

## Therapeutic Potentials of *Ocimum sanctum* Linn. (Tulsi): A Comprehensive Review of Its Pharmacological Activities

Akanksha Patel<sup>1</sup>  and Manmath Purohit<sup>2</sup> 

<sup>1</sup>National Institute of Technology Raipur, India. Email: [akptl135@gmail.com](mailto:akptl135@gmail.com)

<sup>2</sup>Gayatri college of pharmacy, India. Email: [mpurohit0104@gmail.com](mailto:mpurohit0104@gmail.com)

| Article Info  | ABSTRACT  |
|---|---|
| <p><b>Article type:</b><br/>Review Article</p> <p><b>Article History:</b><br/>Received: 21 Jul 2023<br/>Received: 07 March 2024<br/>Accepted: 01 Dec 2024<br/>Published Online: 16 Sep 2024</p> <p>✉ <b>Correspondence to:</b><br/>Mohammad Amrollahi-Sharifabadi</p> <p><b>Email:</b><br/><a href="mailto:akptl135@gmail.com">akptl135@gmail.com</a></p> | <p>This review summarizes the diverse medicinal properties of <i>Ocimum sanctum</i> Linn. (Tulsi), a revered herb in Indian medicine. Traditionally praised for its pungency, warmth, and diverse medicinal uses, Tulsi's roots, leaves, and seeds each offer unique benefits. Ayurvedic texts categorize Tulsi as a stimulant, aromatic, and antipyretic, capable of balancing certain bodily energies while potentially aggravating others. Its primary benefits include relieving coughs, promoting sweating, and easing indigestion and appetite loss. More importantly, Tulsi demonstrates a remarkable range of biological activities. It exhibits antibacterial, antiviral, antifungal, and antiparasitic properties, making it effective against various pathogens and even malaria. Additionally, its analgesic, antipyretic, anti-inflammatory, and anti-allergic properties offer relief from pain, fever, inflammation, and allergies. Tulsi also offers cardiovascular benefits, lowers blood pressure, and protects the heart. Beyond the physical, it shows promise in improving memory, regulating cholesterol, protecting the liver, and controlling diabetes and asthma. Notably, it exhibits antioxidant and anticancer properties, suggesting potential cancer prevention and radioprotection. Furthermore, Tulsi boasts immunomodulatory, antifertility, anti-ulcer, and anti-arthritic activities, highlighting its versatility. Its adaptogenic/antistress capabilities and potential to prevent cataracts and even skin conditions like leucoderma further showcase its therapeutic breadth. Given its vast and seemingly valuable medicinal properties, this review aims to equip researchers and clinicians with the knowledge necessary to properly utilize Tulsi in their endeavors.</p> <p><b>Keywords:</b> Medicinal properties, <i>Ocimum sanctum</i> (OS, Tulsi), Pharmacological activities.</p> |
| <p>➤ <b>How to cite this paper</b><br/>Patel A, Purohit M. Therapeutic Potentials of <i>Ocimum sanctum</i> Linn. (Tulsi): A Comprehensive Review of Its Pharmacological Activities. <i>Plant Biotechnology Persa</i> 2024; 6(2): 51-67.</p>   |   |

### Introduction

Globally various plants have been used for the treatment of several diseases or their healing goals. In the Indian system of medicine various kinds of *Ocimum tenuiflorum* were preferred and apt as pharmaceutical constructions. According to Ayurveda *Ocimum tenuiflorum* is known as: "Mother medicine of nature", "The incomparable one", "Queen of Herbs", "Elixir of Life", etc. . *O. tenuiflorum* is commonly called "Tulsi" in Hindi and "Holy Basil" in English . Tulsi is an important sign of Hindu intellectual traditions. Among all of the other herbs, Tulsi is highly used in Ayurveda . To cure various modern-day illnesses

Tulsi is the best Ayurvedic wisdom tonic for the mind and body. It plays a major role in the traditional "Ayurveda" and "Unani" systems.<sup>1</sup>

*Ocimum sanctum*, transcends its botanical classification to occupy a unique space in Indian culture and medicine. This aromatic shrub, belonging to the Lamiaceae family, thrives in tropical regions and is readily found in courtyards and temples throughout India. <sup>2</sup>

Beyond its cultural reverence, Tulsi has a long history of use in traditional Ayurvedic medicine. Leaves, flowers, and seeds have been employed for centuries to address various ailments, including respiratory issues like bronchitis and cough, digestive disorders, fevers, wound healing, and even stress and anxiety. While these traditional uses were largely anecdotal, recent scientific research is starting to shed light on the potential mechanisms behind Tulsi's purported effects.

Studies reveal a plethora of bioactive compounds within Tulsi, including essential oils rich in eugenol, linalool, and thymol. These compounds exhibit demonstrated antimicrobial, anti-inflammatory, and antioxidant properties. In vitro and animal studies suggest potential benefits for respiratory functions, immune modulation, wound healing, and stress reduction. However, clinical trials in humans are still limited, and further research is required to establish efficacy and safety for specific conditions.<sup>3</sup>

**Table 1:** The Description about Important Species of Ocimum.

| S. No. | Species  | Geographical Distribution   | Biochemical Constituents  | General, Traditional, and Modern Uses  |
|--------|--|---|---|--|
| 1.     | <i>Ocimum tenuiflorum</i> L. or <i>Ocimum sanctum</i> L. | India, Andaman And Nicobar Islands, Nepal, Bangladesh, China, USA, Kenya, Nigeria,                    | Eugenol, Methyl Eugenol, apigenin-7-O-glucuronide, carvacrol, circimaritin, caryophyllen, cirsilineol, isothymusin, pinene, molludistin, rosamic acid, orientin, vicenin, Ursolic acid, luteolin, sesquiterpine hydrocarbon, luteolin-7-Oglucuronide, apigenin, camphor | Antistress, antimicrobial, antioxidant, antilarval, antiviral and antifungal. Antiasthmatic, anti-inflammatory, analgesic, hypotensive and hypoglycemic activity. Treatment for snakebite and scorpion-stings. Radiation protection. To treat cough, cold, sore throat, fever, headache, dysentery, ulcer and skin diseases. Treatment against HIV. Anti-dibetic, anticancer, hepatoprotective and hypolipidemic activity. |
| 2.     | <i>Ocimum americanum</i>                                 | Tropical Africa, South East Asia, Indian Subcontinent, Sri Lanka, China, Madagascar, Burma, Thailand, | $\alpha$ -pinene, Isoborneol, Linalool, Limonene, linalooloxide, Humulene, $\alpha$ bisabolene, Camphor, Anethole, Eugenol, Z- $\beta$ -ocimene,  | To treat conjunctivitis, Fever, Malaria, headache, Indigestion, dysentery, toothache, and migraine. Antioxidant and enzyme inhibitory effects. Antifungal and antibacterial. Insect and pest control.  |
|        |  | Malesia and Indonesia   | t-hydrate-sabinene, Bicyclogermacrene, $\beta$ -caryophyllene, Carvacrol, p-menthadiene,  | Preservative for corpses. Flavoring and fragrance to foods. To treat hypothermia, skin allergy, cancer, and ulcer.   |

[ Downloaded from pbp.medilam.ac.ir on 2025-04-02 ]

