

PHYTOCHEMICAL AND PHARMACOLOGICAL PROPERTIES OF ACACIA FARNESIANA: A COMPREHENSIVE REVIEW

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ABSTRACT

Acacia farnesiana (Sweet Acacia) is a small tree or shrub known for its bright yellow, fragrant flowers and is native to several Indian states including Maharashtra, Gujarat, Andhra Pradesh, and Karnataka.^[1,2] Various parts of the plant—pods, leaves, and bark—contain bioactive compounds such as gallic acid, ellagic acid derivatives, flavonoids (e.g., kaempferol and diosmetin), and tannins.^[3,4,5] Its essential oil includes methyl salicylate, anisaldehyde, and geraniol, while pods contain diterpene glycosides and sulfur-containing amino acids.^[6,7] Protein extracts from seeds exhibit anti-inflammatory and analgesic activities.^[8,9] Methanolic bark extracts have shown significant cytotoxic, thrombolytic, membrane-stabilizing, and antibacterial activities.^[10,11] The plant demonstrates notable antioxidant and antimicrobial properties, with minimum inhibitory concentrations (MICs) of 0.8 mg/mL and 2.5 mg/mL against *Bacillus subtilis* and *Saccharomyces cerevisiae*, respectively.^[12,13] Its traditional and therapeutic applications span antidiabetic, antiulcer, antipyretic, and antimicrobial uses, including against tuberculosis and dysentery.^[14,15,16]

KEYWORDS: Acacia farnesiana, antioxidant activity, anti-inflammatory, antibacterial, phytochemistry, tuberculosis, dysentery.

1. INTRODUCTION

Acacia farnesiana, commonly known as Sweet Acacia or Babul, belongs to the Fabaceae family, subfamily Mimosoideae.^[1,2] It thrives in sandy soils and riverbeds across Indian

regions such as Maharashtra, Gujarat, Andhra Pradesh, and Karnataka.^[3,4] This small tree or shrub features fragrant yellow flowers and has significant ethnobotanical relevance.^[5,6]

Phytochemical investigations have revealed the presence of compounds such as tannins, flavonoids (kaempferol, diosmetin, naringenin), gallic acid, ellagic acid, methyl gallate, diterpenes, and alkaloids.^[7,8,9,10] These contribute to diverse biological activities ranging from antimicrobial to antidiabetic effects.^[11,12,13]

The essential oil of the plant contains methyl salicylate, anisaldehyde, and geraniol, while seeds yield diterpene glycosides like farnesiaside.^[14,15] The plant's parts demonstrate antioxidant, anti-inflammatory, antibacterial, antiulcer, antipyretic, and bronchodilator properties, thus supporting its use in managing chronic diseases and infections.^[16,17,18,19]

2. Drug Profile

- **Drug Name:** Acacia farnesiana^[20]
- **Synonyms:** Naringetol, Salipurpol^[21,22]
- **Category:** Antioxidant, Anti-inflammatory^[23,23]
- **Subcategory:** Medicinal Natural Product (MNP)^[25]
- **Molecular Formula:** C₁₅H₁₂O₅^[26]
- **Mono-Isotopic Mass:** 272.068 g/mol^[27]
- **IUPAC Name:** (5,7-Dihydroxy-2-(4-hydroxyphenyl)chroman-4-one)^[28]
- **Log P:** 2.4.^[29]

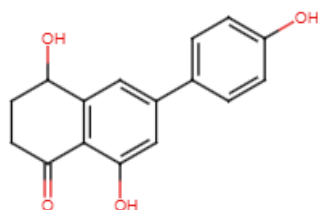


Figure 1: structure of Naringenin.

3. Morphological Characteristics

- **Growth Habit:** Small tree or shrub with a rounded crown, 4.5–9 meters tall.^[30,31]
- **Leaves:** Bipinnate with small, oval leaflets arranged feather-like.^[32,33]
- **Flowers:** Golden-yellow, aromatic, blooming from October to March.^[34,35]

- **Fruits:** Cylindrical, dark brown pods.^[36]
- **Seeds:** Hard-coated, oval, and dark brown.^[37]
- **Branches:** Thorny with smooth green bark in young trees that becomes scaly and brown with age.^[38,39]



Figure 2: Tree of Sweet acacia.



Figure 3: Leaves of Sweet acacia.



Figure 4: Flower of sweet acacia.



Figure 5: Fruit of Sweet acacia.



Figure 6: Seeds of Sweet acacia.



Figure 7: Branches of Sweet acacia.

