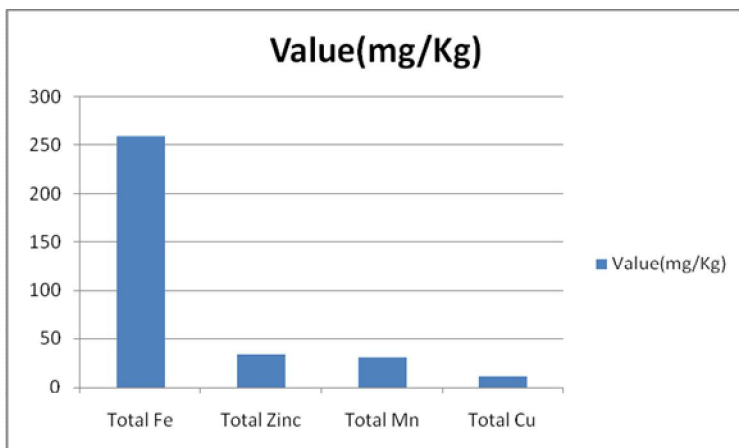


Fig 1. Chebulagicacid, C₄₁H₃₀O₂₇, Exact Mass: 954.66
MSES+

2:TOF

1.05e6

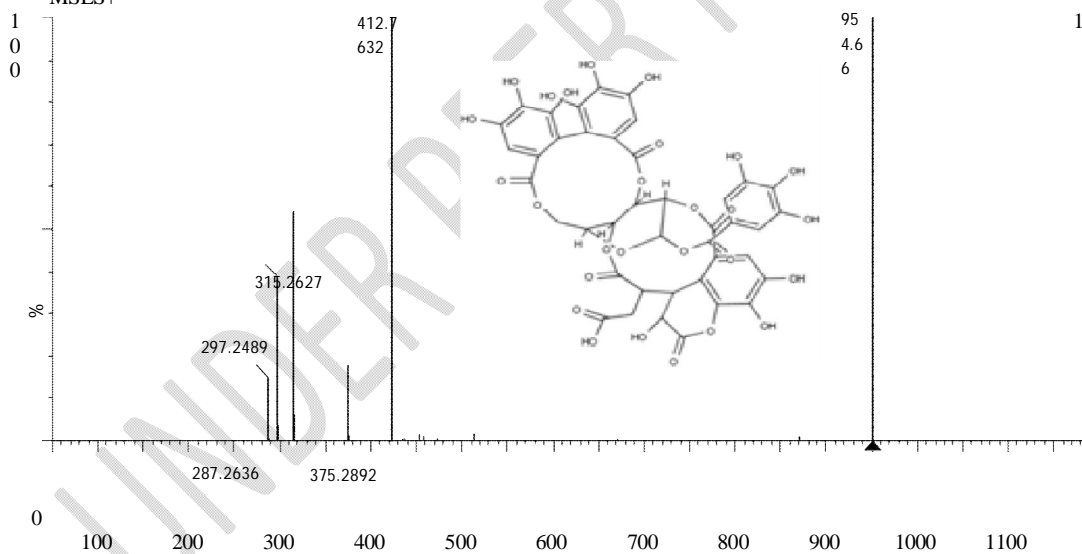


Fig 2: Bellericoside, C₃₆H₅₈O₁₁, Exact Mass: 666.8

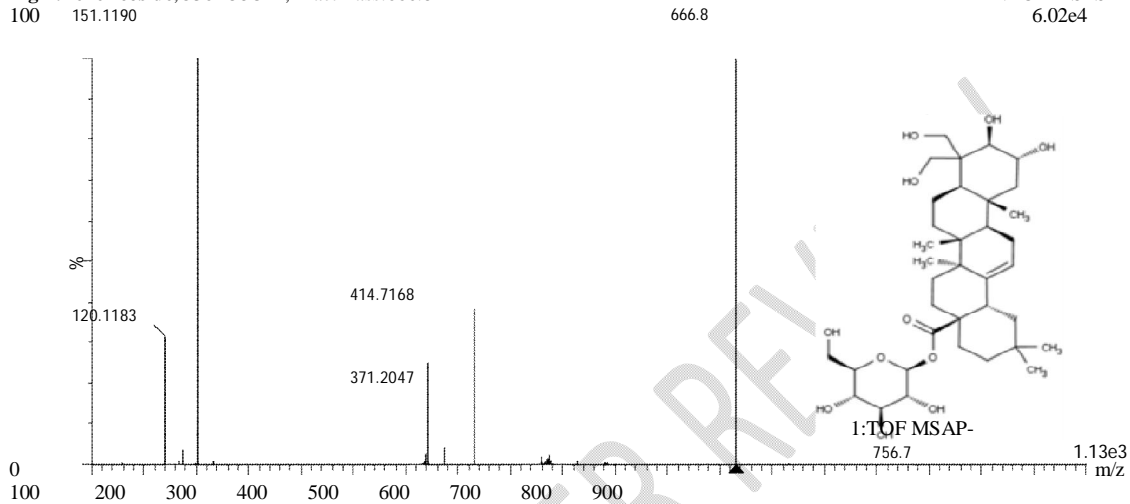
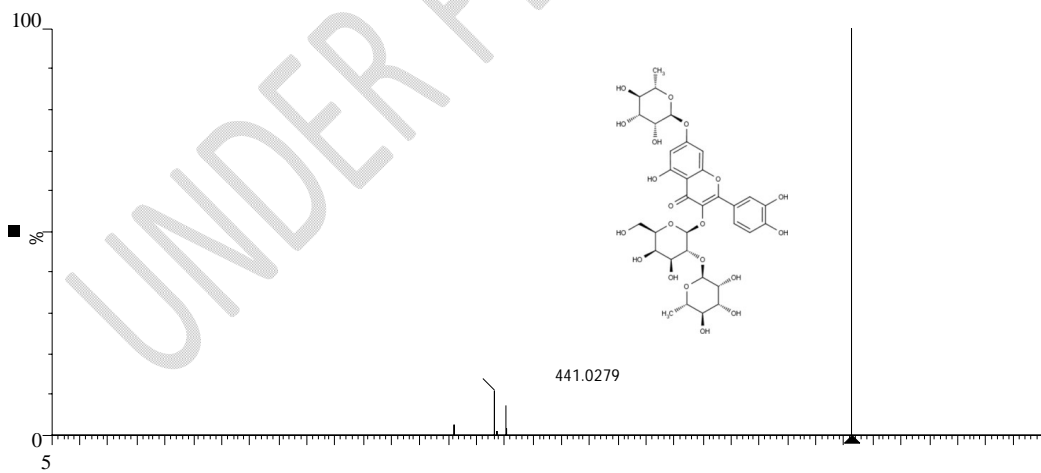


Fig 3: Quercetin-3-O-7-O- α -L-rhamnopyranoside, C₃₃H₄₀O₂₀ Exact Mass: 756.7



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