|    |                                 |  | 3-O-Caffeoylquinicacid, Vitexin   |   |
|----|---------------------------------|--|---|---|
| 3. | <i>Ocimum basilicum</i>         | Bangladesh, Colombia, Bulgaria, Mali, Guinea, Nigeria, Eastern Morocco.                            | Eugenol, Methyl Chavicol, Methyl eugenol, Geraniol, Linalool, Geraniol, Cadinene, methyl cinnamate  | Food and oral care products. Antilarval, Antiviral, Antimicrobial, insecticidal, antiparasitic, antioxidant, immunomodulatory, anti-inflammatory, hepatoprotective, anti-osteoporotic activities.<br>Decrease plasma lipid content.<br>For treating nausea, flatulence, and dysentery.<br>First aid treatment for wasp stings and snakebites.<br>Inhibitory activity against HIV-1 reverse transcriptase. |
| 4. | <i>Ocimum kilimandscharicum</i> | Kenya, Rwanda, Athens, Nigeria, Sudan, Ghana, and India.   | d-camphor, Camphene, Myrcene, d- $\alpha$ -pinene, $\beta$ -pinene, Epicadinol, d-limonene, Linalool, terpinolene, Bornylacetate, Eugenol, Germacrene B, $\beta$ -caryophyllene, $\gamma$ -muurolene and unidentified sesquiterpenes and sesquiterpenes of alcohols | Antioxidant, antimalarial, antimicrobial, antiseptic, allergenic, pesticidal, antitumor, spasmogenic and antiviral activity.<br>Useful in cold and cough, abdominal pains, measles, diarrhea, bronchitis, foul ulcers and wounds, ophthalmopathy and vitiated conditions of 'vata'.<br>CNS activities like: neurotoxic, antineuralgic, CNS stimulant, tranquilizer, Anti-alzheimerian, sedative.          |
| 5. | <i>Ocimum gratissimum</i>       | India, Nigeria, New Zealand, Australia, South America, The Caribbean, South East Asia, and Africa. | Eugenol, thymol, citral, geraniol and linalool, eugenol   | <i>Antifungal, antiviral</i> and antioxidant.<br>Antinociceptive activity.<br>For treatment of rheumatism, paralysis, epilepsy, sunstroke, influenza, Gonorrhoea, high fever, diarrhea, and mental illness  |
| 6. | <i>Ocimum micranthum</i>        | Central and South America, West Indies.  | 1,8-cineol, eugenol element isomers, linalool, $\beta$ -caryophyllene   | Treatment of colds, fever, stomach disturbances, and dysentery. Antimicrobial.<br>To treat epilepsy, nervous trouble, influenza, ear aches.   |
| 7. | <i>Ocimum canum</i>             | India, Rwanda, Israel, Zimbabwe, Somali, Nigeria, USA, Central Africa,                             | Eugenol, linalool, Terpineol, Linalylacetate, camphor, methyl chavicol, Geraniol, Nerol, Geraniol,  | Treat conjunctivitis, malaria and headache<br>Insecticidal and antibacterial and antifungal activity.   |

|  |  |  |  |
|--|--|--|--|
|  |  | Terpinen-4-ol, Methyl cinnamate, 1.8- cineole. |  |
|--|--|--|--|

**Phytoconstituents:**

Comprising a modest 0.7% of the leaf weight, the volatile oil fraction of Tulsi packs a potent punch. Eugenol, with its impressive 71% dominance, reigns supreme as an antiseptic champion. Its close ally, methyl eugenol, contributes another 20%, while carvacrol and caryophyllene bring their unique anti-inflammatory and antimicrobial properties to the table. These volatile constituents likely contribute to Tulsi's traditional use in respiratory and wound healing applications.<sup>4</sup>

Fresh Tulsi leaves and stems go beyond aromatic allure, harboring a wealth of phenolic antioxidants. Compounds like circumaritin, cirsiolol, isothymusin, apigenin, and rosmarinic acid act as cellular guardians, shielding against oxidative damage. Additionally, orientin and vicenin, found in the aqueous leaf extract, join the defensive party with their antioxidant and anti-inflammatory properties. Ursolic acid, another antioxidant and anti-inflammatory warrior, stands alongside apigenin and luteolin, flavonoids with potential anti-cancer and anti-inflammatory effects. Their glycosylated forms,

apigenin-7-O-glucuronide and luteolin-7-O-glucuronide, further enrich the chemical tapestry. Finally, orientin and molludistin, with their antioxidant and antimicrobial properties, complete this diverse array of bioactive molecules.<sup>5</sup>

The symphony of Tulsi's chemistry wouldn't be complete without the enchanting dance of sesquiterpenes and monoterpenes. These aromatic compounds, including bornyl acetate, beta-elemene, neral, alpha- and beta-pinenes, and campesterol, each contribute their unique therapeutic waltz to the overall healing potential of the leaves.<sup>6</sup>

While further research is needed to fully elucidate the individual and synergistic effects of these diverse compounds, the growing body of scientific evidence highlights Tulsi's potential as a source of bioactive molecules with promising health benefits. This knowledge provides a deeper understanding of the cultural significance bestowed upon this revered plant and paves the way for future exploration of its therapeutic potential. (As mentioned in table 2)

**Table 2:** Chemical Constituents of *Ocimum tenuiflorum* and Their Activities.

| Bio-Chemical Constituents | Source      | Characteristics                   | Uses/Activity                                |
|---------------------------|-------------|-----------------------------------|--|
| Eugenol                   | Whole plant | Phenylpropane                     | Antimicrobial, Anti larvacidal, Antidiabetic |
| Camphor                   | Leaves      | Terpenoid                         | Antimicrobial, Antiviral                     |
| Ursolic acid              | Leaves      | Pentacyclic triterpenoid          | Anti-fertility, Antiarthritis, Anti-covid-19 |
| Linalool                  | Whole plant | Terpene alcohol                   | Antilarvacidal, Antimalarial, Antimicrobial  |
| Methylchavicol            | Leaves      | Phenylpropane                     | Free radical scavenging                      |
| Methyl cinnamate          | Whole plant | Methyl ester of cinnamate         | Food additives                               |
| Methyleugenol             | Leaves      | Phenylpropane                     | Antimicrobial                                |
| Caryophyllene             | Leaves      | Sesquiterpene (Natural, bicyclic) | Antimicrobial, Antilarval, Anticataract      |
| Isothymusin               | Aerial part | Flavonoid                         | Anticancer                                   |

|                  |                 |                          |   |
|------------------|-----------------|--------------------------|---|
| Orientin         | Leaves          | Glycosidic flavonoid     | Anticancer, radioprotective, lipid peroxidation                 |
| Vicenin          | Leaves          | Glycosidic flavonoid     | Anticancer, radioprotective, Anti-covid-19                      |
| Rosemarinic acid | Leaves          | Phenolic                 | Anticancer, Hepatoprotective, Antioxidant                       |
| Estragole,       | Whole plant     | Phenylpropane            | Antimicrobial   |
| Pheophytin-a     | Leaves          | Chlorophyll derivative   | Anti-HIV activity   |
| Luteolin         | Leaves, flowers | Flavonoids               | Antiarthritic   |
| Cirisileneol     | Leaves          | Phenolic or biphenyl     | Antioxidant   |
| Linolenic acid   | Seeds           | Omega-3 fatty acid       | Antimicrobial, Antiviral, Antihypertensive                      |
| Oleanolic Acid   | Leaves, Roots   | Pentacyclic triterpenoid | Hepatoprotective, Antioxidant, Antistress                       |
| Quercetin        | Leaves          | Flavonoid                | Antioxidant, Cardioprotective                                   |
| Apigenin         | Flowers, Leaves | Flavone glycosides       | Anti-inflammatory, Antioxidant, Antiarthritic                   |
| Isorientin       | Leaves          | Flavone                  | Anti covid-19 activity  |
| Catechin         | Whole plant     | Phenol                   | Cardioprotective  |
| Isoflavones      | Seeds           | Polyphenolic             | Antioxidant   |
| Sulforaphane     | Leaves          | Isothiocyanate           | Cardioprotective, Antioxidant                                   |
| Disogenin        | Roots           | Phytosteroid saponin     | Antioxidant, hypoglycemia, Anti-inflammatory, Antiproliferative |

### Traditional Uses

Revered as the "elixir of life," Tulsi whispers the promise of longevity through its potent mix of healing properties. Ayurveda and Siddha medicine, ancient systems of wisdom, embrace every part of this sacred herb, weaving its magic into remedies for an astonishing array of ailments. From the common cold's tickle to the searing heat of fever, Tulsi soothes and restores. Its fragrant leaves quell the coughs that wrack your chest, ease the ache of a throbbing head, and silence the rasping whispers of a sore throat. Whether battling bronchitis or the wheezing of asthma, this herbal warrior stands guard against respiratory foes. Tulsi's touch reaches every corner of the body, mending and strengthening. It banishes the chills of malaria fever, neutralizes the venom of serpent and scorpion, eases the churning of colic, and calms the throbbing symphony of a migraine. From skin's gentle whispers of distress to the open cries of wounds, Tulsi's balm offers solace. Mind and spirit, too, find refuge in Tulsi's embrace. Its leaves, chewed thoughtfully, sharpen memory and

soothe tired nerves, chasing away insomnia and fatigue. Even the darkness of night blindness recedes before its light, and diarrhoea's unwelcome grip loosens under its gentle influence. Truly, Tulsi is a guardian angel in every leaf, a testament to the healing power of nature woven into a living tapestry of well-being.<sup>7</sup>

### Morphological types

Shyama Tulsi (Black Holy Basil): Clothed in a cloak of deep purple, this variety, also known as Krishna Tulsi, is a powerful medicinal wonder. Its peppery bite whispers secrets of strength and resilience, making it a popular choice for fighting infections and ailments.<sup>8</sup>

Rama Tulsi (Green Holy Basil): Bathed in emerald hues, this gentle soul, also called Sri Tulsi, reigns supreme in the realm of worship. Its sweet, clove-like fragrance ascends towards the heavens, carrying prayers and offerings with each delicate leaf. But that's not all – Rama Tulsi boasts potent medicinal

properties too, making it a versatile hero in the natural medicine cabinet.<sup>9</sup>

Now, meet the adventurous one – Vana Tulsi (Forest Holy Basil)! This wild spirit, botanically known as *O. gratissimum*, prefers the freedom of the woods. Though not commonly used in worship, its vibrant green leaves and peppery kick pack a powerful punch of medicinal benefits<sup>2-7</sup>.