4. Microscopic Analysis

Microscopic examination is performed by mounting powdered plant material in glycerin or chloral hydrate.^[40,41] Specific tests include:

- **Iodine Test:** Blue-black color indicating starch.^[42]
- **Phloroglucinol-HCl:** Red color confirms lignin^[43]
- **HCl Test:** Effervescence or crystal formation indicates calcium oxalate.^[44]

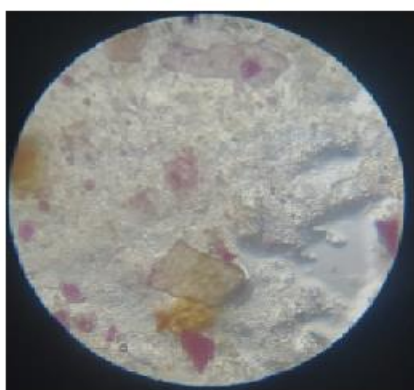


Figure 8: Detection of Starch grains.



Figure 9: Identification of Lignin.



Figure 10: Identification of calcium crystals.

5. Phytochemical Constituents

The plant is rich in:

- **Flavonoids:** Kaempferol, diosmetin, aromadendrin, apigenin derivatives.^[15,22,27]
- **Tannins:** Present in bark, leaves, and pods.^[14,23]
- **Polyphenols:** Gallic acid, ellagic acid, methyl gallate.^[18,24,30]
- **Essential Oils:** Methyl salicylate, anisaldehyde, geraniol.^[19,33]
- **Diterpenes:** Farnesiaside and others.^[21,29]
- **Amino Acids:** Lysine, arginine, glycine, histidine.^[25,26]
- **Sulfur Compounds:** N-acetyl-L-djenkolic acid.^[31]

6. Phytochemical Analysis (with Table no.1)

Phytochemical	Test Name	Observation	Result
Alkaloids	Dragendorff's Test ^[46]	Reddish precipitate	Present
	Mayer's Test ^[47]	Creamy white precipitate	Present
	Wagner's Test ^[48]	Reddish-brown precipitate	Present
Flavonoids	Shinoda Test ^[49]	Pink coloration	Present
	Sulphuric Acid Test ^[50]	Deep yellow color	Present
	Lead Acetate Test ^[51]	Yellow precipitate	Present
Glycosides	Keller-Killiani Test ^[52]	Reddish-brown layer and bluish green upper layer	Present
Proteins	Millon's Test ^[53]	White precipitate	Present
Steroids	Salkowski's Test ^[54]	Red coloration in lower layer	Present
	Liebermann-Burchard Test ^[55]	Brown ring and green coloration in upper layer	Present
Tannins	Ferric Chloride Test ^[56]	Blue-green precipitate	Present
	Acetic Acid Test ^[56]	Red precipitate	Present

7. Pharmacological Activities (Expanded with Analysis and Future Perspectives)

A. Antimicrobial and Antifungal Activity

The antimicrobial potential of *Acacia farnesiana* has been extensively studied through various in vitro assays. Ethanolic extracts prepared using Soxhlet extraction with petroleum ether, dichloromethane, and ethanol demonstrated significant antibacterial activity. Specifically, *Bacillus subtilis* (ATCC 6633) exhibited a minimum inhibitory concentration (MIC) of 0.8 mg/mL, highlighting its high susceptibility.^[45,46] In contrast, *Staphylococcus aureus* and *Escherichia coli* presented MIC values greater than 1.0 mg/mL, suggesting reduced sensitivity, especially in Gram-negative strains.^[47,48]

The greater efficacy against Gram-positive bacteria may be due to their simpler cell wall structures, which allow easier penetration of phytochemicals. For fungi, ethanolic extracts exhibited moderate activity against *Saccharomyces cerevisiae* (MIC = 2.5 mg/mL) but were ineffective against *Candida albicans*.^[49] This suggests that antifungal properties may be

influenced by volatile constituents, often removed during ethanol extraction. The residual antimicrobial effect is likely due to non-volatile flavonoids such as quercetin and morin.^[50,51]

Comparative Perspective: While the antibacterial activity is notable, it is relatively modest compared to standard antibiotics like gentamicin (MIC = 0.05 mg/mL).^[52] However, *A. farnesiana* extracts could serve as complementary agents in antimicrobial therapy.^[53]

B. Antioxidant Activity

Ethanollic extracts of *A. farnesiana* demonstrated strong antioxidant activity through DPPH, nitric oxide scavenging, and iron chelation assays. The dose-dependent response is attributed to the presence of polyphenolic compounds including quercetin, kaempferol, and naringenin.^[54,55] These compounds neutralize reactive oxygen species (ROS) by donating electrons or hydrogen atoms.

