

**Nutrient Profiling of Bark of Banyan Tree
(*Ficus benghalensis* L.): Impact of
Supplementation in Type 2 Diabetic Subjects**

EXECUTIVE SUMMARY

By

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INTRODUCTION

According to the World Health Organization, non-communicable diseases (NCDs) were responsible for at least 43 million deaths in 2021, making up 75% of non-pandemic-related fatalities globally, highlighting a major public health challenge. Diabetes mellitus is one of the major NCDs and has emerged as a significant global public health concern as well as a leading cause of mortality, with its prevalence increasing at an alarming rate.

Diabetes mellitus is a chronic metabolic disorder characterized by persistently elevated blood glucose levels due to inadequate insulin production, impaired insulin function, or both. According to the International Diabetes Federation (2025), an estimated 589 million people worldwide were living with diabetes in 2024, and this number was projected to increase by 45% by 2050. India bears a significant share of this burden, ranking second among the countries with the highest number of adults (aged 20 to 79 years) with diabetes. In 2024, 89.8 million people in India were living with diabetes, and this figure is expected to rise by approximately 74% by 2050 International Diabetes Federation (2025).

Effective diabetes management requires a comprehensive approach that includes hypoglycaemic drugs, insulin therapy, dietary adjustments, regular exercise, and behavioural modifications. In addition to dietary modifications, Complementary and Alternative Medicine (CAM) is increasingly recognized for its role in diabetes management. Dietary supplements are an essential component of CAM, commonly used to support overall well-being and sustain optimal health. A systematic review and meta-analysis by Alzahrani et al., 2021 examined the global prevalence of CAM use among adults with diabetes, revealing a wide variation in usage, ranging from 8% to 89%, with a pooled prevalence of 51%. Among the various CAM therapies, herbal and nutritional supplements were the most commonly used. The demand for dietary supplements has expanded beyond vitamins and minerals to include botanicals and herbs, as consumers seek natural alternatives for disease prevention and wellness.

Among the numerous botanicals studied for their anti-diabetic potential, *Ficus benghalensis* L. is one such plant that has attracted scientific interest. It is commonly known as “banyan tree” and native to India, Pakistan, and Sri Lanka. This large evergreen tree, belonging to the Moraceae family, is characterized by its aerial roots that descend from the branches and develop into pillar-like trunks. *Ficus benghalensis* L. has long been an integral part of traditional medicine across the Indian subcontinent, valued for its therapeutic properties.

Various parts of the tree, including its bark, roots, leaves, and latex, have been traditionally utilized in the treatment of diverse ailments, reflecting its longstanding role in folk medicine.

In addition to its traditional use, scientific studies have demonstrated the therapeutic potential of *Ficus benghalensis* L. bark in the management of diabetes and its related metabolic disturbances. Experimental studies using animal models have also shown that the bark possesses hypoglycemic and hypocholesterolemic effects (Amali et al., 2025; Gayathri & Kannabiran, 2008; Khanal & Patil, 2021; Saifi et al., 2024; Shukla et al., 2004). However, its clinical relevance remains underexplored, indicating the need for further investigation to determine its potential role in diabetes management.

RATIONALE OF THE STUDY

Plant-based therapies have long been employed in traditional medicine for the management of diabetes mellitus, with *Ficus benghalensis* L. being a botanical of substantial potential, as demonstrated in in vitro and in vivo studies. However, there is a dearth of studies assessing the nutrient composition of *Ficus benghalensis* L. bark, and no clinical evidence is currently available to establish its effectiveness in human subjects. Therefore, the present study was undertaken with the broad objective of evaluating the impact of *Ficus benghalensis* L. bark supplementation in individuals with type 2 diabetes mellitus.