### Pharmacological uses

The holy basil plant (*Ocimum tenuiflorum*), revered in the ancient Ayurvedic text "Charaka Samhita," possesses a well-documented history of use in treating various ailments. Extracts from its leaves, prepared through both aqueous and alcoholic methods, demonstrate a broad spectrum of pharmacological activities. These include anti-inflammatory, antipyretic, analgesic, bronchodilatory, antiemetic, antidiabetic, hepatoprotective, hypotensive, hypocholesterolemic, and adaptogenic properties. Additionally, Tulsi leaf essential oil exhibits antibacterial, antioxidant, and anti-inflammatory effects, finding application in the development of topical pharmaceutical products, particularly for skin conditions<sup>10</sup>.

### Antimicrobial activity

*Ocimum tenuiflorum* (commonly known as Tulsi) exhibits a broad spectrum of antimicrobial activity against various bacterial pathogens, including *Candida albicans*, *Staphylococcus aureus*, and *Escherichia coli*. This activity is attributed to the presence of diverse phytoconstituents present throughout the plant's leaves, stems, and seeds.

Scientific investigations have revealed the underlying mechanisms of Tulsi's antimicrobial properties. Studies such as Singh et al. suggest that high concentrations of linoleic acid in the plant's fixed oil may contribute significantly to this effect, particularly against *S. aureus*, *Bacillus pumilus*, and multi-drug resistant *Pseudomonas aeruginosa*. Notably, *S. aureus* appears to be particularly susceptible to Tulsi extracts, highlighting its potential application against this prevalent and potentially serious pathogen.

Further research by Geeta et al. adds another layer of complexity. Their findings demonstrate that the aqueous extract of Tulsi, even at relatively low concentrations, exhibits potent antimicrobial activity against *Klebsiella*, with wider inhibition zones compared to the alcoholic extract. This observation

underscores the importance of investigating diverse extraction methods to potentially unlock the full spectrum of Tulsi's antimicrobial potential.

In conclusion, the antimicrobial activity of Tulsi is not merely anecdotal but supported by a growing body of scientific evidence. These findings suggest the potential of Tulsi as a natural source of antimicrobials in the face of increasing antibiotic resistance. Further research is crucial to fully elucidate the mechanisms of action and explore the potential development of novel and sustainable solutions for combating bacterial infections using this plant.<sup>11-18</sup>

### Action of the chemical components

*Ocimum tenuiflorum* (Tulsi) exhibits potential benefits against various dental ailments attributable to a diverse array of bioactive compounds. These include:

1. **Ursolic acid, eugenol, and carvacrol:** These phytochemicals demonstrate antimicrobial activity against *Streptococcus mutans*, the primary bacterial species implicated in dental caries. In vitro studies suggest that Tulsi extracts at a 4% concentration exhibit the most potent antibacterial activity against this pathogen, highlighting its potential as a natural anti-caries agent.
2. **Eugenol, palmitic acid, gallic acid, vallinin, vitamins A and C:** This combination of phytochemicals exhibits anti-plaque, anti-tartar, antimicrobial, and anti-inflammatory properties, potentially contributing to improved oral hygiene and periodontitis management. Their combined effects may offer benefits for oral health maintenance.
3. **Ursolic acid:** In addition to its potential benefits for oral health, this triterpene acid demonstrates promising anti-cancer activity by modulating various cellular processes associated with tumorigenesis.

Furthermore, chewing Tulsi leaves has been shown to induce a rapid increase in salivary pH, observed both immediately and after 30 minutes. While current research suggests that this effect may be attributed to salivary stimulation, further investigations are warranted to elucidate the underlying mechanisms and potential clinical implications.<sup>19-28</sup>

### Immunomodulatory Activity:

Studies utilizing steam-distilled extracts of fresh Tulsi leaves in *Rattus norvegicus* (albino rats) suggest modulations in their immune response. Potential mechanisms include enhanced antibody production, release of immune mediators, and targeted action on lymphoid organs. These effects might contribute to a strengthened immune system, with Tulsi potentially potentiating both cellular and humoral immunity through increased cell-mediated responses and activation of GABAergic pathways.<sup>29</sup>

### Anti-inflammatory Activity

Tulsi's anti-inflammatory and analgesic properties are supported by preclinical research demonstrating its potential against both acute and chronic inflammation in rats. Various mechanisms may contribute to this effect:

**Methanol extract and aqueous suspension:** At a dose of 500 mg/kg, these extracts exhibit analgesic, antipyretic, and anti-inflammatory effects in rats.<sup>30</sup>

**Fixed oil and linolenic acid:** These components target the arachidonic acid pathway, a key player in inflammation, by inhibiting both lipoxygenase and cyclooxygenase enzymes. This leads to a significant reduction in the production of inflammatory mediators like prostaglandin E<sub>2</sub>, leukotrienes, and arachidonic acid.<sup>31</sup>

**Essential and fixed oils:** Studies show their effectiveness against carrageenan-induced paw edema, croton oil-induced granuloma, and exudate formation in rats. While showing promising anti-edematous activity compared to the standard drug flurbiprofen, Tulsi oils did not reach the same level of efficacy.<sup>32</sup>

**Fixed oil:** Research suggests it can prevent increased vascular permeability and leukocyte infiltration, key features of inflammation, in carrageenan-induced models.<sup>33</sup>

Extensive scientific literature, including scholarly publications and databases, delves deeper into the anti-inflammatory properties of Tulsi. This research explores mechanisms beyond the arachidonic acid pathway, such as modulation of inflammatory cytokines and antioxidant activity.

### Anticancer Activity:

Tulsi's potential as an antineoplastic agent is supported by preclinical studies demonstrating its ability to target various aspects of tumorigenesis:

**Detoxification Enhancement:** The alcoholic extract of Tulsi leaves modulates the activity of key detoxification enzymes such as cytochrome P450 and glutathione S-transferase. These enzymes play a crucial role in neutralizing carcinogens and mutagens, potentially preventing their ability to induce cellular damage and promote cancer development.

**Direct Cytotoxic Activity:** In vitro studies using Tulsi extracts (AIE) at concentrations exceeding 50 µg/mL exhibit cytotoxic effects on cancer cells. This translates to apoptosis, characterized by cytoplasmic shrinkage and nuclear condensation. Further analysis reveals DNA fragmentation, suggesting Tulsi's capacity to disrupt the integrity of cancer cells.

**Tumor Reduction in Animal Models:** The in vivo efficacy of Tulsi has been demonstrated in models where mice were exposed to lung and liver carcinogens. Treatment with Tulsi extracts resulted in a significant reduction in tumor incidence.

**Skin Cancer Inhibition:** Research explores the potential of Tulsi in preventing skin cancer. Application of Tulsi leaf extract significantly reduced tumor burden in mice with chemically induced papillomas. This protective effect is attributed to enhanced antioxidant activity and upregulation of detoxification enzymes. Notably, eugenol, a naturally occurring compound in Tulsi, plays a key role in this process.

**Early Intervention Potential:** Studies suggest that Tulsi may be beneficial even in the early stages of tumor development. Application of fresh Tulsi leaf paste to the oral cavity of mice effectively prevented the progression of pre-cancerous lesions induced by a carcinogen.

**Proposed Mechanism of Action:** The antineoplastic properties of Tulsi are likely linked to its ability to inhibit or suppress the metabolic activation of carcinogens. This effectively hinders their potential to initiate and promote cancer development.

**Support from Established Tumor Models:** Studies utilizing established tumor models in mice, such as Ehrlich ascites carcinoma and S180, further support the broad-spectrum anticancer potential of Tulsi extracts. These studies demonstrate

positive results, highlighting the need for further investigation into the mechanisms and efficacy of Tulsi in humans compared to conventional therapies.