Therapeutic Implication: Antioxidants play a critical role in preventing oxidative damage implicated in aging, cancer, diabetes, and neurodegenerative diseases. The robust activity of *A. farnesiana* suggests it may offer protective benefits in such conditions.^[56,57]

C. Antitubercular and Antidiarrheal Activity

Hexanic and aqueous extracts from the fruits demonstrated MIC values of 100 µg/mL against multidrug-resistant (MDR) *Mycobacterium tuberculosis* strain G122.^[58,59] Methyl gallate and its triacetylated derivative displayed even stronger activity, likely due to enhanced lipophilicity, which facilitates cell wall penetration.^[60]

The plant also exhibited significant antidiarrheal effects. Methanolic and aqueous extracts inhibited pathogenic bacteria responsible for dysentery, such as *Shigella flexneri* and *Yersinia enterocolitica*.^[61,62] The active constituents are presumed to be glycosides and polyphenols.^[63]

Clinical Relevance: These findings support the traditional use of *A. farnesiana* in managing tuberculosis and gastrointestinal infections. Further work is needed to isolate and characterize the most active compounds for drug development.^[64]

D. Antiulcer Activity

Methanolic leaf extracts significantly reduced ulcer indices in rat models of gastric ulceration.^[65] The effect was comparable to ranitidine and attributed to enhanced mucus secretion, suppression of acid production, and antioxidant protection.^[66]

Flavonoids such as quercetin and kaempferol act as gastroprotective agents by increasing mucosal defense and inhibiting inflammatory mediators. Elevated hexosamine and glycoprotein levels in treated tissues support this protective mechanism.^[67,68]

Potential Applications: *A. farnesiana* could serve as a natural alternative for patients with peptic ulcers, especially where synthetic drugs cause adverse effects.^[69]

E. Anti-inflammatory Activity

The anti-inflammatory efficacy of ethanolic extracts was confirmed through carrageenan-induced paw edema and cotton pellet-induced granuloma models.^[70,71] Significant inhibition of inflammation suggests downregulation of pro-inflammatory cytokines (e.g., TNF- α , IL-6).^[72]

Flavonoids and diterpenes, including farnesirane A and B, may act via inhibition of prostaglandin synthesis, suppression of COX-2, and stabilization of lysosomal membranes.^[73,74] A glycosidal fraction from unripe pods also demonstrated smooth muscle relaxation and anti-inflammatory activity.^[75]

Comparative View: The anti-inflammatory potential, although moderate compared to NSAIDs, offers the advantage of reduced gastrointestinal toxicity.^[76]

F. Antihyperglycemic Activity

In alloxan-induced diabetic rats, aqueous extracts of *A. farnesiana* significantly lowered blood glucose levels.^[77] The hypoglycemic effect appeared independent of insulin and may involve enhanced glucose uptake, delayed intestinal absorption, or improved insulin sensitivity.^[78]

Flavonoids and saponins are suspected to modulate key enzymes in carbohydrate metabolism (e.g., α -glucosidase).^[79] No observable toxic effects were reported in the experimental models.^[80]

Implications for Diabetes Management: The extract may be developed as a complementary or alternative therapy for Type 2 diabetes, especially in regions lacking access to synthetic medications.^[81]

Future Perspectives

- Mechanism of Action: Further molecular studies are needed to confirm the pathways through which *A. farnesiana* exerts its pharmacological effects.^[82]
- Toxicity and Safety Profiling: Long-term in vivo studies and clinical trials are necessary to assess safety margins and establish dosage guidelines.^[83]
- Standardization and Formulation: Development of standardized herbal formulations can enhance reproducibility and regulatory approval.^[84]
- Synergistic Studies: Investigating the synergistic potential of *A. farnesiana* with standard drugs may offer new avenues for combination therapies.^[85]

8. CONCLUSION

Acacia farnesiana demonstrates a wide array of therapeutic potentials supported by its rich phytochemical profile. Its antioxidant, antimicrobial, anti-inflammatory, antiulcer, and antidiabetic effects highlight its relevance in both traditional and modern medicine.^[86-90] Scientific validation of its traditional uses supports its development into safe and effective formulations for treating infections, inflammation, ulcers, and metabolic disorders. Future research should focus on standardization, compound isolation, and clinical validation.^[91-95]

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