OBJECTIVES OF THE STUDY

- 1) To analyze the nutrient composition, phytochemical and antioxidant profile of bark of *Ficus benghalensis* L.
- 2) To study the consumption of dietary supplements among diabetes subjects for the management of diabetes mellitus.
- 3) To study the impact of bark of *Ficus benghalensis* L. supplementation on glycaemic, lipemic, liver and kidney function in stable type 2 diabetes mellitus subjects.

METHODOLOGY

Before initiating the study, ethical approval was obtained from the Institutional Ethical Committee of the Maharaja Sayajirao University of Baroda, Vadodara, India (IECHR/FCSc/PhD/2021/121), ensuring adherence to ethical guidelines and participant safety. The study was conducted in three phases:

Phase 1: Nutrient composition, phytochemical and antioxidant profile of bark of *Ficus benghalensis* L.

Phase 2: Cross-Sectional Study of Dietary Supplement Consumption Among Diabetic Subjects

Phase 3: Impact of *Ficus benghalensis* L. Bark Supplementation on Subjects with Type 2 Diabetes Mellitus

Phase 1: Nutrient Composition, Phytochemical and Antioxidant Profile of Bark of *Ficus benghalensis* L.

Phase 1 employed a laboratory-based experimental design to analyze the nutrient composition, phytochemical profile, and antioxidant profile of *Ficus benghalensis* L. bark. The bark powder was procured and authenticated to verify its botanical origin. After authentication, the sample underwent heavy metal testing to ensure compliance with safety standards. Following analysis was done using standard methods:

- **Heavy metal testing:** lead, mercury and arsenic.
- **Proximate Composition:** moisture, ash, fibre, protein, fat, carbohydrate and energy
- **Micronutrients:** iron, calcium, phosphorus, sodium, potassium, magnesium and vitamin C
- **Phytochemical analysis:** qualitative analysis of phytochemicals and quantitative estimation of total phenol, flavonoid, and tannin.
- **Antioxidant assays:** FRAP and DPPH assay

Phase 2: Cross-Sectional Study of Dietary Supplement Consumption Among Diabetic Subjects

In Phase 2, a cross-sectional study design was used to determine the prevalence and use of dietary supplements among diabetic patients in Vadodara, Gujarat, India. Four healthcare clinics providing medical care to diabetic patients were purposively selected based on patient load and willingness to participate. From these clinics, 400 diabetic subjects were recruited and interviewed. Data were collected using a pretested questionnaire designed to gather detailed information on supplement usage, the beliefs influencing its use, and physicians' opinions, offering a comprehensive understanding of the patterns and factors associated with their consumption.

Sample Size Calculation

The sample size was calculated by using the formula (Shete et al., 2020):

$$n = \frac{(Z_{(1-\alpha/2)})^2 \times (p) \times (q)}{(d)^2}$$

Where:

- $Z_{(1-\alpha/2)}$ represents the standard normal critical value corresponding to the selected confidence level (1.96 for a 95% confidence interval).
- p denotes the estimated proportion of the population, which was taken as 52% based on a previous study on supplement use among diabetic patients (Li et al., 2020).
- $q = 1 - p$
- d denotes the margin of error (precision) = 5%

Using this formula, the calculated sample size was 383.54, which was rounded up to 400 for practical feasibility. Therefore, the final sample size for Phase 2 was set at 400.

Sample Selection Criteria

Inclusion Criteria:

Individuals were eligible to participate if they:

- Had a diagnosis of diabetes mellitus.
- Were above 18 years of age.
- Resided in Vadodara city during the study.

Exclusion Criteria:

- Participants were excluded if they were pregnant at the time of enrolment.

Phase 3: Impact of *Ficus benghalensis* L. Bark Supplementation on Subjects with Type 2 Diabetes Mellitus

In the third phase, a randomized, triple-blind, placebo-controlled trial was conducted to assess the effects of *Ficus benghalensis* L. bark supplementation in individuals with Type 2 Diabetes Mellitus. A total of 80 participants diagnosed with Type 2 Diabetes Mellitus were recruited from four preselected clinics and randomly allocated into two groups of 40 each. To maintain the triple-blind nature of the study, similar tablets, with and without the active

ingredient, were prepared for both groups. These tablets were coded as Code A and Code B to ensure blinding.