While further research is crucial, Tulsi's potential as a natural approach in the fight against cancer warrants deeper exploration.<sup>34,35,36,37,38,39,40</sup>

### Anticoagulant Activity

Preliminary research suggests that Tulsi may offer a natural alternative to conventional antithrombotic agents in managing blood clotting. Studies explore the potential anticoagulant activity of Tulsi's fixed oil, demonstrating its ability to prolong blood clotting time.

**Mechanism of Action:** The proposed mechanism involves the linolenic acid present in Tulsi oil. This fatty acid can be metabolized into eicosapentaenoic acid (EPA), which then enters the cyclooxygenase pathway. This pathway leads to the production of two prominent metabolites:

**Prostaglandin I3 (PGI3):** This molecule exhibits antiplatelet aggregation properties, acting as an endogenous anticoagulant.

**Thromboxane A3 (TXA3):** Unlike its counterpart TXA2, which promotes platelet aggregation, TXA3 displays minimal platelet aggregation activity, further contributing to the anticoagulant effect.

The combined effect of elevated PGI3 and TXA3 levels while simultaneously suppressing TXA2 production is believed to contribute to the anticoagulant properties of Tulsi's fixed oil. In vitro studies support this hypothesis, demonstrating that a 3 ml/kg dose of the oil exhibits similar blood clotting time prolongation as 100 mg/kg of aspirin.<sup>41,42,43,44,45,46,47,48,49,50</sup>

### Antifungal Activity

Beyond its aesthetic appeal, Tulsi exhibits broad-spectrum antifungal activity against various filamentous fungi, including established pathogens like *Aspergillus* and *Penicillium* species, as well as emerging threats such as *Fusarium* and *Rhizomucor*. This activity is attributed to a diverse array of bioactive compounds present within Tulsi extracts:

**Methyl chavicol and linalool:** These volatile compounds demonstrate direct antifungal activity, contributing significantly to Tulsi's antifungal potential.

**Secondary metabolites:** This group, encompassing alkaloids, glycosides, saponins, tannins, and ascorbic acid, plays a synergistic role in the antifungal effect. Notably, eugenol, previously mentioned, is a particularly potent member of this class.

### Recent Research

A 2023 study published in "Mycological Progress" investigated the efficacy of Tulsi extracts against five prevalent dermatophytic fungi, known to cause skin infections. The study confirmed their antifungal activity, with a minimum inhibitory concentration (MIC) of 200 µg/mL, signifying their ability to inhibit fungal growth at this concentration.

Ongoing research at the National Institute of Plant Health Management in India delves into the potential of Tulsi extracts for managing plant pathogens like *Fusarium oxysporum*, a significant threat to agricultural crops.

**Future Directions:**

In the face of rising concerns regarding antifungal resistance, researchers are actively exploring natural alternatives like Tulsi.

Investigating the synergistic effects of combining Tulsi extracts with conventional antifungal drugs holds promise for developing more effective and sustainable treatment options.

Elucidating the specific mechanisms of action of individual antifungal compounds within Tulsi is crucial to fully harnessing their potential.<sup>50,51,52,53,54,55,56,57,58</sup>

### Radioprotective Activity

Since its discovery in 1995, Tulsi has attracted significant interest for its potential radioprotective properties. Studies demonstrate its ability to mitigate the harmful effects of radiation on human lymphocytes, even at low doses. Notably, orientin and vicenin, two flavonoids present in Tulsi leaves, exhibit superior radioprotective activity compared to synthetic protectors.<sup>58</sup>

Beyond its individual efficacy, Tulsi extracts demonstrate synergistic effects when combined with established



radioprotectors like WR-2721. This combination not only enhances bone marrow cell protection but also reduces the toxicity associated with higher doses of WR-2721, highlighting its potential for improved therapeutic strategies.<sup>59</sup>

**Comparative Studies:** Research suggests that the aqueous extract of Tulsi offers superior radioprotective activity compared to the alcoholic extract in animal models. The optimal dose for this effect has been established at 10 mg/kg.<sup>60</sup>

**Ongoing Investigations:**

Eugenol, a natural oil component, and ursolic acid, a bioactive compound present in Tulsi, are currently under investigation for their potential radioprotective properties.<sup>61</sup>

Research is actively exploring the potential of Tulsi in mitigating radiation-induced side effects like nausea and fatigue, offering a more holistic approach to radiation protection beyond cellular damage management.<sup>62</sup>

**Future Considerations:**

While preliminary findings are promising, further research is crucial to translate these observations into safe and effective clinical applications for humans. Nonetheless, Tulsi's potential as a natural radioprotective agent presents an exciting avenue for future exploration in healthcare.

## Antioxidant Activity

Research suggests that Tulsi exhibits potent antioxidant activity due to its rich array of bioactive compounds. These compounds help combat the detrimental effects of free radicals on cellular structures.

**Flavonoids:** Notably, orientin and vicenin, two prominent flavonoids present in Tulsi, have demonstrated significant antioxidant activity in vivo studies. They effectively reduce radiation-induced lipid peroxidation in mouse livers, highlighting their ability to protect cell membranes from oxidative damage.

**Free Radical Scavenging:** Tulsi extracts exhibit free radical scavenging activity, directly neutralizing highly reactive free radicals and preventing them from inflicting cellular damage.

**Diverse Antioxidant Arsenal:** Tulsi possesses a diverse range of phenolic compounds, including cirsilineol, cirsimaritin, isothymusin, apigenin, and rosmarinic acid, along with

significant amounts of eugenol from its essential oil. These compounds collectively contribute to the plant's antioxidant potential, each offering unique mechanisms of action against free radicals.<sup>63,64,65,66,67,68,69</sup>

## Antihypertensive and Cardioprotective Activities

Tulsi's potential benefits extend beyond its aesthetic appeal, demonstrating promising neuroprotective and cardioprotective properties in preclinical research.

**Neuroprotective Effects:**

**Transient cerebral ischemia and chronic cerebral hypoperfusion:** Recent studies investigate Tulsi's potential to mitigate the detrimental effects of transient cerebral ischemia (TCI) and chronic cerebral hypoperfusion (CCH). Both conditions can lead to neurological deficits characterized by cellular swelling, scarring, and inflammation. However, research suggests that Tulsi may offer neuroprotective benefits by mitigating these detrimental effects.

**Cardioprotective Effects:**

**Hypotensive activity:** The intravenous administration of Tulsi's fixed oil exhibits a significant hypotensive effect, lowering blood pressure. This effect is likely attributed to its ability to induce peripheral vasodilation, facilitating smoother blood flow and potentially reducing strain on the heart.

**Essential fatty acids and vasodilation:** The essential fatty acids present in Tulsi oil, linoleic acid and linolenic acid, play a crucial role in this vasodilation process. These fatty acids can be metabolized into vasodilatory prostaglandins (PGE1 and PGE3) of the series 1 and 3 families, while simultaneously suppressing the formation of vasoconstricting prostaglandin (PGE2) of the series 2 family.<sup>70,71,72,73,74</sup>

**Emerging Research Areas:**

**Cognitive function and memory:** Ongoing research explores the potential of Tulsi extracts to improve cognitive function and memory, offering potential therapeutic options for neurodegenerative diseases like Alzheimer's.

Stroke-induced brain damage: Studies are actively investigating Tulsi's ability to protect against brain damage caused by stroke, further supporting its potential as a neuroprotective agent.

### Central Nervous System Depressant Activity

Tulsi exhibits potential psychotropic effects, interacting with the central nervous system (CNS) in various ways, as evidenced by preclinical research.

### Calming and Anxiolytic Activity

Alcoholic extract (AIE): Studies demonstrate that AIE can delay the loss of reflexes in mice under the influence of the sedative pentobarbital. Additionally, it reduces the duration and severity of seizures induced by electroshock or pentylenetetrazole. These findings suggest that Tulsi might possess calming and anxiolytic properties by modulating an overactive nervous system.

Behavioral effects: Mice administered AIE displayed reduced aggression and activity in open field tests, indicating potential anti-anxiety and anti-hyperactivity effects.

### Mood-Boosting Potential

At higher doses, AIE was observed to prolong swimming time in mice, which may suggest stimulant or anti-stress properties. This effect was comparable to desipramine, a known antidepressant, hinting at Tulsi's potential role in mood regulation.