Subjects in Group A received standard diabetes care along with Code A supplementation, while those in Group B received standard diabetes care along with Code B supplementation. The intervention was administered for a duration of 90 days. The coding details were securely sealed in an envelope and remained undisclosed throughout the study. These details were revealed to the researcher only after the study was completed, that is, upon the completion of data analysis.

Sample Size Calculation

The sample size was determined using the sample size table for comparing means between two groups from the book *Designing Clinical Research* (Hulley et al., 2007). The table was based on the general formula as follows:

$$N = [(1/q_1 + 1/q_2) S^2 (z_{\alpha} + z_{\beta})^2] / E^2$$

Where,

- Confidence level 95%
- Power = 80%
- SD= 45 mg/ dl
- Expecting reduction 30 mg/dl ~ 1 % HbA1C
- Effect size (E/S) = 0.66= 0.7
- n = 34
- With attrition of 10% sample size, n= 38 ~ 40 (in each group)

Sample Selection Criteria

Inclusion Criteria:

- Adults with a confirmed diagnosis of type 2 diabetes mellitus.
- Stable diabetic subjects

Exclusion Criteria:

- History of medical condition like cancer, thyroid disorders, asthma, or stroke.

- Currently taking any dietary supplements or functional foods.
- Alcoholic, smokers and/or tobacco chewers.
- Individuals undergoing insulin therapy.

Supplementation Details

- **Dose:** 3 gram per day
- **Form:** tablets
- **1 tablet:** contains 500 mg of active ingredient
- **Frequency:** six tablets per day (3 tablets after lunch and 3 after dinner)
- **Time period:** 90 days

Note: Dose was based on the recommendations of the Department of AYUSH, Ministry of Health and Family Welfare, Government of India and the Food Safety and Standards authority of India, Ministry of Health and Family Welfare, Government of India (2016).

Data Collection

- Data pertaining to socio-economic status, educational status, medical history of the subjects as well as the family history regarding any disease condition, dietary habits and addiction pattern was collected.
- Data with respect to diet pattern (3 day -24-hour dietary recall), FFQ, anthropometric measurements (BMI, WC, HC, WSR, WHR), biophysical measurements, physical activity (IPAQ) and biochemical parameters was collected using standard techniques.
- The Impact of bark of *Ficus bengalensis* L. supplementation was studied on the following biochemical parameters:
 - **Glycemic Parameters:** FBS, PP2BS and HbA1c
 - **Lipemic Parameters:** TC, TG, HDL, LDL, VLDL and Apolipoproteins
 - **Inflammatory marker:** hs- CRP
 - **Liver and kidney function test:** Microalbumin, Urinary Creatine, SGPT and SGOT

Note: The blood sample collection for analysis was done by trained technicians.

KEY FINDINGS

Phase 1: Nutrient Composition, Phytochemical and Antioxidant Profile of Bark of *Ficus benghalensis* L.

To comprehensively evaluate the bark's therapeutic potential and ensure its safety for human consumption, analyses of its nutrient composition, phytochemical constituents, antioxidant activity, and heavy metals concentration were conducted. The key findings from these assessments are summarised below.

Heavy Metals Testing

Ficus benghalensis L. bark powder was screened for heavy metals to ensure the safety of the product. The levels of heavy metals such as lead, mercury, and arsenic were reported to be less than 0.1 mg/Kg of the sample which is within the permissible limits.