### Contrasting Effects of Fixed Oil

Interestingly, Tulsi's fixed oil appears to have opposed effects, extending pentobarbital-induced sleep duration in rats. This observation may be attributed to its interaction with the drug's metabolic pathways or clearance mechanisms.

### Unraveling the Complexity

These seemingly conflicting findings suggest a complex interaction between Tulsi and the CNS. Ongoing research endeavors to decipher this intricate interplay, focusing on identifying the specific compounds and mechanisms responsible for Tulsi's calming, mood-boosting, and potentially sleep-promoting effects.<sup>75,76,77,78,79,80,81,82</sup>

### Analgesic Activity

Despite limitations observed in conventional pain models, Tulsi oil exhibited potential analgesic activity in a study investigating acetic acid-induced writhing in mice. This suggests a possible

role in pain management through the inhibition of prostaglandins, histamine, and acetylcholine, which are known to participate in pain pathways.

Emerging Applications: Current research is exploring the potential of topical application of Tulsi extracts for managing joint and muscle pain.<sup>83,84,85,86,87</sup>

### Antipyretic Activity

Tulsi oil's fixed oil demonstrates antipyretic activity, effectively reducing fever in rats injected with a typhoid vaccine. This effect is comparable to that of aspirin. The proposed mechanism involves the suppression of prostaglandin synthesis, as these molecules play a crucial role in promoting fever.

Emerging Applications: Current research is exploring the potential of Tulsi extracts as natural antipyretics for children, offering a potentially safer alternative to conventional medications.<sup>88,89,90</sup>

### Cognitive Enhancement

Preclinical studies using Tulsi's alcoholic extract (AIE) suggest potential cognitive benefits, particularly in memory improvement. AIE administration has been shown to enhance memory in both scopolamine-treated mice and aging mice, hinting at possible therapeutic applications in neurodegenerative disorders like Alzheimer's and dementia. Current research is actively investigating the underlying mechanisms responsible for Tulsi's memory-boosting effects, exploring its interactions within the brain.<sup>91,92,93</sup>

### Hepatoprotective Activity

Tulsi exhibits hepatoprotective properties, demonstrating the ability to protect the liver from damage. Studies have shown that extracts from Tulsi leaves can shield rats from paracetamol-induced liver injury, highlighting its potential for safeguarding this vital organ. Ongoing research delves deeper into Tulsi's potential to combat other forms of liver damage, including those caused by viral infections and fatty liver disease.<sup>94,95,96,97</sup>

### Fertility Regulation

While studies have observed reduced sperm count and motility in male rats administered specific doses of Tulsi extracts, further research is crucial to comprehensively understand its impact on human fertility. It is important to emphasize that this

information should be approached with caution and not interpreted as a definitive guide for fertility control.<sup>98,99,100,101</sup>

### Antidiabetic Potential

Tulsi demonstrates promising antidiabetic properties, exhibiting the ability to lower blood sugar levels in both animal and human studies. This suggests potential applications in the management of diabetes. Further research is underway to elucidate how Tulsi interacts with the body's insulin system and explore its potential long-term benefits for diabetic patients.<sup>102,103,104</sup>

### Antiulcer Activity

Tulsi's fixed oil displays antiulcer properties, acting as a natural shield against various ulcer-inducing agents such as aspirin and alcohol. Its lipooxygenase-inhibiting, histamine-antagonistic, and antisecretory properties are believed to contribute to its effectiveness in combating ulcers. Current research explores the potential of combining Tulsi extracts with conventional ulcer medications to achieve potentially enhanced efficacy and reduced side effects.<sup>105,106</sup>

### Antiarthritic Potential

Preclinical studies suggest that Tulsi's fixed oil possesses significant antiarthritic activity in rats with formaldehyde-induced arthritis. Its ability to inhibit inflammatory mediators like serotonin and prostaglandins suggests potential for alleviating arthritic pain and inflammation. Future research holds promise in investigating how Tulsi extracts can be safely and effectively integrated into arthritis treatment plans.<sup>107,108,109,110</sup>

### Immunomodulatory and Adaptogenic Activity:

Tulsi's potential extends beyond its adaptogenic activity, encompassing immunomodulatory properties as well. Research suggests that it may play a role in enhancing the immune system's response to various stressors, both physical and mental.<sup>111</sup>

### Potential Mechanisms:

**Immunostimulant activity:** Tulsi's immunostimulatory effects are believed to contribute to its adaptogenic properties. This implies its potential to bolster the body's innate defense mechanisms against diverse stressors.

**Enhanced physical endurance:** Studies in mice have demonstrated that Tulsi's alcoholic extract (AIE) can prolong their swimming time, suggesting potential benefits in improving energy levels and resilience.

**Stress modulation:** AIE has been shown to exhibit anti-stress properties in rats, preventing the formation of stress-induced ulcers. This suggests that Tulsi may help mitigate the body's response to stressful situations, potentially safeguarding physical health.

**Regulation of immune response:** Administration of the same extract in mice countered the increase in white blood cells triggered by milk (a foreign substance). This observation points towards Tulsi's potential to regulate the immune system's response to potential threats, preventing excessive reactions.<sup>112,113,114</sup>

### Holistic Approach to Well-being

These findings collectively portray Tulsi as a potential holistic immune system booster. It may offer various benefits, including enhanced physical stamina, mitigation of stress-induced damage, and regulation of immune responses, contributing to overall well-being.

### Toxicity of Tulsi Fixed Oil

Research suggests that Tulsi fixed oil, a concentrated extract obtained from the leaves, exhibits relatively good tolerability at moderate doses. Here's a summary of the key findings:

**Median Lethal Dose (LD50):** Studies in mice have established the LD50 of the oil to be approximately 42.5 ml/kg when administered via intraperitoneal injection. This value indicates relative safety at lower doses.

**High-Dose Tolerance:** Intraperitoneal administration of doses as high as 30 ml/kg in mice did not induce any observable negative effects, suggesting the oil's potential for safe usage within specific dose ranges.

**Subacute Toxicity Assessment:** A 14-day subacute toxicity study in rats receiving daily doses of 3 ml/kg of the oil revealed no adverse effects, further supporting its safety profile at moderate doses.<sup>115,116</sup>

## Ongoing Research

**Long-Term Safety:** Current research is actively investigating the long-term safety of Tulsi oil consumption, particularly focusing on its potential impact on various organs and metabolic functions.

**Drug Interactions:** Studies are also underway to evaluate the potential interactions between Tulsi oil and other medications, ensuring its safe integration into existing treatment plans.

## Conclusion

Tulsi (*Ocimum tenuiflorum*) presents itself as a multifaceted herb with a diverse array of potential health benefits. Scientific evidence increasingly supports its traditional uses, demonstrating its antimicrobial, anti-inflammatory, and anticancer properties. Additional research continues to explore its potential benefits in managing conditions ranging from diabetes and ulcers to cognitive decline and arthritis.

Recent advancements highlight the broad spectrum of Tulsi's potential. Studies investigating its interaction with the central nervous system suggest possibilities for managing anxiety, mood regulation, and pain perception. Additionally, its potential as a natural radioprotective agent and its ability to modulate the immune system warrant further exploration.

While the long-term safety of Tulsi appears promising at moderate doses, ongoing research is crucial to fully elucidate its mechanisms of action, potential drug interactions, and long-term effects. Considering the growing body of evidence, Tulsi emerges as a promising natural resource deserving further investigation in the pursuit of holistic well-being.

## Declarations

### Acknowledgement

We are grateful to the Gayatri College of Pharmacy and the Tagore Institute of Pharmacy and Research for their invaluable support in providing access to resources and guidance.

### Ethical Approval

Not Applicable. This manuscript does not report on a clinical trial and did not involve any human or animal subjects.

### Funding

Not Applicable. No funding was received for this.

## Availability of data and materials

Not Applicable.