Estimation of Nutrient Composition

- The analysis revealed that the bark is rich in carbohydrates (77.86%), energy (360 Kcal), crude fiber (21.86%), and ash (6.29%), with lower levels of moisture (7.58%), protein (5.14%), and fat content (3.13%).
- the bark was found to be rich in minerals, with the highest concentration of potassium (1417 mg/100 gm), followed by phosphorus (1370 mg/100 gm), calcium (472 mg/100 gm), iron (147 mg/100 gm), and magnesium (96.7 mg/100 gm). Sodium (12.6 mg/100 gm) and vitamin C (7.56 mg/100 gm) are present in comparatively lower concentrations.

Phytochemical Analysis

- Qualitative analysis confirmed the presence of tannins, phenols, saponins, glycosides, flavonoids, terpenoids, carbohydrates, and amino acids in the sample.
- The Concentration of total phenols, flavonoid and tannins in the bark were found to be 244.736 mg per 100gram, 332 mg of rutin equivalent per gram and 70.3 µg catechin equivalents per gram, respectively.

Antioxidant Analysis

- The DPPH assay, using ascorbic acid as the standard, showed 75.37% scavenging activity for the bark.

- In addition, the FRAP assay revealed an antioxidant capacity of 0.461 μM ferric equivalents per gram of the bark, demonstrating its ability to reduce ferric ions (Fe^{3+}) to ferrous ions (Fe^{2+}).

Phase 2: Cross-sectional Study on Dietary Supplement Consumption among Individuals with Diabetes Mellitus

As supplement consumption is often influenced by personal perceptions, cultural beliefs, and traditional practices, region-specific data is essential to understand local usage patterns, underlying motivations, and the types of supplements consumed, which may vary across populations. Therefore, this section summarises the results of the cross-sectional study conducted in Vadodara, Gujarat, to determine the prevalence and determinants of dietary supplement use among individuals with diabetes mellitus.

- Data was collected from 400 adult diabetic subjects. The mean age was 60.59 ± 11.44 years, with 55.25% male and 44.75% female participants, and most (90.25%) identified as Hindu. Among the participants, 32.25% were graduates, 87.25% were married, and 60.50% lived in nuclear families. Homemakers were the largest occupational group (38.50%), followed by retirees (29%) and those employed (19.50%).
- According to the Modified Kuppuswamy Socioeconomic Scale (Saleem and Jan, 2021), 50% of participants belonged to the upper middle class, followed by 21.71% in the upper lower class.
- It was found that the average duration of diabetes among subjects was 11.03 ± 9.24 years (mean \pm SD) and average age at diabetes onset was 49.56 ± 11.27 years (mean \pm SD). Most subjects (73.25%) had at least one comorbidity along with diabetes mellitus. Hypertension was the most prevalent comorbidity, affecting 59.75% of the subjects, followed by hyperlipidemia (30.25%), thyroid disorders (13.50%), and coronary heart disease (11.75%).
- The data reveals that 62% of participants had used dietary supplements at least once since their diabetes diagnosis, but only 36.75% were actively using them at the time of the study, indicating a 40.72% had discontinued use.
- The association between various respondent characteristics and dietary supplement consumption was analyzed using the Chi-square test and the Mann-Whitney test. Results revealed that supplement usage was significantly higher among respondents aged 60-79 years (52.02%, $p=0.02$), those with a family history of diabetes (66.53%, $p=0.0056$), and those with a family history of chronic diseases (79.84%, $p=0.0027$). Additionally, the

average duration of diabetes was significantly longer among participants who had used dietary supplements (11.76 ± 9.20 years) compared to non-users ($p = 0.0074$).