## References

1. Khurana, P. A Review On Medicinal Uses of *Ocimum Tenuiflorum* Linn (TULSI).
2. Singh, N., Hoette, Y., and Miller, D. R. (2002). Tulsi: The mother medicine of nature. International Institute of Herbal Medicine.
3. Pandey, G., and Madhuri, S. (2010). Pharmacological activities of *Ocimum sanctum* (tulsi): a review. *Int J Pharm Sci Rev Res*, 5(1), 61-66.
4. Cohen M. M. (2014). Tulsi - *Ocimum sanctum*: A herb for all reasons. *Journal of Ayurveda and integrative medicine*, 5(4), 251-259.
5. Kumar, K. P., Bhowmik, D., Tripathi, K. K., and Chandira, M. (2010). Traditional Indian Herbal Plants Tulsi and Its Medicinal Importance. *Research Journal of Pharmacognosy and Phytochemistry*, 2(2), 93-101.
6. Pattanayak P, Behera P, Das D, Panda SK. *Ocimum sanctum* Linn. A reservoir plant for therapeutic applications: An overview. *Pharmacogn Rev*. 2010 Jan;4(7):95-105.
7. Singh, E., Sharma, S., Dwivedi, J., and Sharma, S. (2012). Diversified potentials of *Ocimum sanctum* Linn (tulsi): an exhaustive survey. *Journal of Natural Product and Plant Resources*, 2(1), 39-48.
8. Bhooshitha, A. N., Ghosh, A. R., Chandan, H. M., Nandhini, H. S., Pramod, B. R., and Krishna, K. L. (2020). Review On Nutritional, Medicinal and CNS Activities of Tulsi (*Ocimum. Sanctum*). *Journal of Pharmaceutical Sciences and Research*, 12(3), 420-426.
9. Evans, W. C. (1996). Phenols and phenolic glycosides. *Trease and Evans Pharmacognosy*, 14th ed. Noida: Gopsons Papers Limited, 218.
10. Pooja, Anil Kumar. A Systemic Review of Tulsi (*Ocimum tenuiflorum* or *Ocimum sanctum*): Phytoconstituents, Ethnobotanical and Pharmacological Profile. *Research Journal of Pharmacognosy and Phytochemistry*. 2023; 15(2):179-8.
11. Sajjadi, S. E. (2006). Analysis of the essential oils of two cultivated basil (*Ocimum basilicum* L.) from Iran. *DARU Journal of Pharmaceutical Sciences*, 14(3), 128-130.
12. Kumar, P. K., Kumar, M. R., Kavitha, K., Singh, J., and Khan, R. (2012). Pharmacological actions of *Ocimum sanctum*—review article. *International Journal of Advances in Pharmacy, Biology and Chemistry*, 1(3), 2277-4688.
13. Wangcharoen, W., and Morasuk, W. (2007). Antioxidant capacity and phenolic content of holy basil. *Songklanakarinn J Sci Technol*, 29(5), 1407-1415.

14. Atal, C. K., and Kapur, B. M. (1982). Cultivation and utilization of medicinal plants. Jammu- Tawi: Regional Research Laboratory, Council of Scientific and Industrial Research.
15. Gupta, P. A Review on Medicinal Uses of *Ocimum Tenuiflorum* LINN (TULSI).
16. Prajapati, N. D., Purohit, S. S., Sharma, A. K., and Kumar, T. (2003). A handbook of medicinal plants: A complete source book. In A handbook of medicinal plants: a complete source book (pp. 554-554).
17. Gupta, S. K., Prakash, J., and Srivastava, S. (2002). Validation of traditional claim of Tulsi, *Ocimum sanctum* Linn.as a medicinal plant.
18. Tewari, D., Sah, A. N., Pandey, H. K., Meena, H. S., Meena, R., Ramaswamy, R. S., ... and Murthy, P. H. (2012). A review on phytoconstituents of *Ocimum* (Tulsi). *International Journal of Ayurvedic Medicine*, 3(1), 1-9.
19. Chauhan, N. S. (1999). Medicinal and aromatic plants of Himachal Pradesh. Indus publishing.
20. Shah, C. S., and Qadry, J. S. A Textbook of Pharmacognosy. 1998. Back to cited text, (32), 216.
21. Yanpallewar, S. U., Rai, S., Kumar, M., and Acharya, S. B. (2004). Evaluation of antioxidant and neuroprotective effect of *Ocimum sanctum* on transient cerebral ischemia and long- term cerebral hypoperfusion. *Pharmacology Biochemistry and Behavior*, 79(1), 155-164.
22. Nair, A. R., Gunasegaran, R., and Joshi, B. S. (1982). Chemical investigation of certain south Indian plants. *Indian Journal of Chemistry Section B-Organic Chemistry Including MEDICINAL Chemistry*, 21(10), 979-980.
23. Patani, A. (2002). *Indian Herbal Pharmacopoeia*. Mumbai: Indian Drug Manufactures Association.
24. Singh, S., Majumdar, D. K., and Rehan, H. M. S. (1996). Evaluation of anti-inflammatory potential of fixed oil of *Ocimum sanctum* (Holybasil) and its possible mechanism of action. *Journal of Ethnopharmacology*, 54(1), 19-26.
25. Rucinke, D. S., Ferreira, P. F., Leme, P. R. P., Lapa-Guimarães, J., and Viegas, E. M. M. (2021). *Ocimum americanum* and *Lippia alba* essential oils as anaesthetics for Nile tilapia: Induction, recovery of apparent unconsciousness and sensory analysis of filets. *Aquaculture*, 531, 735902.
26. Zengin, G., Ferrante, C., Gnapi, D. E., Sinan, K. I., Orlando, G., Recinella, L., ... and Menghini, L. (2019). Comprehensive approaches on the chemical constituents and pharmacological properties of flowers and leaves of American basil (*Ocimum americanum* L). *Food Research International*, 125, 108610.
27. Sen, S. K., and Behera, L. M. (2008). Ethnomedicinal plants used by the tribals of Bargarh district to cure diarrhoea and dysentery.
28. Shah, A., and Rahim, S. (2017). Ethnomedicinal uses of plants for the treatment of malaria in Soon Valley, Khushab, Pakistan. *Journal of ethnopharmacology*, 200, 84-106.
29. Selvi, M. T., Thirugnanasampandan, R., and Sundarammal, S. (2015). Antioxidant and cytotoxic activities of essential oil of *Ocimum canum* Sims. from India. *Journal of Saudi Chemical Society*, 19(1), 97-100.
30. Adomako-Bonsu, A. G., Chan, S. L., Pratten, M., and Fry, J. R. (2017). Antioxidant activity of rosmarinic acid and its principal metabolites in chemical and cellular systems: Importance of physico-chemical characteristics. *Toxicology in vitro*, 40, 248-255.
31. Oyedemi, S. O., Oyedemi, B. O., Cooposamy, R. M., Prieto, J. M., Stapleton, P., and Gibbons, S. (2017). Antibacterial and norfloxacin potentiation activities of *Ocimum americanum* L. against methicillin resistant *Staphylococcus aureus*. *South African Journal of Botany*, 109, 308-314.
32. Pandey, R., Chandra, P., Kumar, B., Dutt, B., and Sharma, K. R. (2016). A rapid and highly sensitive method for simultaneous determination of bioactive constituents in leaf extracts of six *Ocimum* species using ultra high performance liquid chromatography-hybrid linear ion trap triple quadrupole mass spectrometry. *Analytical Methods*, 8(2), 333-341.
33. Dhama, K., Sharun, K., Gugjoo, M. B., Tiwari, R., Alagawany, M., Iqbal Yatoo, M., ... and Farag, M. R. (2021). A Comprehensive Review on Chemical Profile and Pharmacological Activities of *Ocimum basilicum*. *Food Reviews International*, 1-29.
34. Surburg, H., and Panten, J. *Common Fragrance and Flavor Materials*.
35. Chiang, L. C., Ng, L. T., Cheng, P. W., Chiang, W., and Lin, C. C. (2005). Antiviral activities of extracts and selected pure constituents of *Ocimum basilicum*. *Clinical and Experimental Pharmacology and Physiology*, 32(10), 811-816.
36. Hussain, A. I., Anwar, F., Sherazi, S. T. H., and Przybylski, R. (2008). Chemical composition, antioxidant and antimicrobial activities of basil (*Ocimum basilicum*) essential oils depends on seasonal variations. *Food chemistry*, 108(3), 986-995.
37. Baytop, T. (1984). *Therapy with Medicinal Plants in Turkey (Past and Present)* No: 3255. Istanbul: Publications of the Istanbul University, 359.
38. Mondello, L., Zappia, G., Cotroneo, A., Bonaccorsi, I., Chowdhury, J. U., Yusuf, M., and Dugo, G. (2002). Studies on the essential oil-bearing plants of Bangladesh. Part VIII. Composition of some *Ocimum* oils *O. basilicum* L. var. *purpurascens*; *O. sanctum* L. green; *O. sanctum* L. purple; *O. americanum* L., citral type; *O. americanum* L., camphor type. *Flavour and fragrance journal*, 17(5), 335-340.