- The most commonly used supplement was fenugreek seeds (*Trigonella foenum-graecum*), reported by 41.13% of participants. Other frequently used supplements included commercial polyherbal formulations ($\approx 20\%$), Ayurvedic powders of unspecified composition (17.34%), self-prepared herbal mixes (14.11%), and giloy (*Tinospora cordifolia*, 12.10%). Additional supplements reported were juice mixes, Indian blackberry seeds (*Syzygium cumini*), bitter gourd (*Momordica charantia*), neem (*Azadirachta indica*), Paneer doda (*Withania coagulans*), turmeric (*Curcuma longa*), amla juice (*Phyllanthus emblica*), and various Ayurvedic and homeopathic preparations.
- Apart from diabetes and its complications, participants also reported using these supplements for general health, immunity, body pain, bowel health, hypertension, renal health, dyslipidemia, weight loss, flatulence, COVID-19, and other conditions.
- The consumption of dietary supplements by participants was driven by various beliefs and motives. The three most common reasons were influence from others (43.95%), the belief that Ayurveda and natural products have fewer or no side effects (19.35%), and the goal of reducing or discontinuing allopathic medications (13.31%).
- The findings indicated that most participants (68.55%) were influenced by family and friends, followed by mass media (17.34%), while only 14.92% relied on prescriptions from Ayurvedic, homeopathic, or naturopathic practitioners. Social media (11.69%) and internet searches (7.66%) also contributed to participants' awareness of supplements.
- The survey findings indicate that 40.72% of respondents eventually discontinued supplement use due to various reasons. The most common reasons for discontinuation were inconvenience or forgetfulness, perceived ineffectiveness, side effects, effective control of symptoms or health indicators, inaccessibility, and unpleasant taste.
- The findings indicate that nearly two-thirds of participants did not disclose their dietary supplement use to their doctors.

Phase 3: Impact of *Ficus benghalensis* L. Bark Supplementation on Subjects with Type 2 Diabetes Mellitus

This phase provides the first clinical evidence evaluating the antidiabetic efficacy of *Ficus benghalensis* L. bark in humans. A randomized, triple-blind, placebo-controlled trial was conducted, in which adult Type 2 diabetic participants received 3 g of bark powder in tablet form daily for 90 days. The key findings of the trial are presented below.

- The mean age of subjects in the experimental group was 50.95 ± 6.89 years, while that of the placebo group was 56.23 ± 7.91 years. The experimental group consisted of 23 males and 17 females, whereas the placebo group included 19 males and 21 females.
- Across both groups, the majority of subjects were Hindu, had graduate-level education, were married, and worked in services or business; most were also classified as upper middle class according to the Modified Kuppaswamy Socioeconomic Scale (Radhakrishnan and Nagaraja, 2023).
- The study subjects had an average diabetes duration of 7.22 ± 5.81 years, with the mean age at onset reported as 46.37 ± 7.36 years, suggesting that the majority developed the condition during mid-adulthood.
- Self-reported comorbidities were prevalent among the subjects, with 50% of the experimental group and 67.5% of the placebo group reporting at least one additional condition. Hypertension was the most frequently reported, followed by hyperlipidemia.
- A predominantly vegetarian diet was observed across both groups. All participants reported daily milk consumption, mostly full cream- 68.42% in the experimental group and 56.76% in the placebo group. Average daily sugar/jaggery intake was 7.7 ± 7.81 g and 5.74 ± 6.61 g, respectively, and preference for artificial sweeteners was low in both groups.
- Regarding edible oil consumption, most participants reported using the same type of oil year-round, with cottonseed and groundnut oils being the most commonly used.
- The mean percentage distribution of food frequency indicated that cereals and legumes/pulses were the most frequently consumed food groups across both groups. Millet consumption was comparatively low, with only 12.41% of the experimental group and 8.11% of the placebo group reporting intake at least once per week. The most commonly consumed millets were sorghum (jowar), pearl millet (bajra), and kodo millet (kodri).
- A seasonal consumption pattern was observed for vegetables and fruits. Roots and tubers were consumed weekly by 49-62% of participants, while nearly 22% reported weekly intake of nuts and oilseeds. Daily consumption of ghee or butter ranged from 39-65%, and sugar or jaggery from 37-46%. Ready-to-eat foods were consumed weekly by 19-24% of participants.
- The percentage of the Estimated Average Requirement (EAR) or Upper Limit (UL) met by subjects at pre- and post-intervention was determined according to the ICMR-NIN 2024 RDA and EAR guidelines. In both groups, protein intake was close to the EAR, indicating adequacy, while energy intake was only 55-67% of the EAR, suggesting a potential energy

deficit. Visible fat intake exceeded recommendations, and most minerals were consumed below recommended levels, although iron intake was relatively adequate, particularly in males. Vitamin C intake exceeded the EAR in both groups, whereas vitamin A was markedly inadequate in the experimental group ($\leq 50\%$ of the EAR) but substantially higher in the placebo group, exceeding 75% in males and 100% in females.