39. Viña, A., and Murillo, E. (2003). Essential oil composition from twelve varieties of basil (*Ocimum* spp) grown in Colombia. *Journal of the Brazilian chemical society*, 14(5), 744- 749.
40. Kéita, S. M., Vincent, C., Schmit, J. P., and Bélanger, A. (2000). Essential oil composition of *Ocimum basilicum* L., *O. gratissimum* L. and *O. suave* L. in the Republic of Guinea. *Flavour and fragrance journal*, 15(5), 339-341.
41. Charles, D. J., and Simon, J. E. (1992). Essential oil constituents of *Ocimum kilimandscharicum* Guerke. *Journal of essential oil Research*, 4(2), 125-128.
42. Barker, C., Dunn, H. C., and Hilditch, T. P. (1950). African drying oils. V. Some Nigerian and Sudanese drying oils. *Journal of the Society of Chemical Industry*, 69(3), 71-75.
43. Sastri, B. N. (1962). *The Wealth of India. A Dictionary of Indian Raw Materials and Industrial Products. Raw Materials, Vol. 6: LM. The Wealth of India. A Dictionary of Indian Raw Materials and Industrial Products. Raw Materials, Vol. 6: LM.*
44. Soni, N., Gill, D., Sagar, B. S., Raheja, S., and Agrawal, S. (2012). *Ocimum kilimandscharicum*: A systematic review. *Journal of Drug Delivery and Therapeutics*, 2(3).
45. Ntezurubanza, L. J. J. C., Scheffer, J. J. C., Looman, A., and Svendsen, A. B. (1984). Composition of Essential Oil of *Ocimum kilimandscharicum* Grown in Rwanda. *Planta medica*, 50(05), 385-388.
46. Prakash, P. A. G. N., and Gupta, N. (2005). Therapeutic uses of *Ocimum sanctum* Linn (Tulsi) with a note on eugenol and its pharmacological actions: a short review. *Indian journal of physiology and pharmacology*, 49(2), 125.
47. Nadkarni, K., and Nadkarni, A. K. (1976). *Indian Materia Medica*, Popular Prakashan Pvt. Ltd., Bombay, 1, 799.
48. Warriar, P. K., Nambiar, V. P. K., Ramankutty, C., and Indian Medi-cinal Plants, A. (1995). *A compendium of 500 species. Indian Medicinal Plants. Orient Longman Publisher*, 4, 157-168.
49. Kweka, E. J., Mosha, F. W., Lowassa, A., Mahande, A. M., Mahande, M. J., Massenga, C. P., ... and Temu, E. A. (2008). Longitudinal evaluation of *Ocimum* and other plants effects on the feeding behavioral response of mosquitoes (Diptera: Culicidae) in the field in Tanzania. *Parasites and Vectors*, 1(1), 1-8.
50. Singh, P., Kalunke, R. M., Shukla, A., Tzfadia, O., Thulasiram, H. V., and Giri, A. P. (2020). Biosynthesis and tissue-specific partitioning of camphor and eugenol in *Ocimum kilimandscharicum*. *Phytochemistry*, 177, 112451.
51. Awogbindin, I. O., Tade, O. G., Metibemu, S. D., Olorunsogo, O. O., and Farombi, E. O. (2014). Assessment of flavonoid content, free radical scavenging and hepatoprotective activities of *Ocimum gratissimum* and *Spondias mombin* in rats treated with dimethylnitrosamine. *Arch Basic Appl Med*, 2, 45-54.
52. Dubey, N. K., Tiwari, T. N., Mandin, D., Andriamboavonjy, H., and Chaumont, J. P. (2000). Antifungal properties of *Ocimum gratissimum* essential oil (ethyl cinnamate chemotype). *Fitoterapia*, 71(5), 567-569.
53. Akinmoladun, A. C., Ibukun, E. O., Afor, E., Obuotor, E. M., and Farombi, E. O. (2007). Phytochemical constituent and antioxidant activity of extract from the leaves of *Ocimum gratissimum*. *Scientific Research and Essays*, 2(5), 163-166.
54. Dhawan, B. N., Patnaik, G. K., Rastogi, R. P., Singh, K. K., and Tandon, J. S. (1977). Screening of Indian plants for biological activity: part VI.
55. Rabelo, M., Souza, E. P., Soares, P. M. G., Miranda, A. V., Matos, F. J. A., and Criddle, D. N. (2003). Antinociceptive properties of the essential oil of *Ocimum gratissimum* L. (Labiatae) in mice. *Brazilian Journal of Medical and Biological Research*, 36(4), 521- 524.
56. Edeoga, H. O., Omosun, G., and Uche, L. C. (2006). Chemical composition of *Hyptis suaveolens* and *Ocimum gratissimum* hybrids from Nigeria. *African Journal of Biotechnology*, 5(10).
57. Charles, D. J., Simon, J. E., and Wood, K. V. (1990). Essential oil constituents of *Ocimum micranthum* Willd. *Journal of agricultural and food chemistry*, 38(1), 120-122.
58. Sacchetti, G., Medici, A., Maietti, S., Radice, M., Muzzoli, M., Manfredini, S., ... and Bruni, R. (2004). Composition and functional properties of the essential oil of Amazonian basil, *Ocimum micranthum* Willd., Labiatae in comparison with commercial essential oils. *Journal of Agricultural and Food Chemistry*, 52(11), 3486-3491.
59. Vieira, P. R., de Morais, S. M., Bezerra, F. H., Ferreira, P. A. T., Oliveira, Í. R., and Silva, M. G. V. (2014). Chemical composition and antifungal activity of essential oils from *Ocimum* species. *Industrial Crops and Products*, 55, 267-271.
60. de Vasconcelos Silva, M. G., Matos, F. J. A., Machado, M. I. L., and de Oliveira Silva, F. (2004). Essential Oil Composition of the Leaves of *Ocimum micranthum* Willd. *Journal of Essential Oil Research*, 16(3), 189-190.
61. Caamal-Herrera, I. O., Carrillo-Cocom, L. M., Escalante-Réndiz, D. Y., Aráiz-Hernández, D., and Azamar-Barrios, J. A. (2018). Antimicrobial and antiproliferative activity of essential oil, aqueous and ethanolic extracts of *Ocimum micranthum* Willd leaves. *BMC complementary and alternative medicine*, 18(1), 1-9.
62. Sen, P. (1993). Therapeutic potentials of Tulsi: from experience to facts. *Drugs News and Views*, 1(2), 15-21.
63. Hasan, S. B., and Deo, P. S. (1994). *Ocimum sanctum* seeds for mosquito control. *International pest control*, 36(1), 20-21.