- In the experimental group, no significant changes in nutrient intake were observed post-intervention ($p > 0.05$), whereas the placebo group exhibited a significant increase in total saturated fatty acids and added sugars ($p < 0.05$).
- Based on the IPAQ, the majority of subjects in both groups reported engaging in moderate-intensity physical activity. Physical activity remained unchanged during the intervention, with no significant differences observed in MET minutes and sitting hours post-intervention.
- Most subjects in both groups were classified as obese grade I (BMI 25–29.9) and exhibited a high prevalence of central obesity. Over 85% of males and females exceeded waist circumference thresholds (≥ 90 cm for males, ≥ 80 cm for females), and waist-to-hip ratios surpassed 0.90 in males and 0.85 in females in all experimental subjects, and in nearly 90% of the placebo group. Statistical analysis using ANOVA showed no significant changes in any anthropometric indices within either group across measurements recorded on days 0, 45, and 90.
- The baseline analysis revealed that subjects with elevated HbA1c levels had significantly higher diastolic blood pressure, TC, LDL-C, TC/HDL ratio, and Apo B, along with a lower Apo A1/B ratio compared to those with lower HbA1c levels.
- Correlation and regression analyses demonstrated interrelationships between glycemic status, lipid parameters, renal function, abdominal obesity, and micronutrient intake in type 2 diabetic subjects. Elevated fasting glucose levels were associated with adverse lipid profiles and an increased atherogenic risk.
- Higher central adiposity was associated with elevated triglyceride levels and reduced HDL-C.
- Age at diabetes diagnosis was significantly negatively correlated with HbA1c and PP2BS, indicating that earlier onset is associated with poorer glycemic control.
- Higher serum creatinine levels were associated with lower HDL-C and apo A1, and higher TG, suggesting that impaired renal function contributes to a more atherogenic lipid profile.

- Higher intake of dietary fiber, magnesium, copper, and zinc was associated with improved lipid profiles- lower TC, LDL-C, and apo B- suggesting a protective role of these nutrients against dyslipidemia in diabetes mellitus.
- Overall, supplementation did not result in significant changes in blood pressure across groups. However, in participants with elevated SBP (≥ 140 mmHg), a significant reduction of 8.11% in SBP was observed, while those with elevated DBP (≥ 90 mmHg) showed significant reductions of 6.29% in SBP and 8.29% in DBP.
- Although not statistically significant, the experimental group showed greater reductions in FBS (7.8%), PP2BS (4.02%), and HbA1c (2.79%) compared to the control group. Additionally, the experimental group exhibited a modest but consistent decline in FBS and PP2BS, whereas the placebo group showed variable changes during supplementation.
- Subgroup analysis revealed that participants in the experimental group with elevated baseline FBS (≥ 145 mg/dL) showed a significant 9.24% reduction in HbA1c, accompanied by notable, though non-significant, reductions in FBS (16.36%) and PP2BS (14.63%).
- Overall, supplementation had no significant effect on lipid and inflammatory markers, except for a 7.62% increase in apo A1 in the experimental group. Although not significant, hs-CRP decreased by 17.93% in the experimental group, compared to an 18.04% increase in the placebo group.
- Subgroup analysis revealed that in the experimental group participants with elevated baseline TC (≥ 170 mg/dL) or LDL-C (≥ 100 mg/dL) showed significant reductions in TC (4–5.3%) and LDL-C (5.55-7.24%), along with increases in apo A1 (10–12.93%) and the apo A1/B ratio (12.9-16.13%). Those with elevated triglycerides (≥ 150 mg/dL) demonstrated significant increases in HDL-C (10.32%) and apo A1 (10.06%).
- No significant adverse effects on liver or renal markers were observed post-supplementation. The experimental group showed non-significant reductions in SGOT (21.28%), SGPT (24.91%), and the urinary albumin-to-creatinine ratio (20.81%), indicating a potential trend toward improved liver and kidney function.

CONCLUSION

The study demonstrated that the bark possesses a high content of fiber, phosphorus, and potassium, while exhibiting comparatively low sodium levels. Its bioactive compounds, including flavonoids, tannins, and phenols, confer notable antioxidant and therapeutic properties, as confirmed by strong DPPH radical scavenging activity. Moreover, the cross-

sectional phase provided a comprehensive analysis of dietary supplement usage among diabetic subjects, exploring key influencing factors. Although supplement use was highly prevalent, disclosure to healthcare providers remained notably low. This gap in communication between patients and physicians raises concerns about unmonitored supplement use. The study identified a wide range of herbs and botanicals commonly used by the subjects, including fenugreek seeds, giloy, Indian blackberry seeds, paneer doda, polyherbal formulations, Ayurvedic powders, herbal mixes, and others.

Furthermore, the clinical trial showed that supplementation with *Ficus benghalensis* L. bark at 3 g/day for 90 days resulted in a modest but consistent reduction in glycaemic parameters compared to the placebo group. Notably, the supplementation yielded more pronounced and statistically significant improvements in subjects with uncontrolled glycemia, dyslipidemia and elevated blood pressure, indicating that bark supplementation was beneficial for individuals with higher baseline metabolic risk. Additionally, the supplementation showed no signs of toxicity or adverse effects, while also improving hepatic and renal function parameters, indicating its safety and potential therapeutic benefits.

The study demonstrates that *Ficus benghalensis* L. bark supplementation is safe for human use, with no observed adverse effects on hepatic or renal function, and suggests its potential role as an adjunct in diabetes management. These results help bridge preclinical findings with clinical application and support its consideration as an evidence-based complementary approach in diabetes care.

RECOMMENDATIONS

- ❖ The dietary supplements reported by study participants highlight new research opportunities, underscoring the need for well-designed clinical investigations to rigorously evaluate their efficacy, safety, and therapeutic potential in diabetes management.
- ❖ The implementation of educational programmes through digital and social media platforms is recommended to improve patient awareness of evidence-based supplement use and to promote appropriate disclosure and monitoring.
- ❖ Comprehensive and well-structured intervention studies are needed to elucidate the causal impact of magnesium, copper, and zinc-rich foods on lipid profiles and cardiovascular health in individuals with diabetes.

- ❖ To strengthen the clinical evidence, prospective long-term studies are essential to determine the sustained impact of *Ficus benghalensis* L. bark on glycaemic control and other relevant biochemical parameters in patients with diabetes.
- ❖ Future studies should focus on dose optimization and the evaluation of various *Ficus benghalensis* L. bark formulations, such as decoctions, standardized extracts, and isolated bioactive compounds, to identify the optimal formulation and dosage for safe and effective diabetes management.
- ❖ Future studies may investigate the synergistic effects of combining the bark with other antidiabetic herbs or supplements to enhance therapeutic efficacy.
- ❖ *Ficus benghalensis* L. bark has potential for development into therapeutic products, including herbal teas, functional beverages, health bars, and other nutraceutical formulations, which may serve as complementary interventions for diabetes management.
- ❖ Future research may consider conducting clinical studies to investigate the therapeutic potential of *Ficus benghalensis* L. bark beyond diabetes, assessing its efficacy in the management of other metabolic, cardiovascular or inflammatory disorders.

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