64. Mukherjee, P. K., Maiti, K., Mukherjee, K., and Houghton, P. J. (2006). Leads from Indian medicinal plants with hypoglycemic potentials. *Journal of ethnopharmacology*, 106(1), 1- 28.
65. Jena, J., Ranjan, R., Ranjan, P., and Sarangi, M. K. (2012). A study on natural anticancer plants. *Int J Pharmaceut Chem Sci*, 1(1), 365-8.
66. Nadkarni, K. M. (1996). [Indian materia medica]; Dr. KM Nadkarni's Indian materia medica: with Ayurvedic, Unani-Tibbi, Siddha, allopathic, homeopathic, naturopathic and home remedies, appendices and indexes. 1 (Vol. 1). Popular Prakashan.
67. Ngassoum, M. B., Ousmaila, H., Ngamo, L. T., Maponmetsem, P. M., Jirovetz, L., and Buchbauer, G. (2004). Aroma compounds of essential oils of two varieties of the spice plant *Ocimum canum* Sims from northern Cameroon. *Journal of Food Composition and Analysis*, 17(2), 197-204.
68. Martins, A. P., Salgueiro, L. R., Vila, R., Tomi, F., Cañigueral, S., Casanova, J., ... and Adzet, T. (1999). Composition of the essential oils of *Ocimum canum*, *O. gratissimum* and *O. minimum*. *Planta medica*, 65(02), 187-189.
69. Ravid, U., Putievsky, E., Katzir, I., and Lewinsohn, E. (1997). Enantiomeric composition of linalol in the essential oils of *Ocimum* species and in commercial basil oils. *Flavour and fragrance journal*, 12(4), 293-2
70. Euloge, S. A., Kouton, S., Dahouenon-Ahoussi, E., Sohounhloue, D. C. K., and Soumanou, M. M. (2012). Antifungal activity of *Ocimum canum* essential oil against toxinogenic fungi isolated from peanut seeds in post-harvest in Benin. *International Research Journal of Biological Sciences*, 1(7), 20-26.
71. Dewangan, A., Sahu, B. P., and Meher, B. (2020). Review on Pharmacological Potential of *Ocimum sanctum* L. *Advanced Journal of Bioactive Molecules*, 17-24.
72. Madhuri, S., and Govind, P. (2010). Effect of ProImmu, a herbal drug on estrogen caused uterine and ovarian cytotoxicity. *Biomed*, 5(1), 57-62.
73. Saija, A., Scalese, M., Lanza, M., Marzullo, D., Bonina, F., and Castelli, F. (1995). Flavonoids as antioxidant agents: importance of their interaction with biomembranes. *Free Radical Biology and Medicine*, 19(4), 481-486.
74. Chattopadhyay, R. R. (1993). Hypoglycemic effect of *Ocimum sanctum* leaf extract in normal and streptozotocin diabetic rats. *Indian journal of experimental biology*, 31(11), 891-893.
75. Hannan, J. M. A., Marenah, L., Ali, L., Rokeya, B., Flatt, P. R., and Abdel-Wahab, Y. H. A. (2006). *Ocimum sanctum* leaf extracts stimulate insulin secretion from perfused pancreas, isolated islets and clonal pancreatic  $\beta$ -cells. *Journal of Endocrinology*, 189(1), 127-136.
76. Nagarajan, S., Jain, H. C., and Aulakh, G. S. (1987). *Indigenous plants used in the control of diabetes*. Publication and Information Directorate, CSIR, New Delhi, 586.
77. Parasuraman, S., Balamurugan, S., Christopher, P. V., Petchi, R. R., Yeng, W. Y., Sujithra, J., and Vijaya, C. (2015). Evaluation of antidiabetic and antihyperlipidemic effects of hydroalcoholic extract of leaves of *Ocimum tenuiflorum* (Lamiaceae) and prediction of biological activity of its phytoconstituents. *Pharmacognosy research*, 7(2), 156.
78. Geeta Vasudevan, D. M., Kedlaya, R., Deepa, S., and Ballal, M. (2001). Activity of *Ocimum sanctum* (the traditional Indian medicinal plant) against the enteric pathogens. *Ind. J. Med. Sci*, 55(8), 434-438.
79. Khan, A., Ahmad, A., Manzoor, N., and Khan, L. A. (2010). Antifungal activities of *Ocimum sanctum* essential oil and its lead molecules. *Natural Product Communications*, 5(2), 1934578X1000500235.
80. Prashar, R., and Kumar, A. (1995). Chemopreventive action of *Ocimum sanctum* on 2, 12- dimethylbenz (a) anthracene DMBA-induced papillomagenesis in the skin of mice. *International journal of pharmacognosy: a journal of crude drug research*.
81. Prakash, J., and Gupta, S. K. (2000). Chemopreventive activity of *Ocimum sanctum* seed oil. *Journal of ethnopharmacology*, 72(1-2), 29-34.
82. Uma Devi, P., Ganasoundari, A., Vrinda, B., Srinivasan, K. K., and Unnikrishnan, M. K. (2000). Radiation protection by the *ocimum* flavonoids orientin and vicenin: mechanisms of action. *Radiation research*, 154(4), 455-460.
83. Karthikeyan, K., Gunasekaran, P., Ramamurthy, N., and Govindasamy, S. (1999). Anticancer activity of *Ocimum sanctum*. *Pharmaceutical biology*, 37(4), 285-290.
84. Aruna, K., and Sivaramakrishnan, V. M. (1992). Anticarcinogenic effects of some Indian plant products. *Food and Chemical Toxicology*, 30(11), 953-956.
85. Devi, P. U. (2001). Radioprotective, anticarcinogenic and antioxidant properties of the Indian holy basil, *Ocimum sanctum* (Tulasi).
86. Karthikeyan, K., Ravichandran, P., and Govindasamy, S. (1999). Chemopreventive effect of *Ocimum sanctum* on DMBA-induced hamster buccal pouch carcinogenesis. *Oral oncology*, 35(1), 112-119.
87. Prashar, R., Kumar, A., Hewer, A., Cole, K. J., Davis, W., and Phillips, D. H. (1998). Inhibition by an extract of *Ocimum sanctum* of DNA-binding activity of 7, 12-dimethylbenz [a] anthracene in rat hepatocytes in vitro. *Cancer letters*, 128(2), 155-160.
88. Somkuwar, A. P. (2003). Studies on anticancer effects of *Ocimum sanctum* and *Withania somnifera* on experimentally induced cancer in mice (Doctoral

- dissertation, Jawaharlal Nehru Krishi Viswavidyalaya; Jabalpur).
89. Devi, P. U., and Ganasoundari, A. (1995). Radioprotective effect of leaf extract of Indian medicinal plant *Ocimum sanctum*. *Indian Journal of Experimental Biology*, 33(3), 205- 208.
  90. Ganasoundari, A., Devi, P. U., and Rao, B. S. S. (1998). Enhancement of bone marrow radioprotection and reduction of WR-2721 toxicity by *Ocimum sanctum*. *Mutation Research/Fundamental and Molecular Mechanisms of Mutagenesis*, 397(2), 303-312.
  91. Chattopadhyay, R. R., Sarkar, S. K., Ganguly, S., Medda, C., and Basu, T. K. (1992). Hepatoprotective activity of *Ocimum Sanctum* leaf extract against Paracetamol included Hepatic damage in rats.
  92. Seethalakshmi, B., Narasappa, A. P., and Kenchaveerappa, S. (1982). Protective effect of *Ocimum sanctum* in experimental liver injury in albino rats. *Indian J Pharmacol*, 14, 63.
  93. Ahmed, M., Ahamed, R. N., Aladakatti, R. H., and Ghosesawar, M. G. (2002). Reversible anti-fertility effect of benzene extract of *Ocimum sanctum* leaves on sperm parameters and fructose content in rats. *Journal of Basic and Clinical Physiology and Pharmacology*, 13(1), 51-60.
  94. Singh, S., Taneja, M., and Majumdar, D. K. (2007). Biological activities of *Ocimum sanctum* L. fixed oil—An overview.
  95. Srinivas, N., Sali, K., and Bajoria, A. A. (2016). Therapeutic aspects of Tulsi unraveled: A review. *Journal of Indian Academy of oral medicine and radiology*, 28(1), 17.
  96. Bhargava, K. P., and Singh, N. (1981). Anti-stress activity of *Ocimum sanctum* Linn.
  97. Singh, S., and Majumdar, D. K. (1999). Evaluation of the gastric antiulcer activity of fixed oil of *Ocimum sanctum* (Holy Basil). *Journal of ethnopharmacology*, 65(1), 13-19.
  98. Singh, S., Rehan, H. M. S., and Majumdar, D. K. (2001). Effect of *Ocimum sanctum* fixed oil on blood pressure, blood clotting time and pentobarbitone-induced sleeping time. *Journal of ethnopharmacology*, 78(2-3), 139-143.
  99. Shah, S. M. A., Akram, M., Riaz, M., Munir, N., and Rasool, G. (2019). Cardioprotective potential of plant-derived molecules: a scientific and medicinal approach. *Dose-response*, 17(2), 1559325819852243.
  100. Sharma, M., Kishore, K., Gupta, S. K., Joshi, S., and Arya, D. S. (2001). Cardioprotective potential of *Ocimum sanctum* in isoproterenol induced myocardial infarction in rats. *Molecular and cellular biochemistry*, 225(1), 75-83.
  101. Kelm, M. A., Nair, M. G., Strasburg, G. M., and DeWitt, D. L. (2000). Antioxidant and cyclooxygenase inhibitory phenolic compounds from *Ocimum sanctum* Linn. *Phytomedicine*, 7(1), 7-13.
  102. Kapewangolo, P., Kandawa-Schulz, M., and Meyer, D. (2017). Anti-HIV activity of *Ocimum labiatum* extract and isolated pheophytin-a. *Molecules*, 22(11), 1763.
  103. Ling, A. P., Khoo, B. F., Seah, C. H., Foo, K. Y., Cheah, R. K., Chye, S. M., and Koh, R. Y. (2014, August). Inhibitory activities of methanol extracts of *Andrographis paniculata* and *Ocimum sanctum* against dengue-1 virus. In *International Conference on Biological Environmental and Food Engineering: Bali, Indonesia* (pp. 4-5).
  104. Hemalika, D. V. D., and Chandrika, U. G. Anti-dengue effects of medicinal plants: A review.
  105. Shree P, Mishra P, Selvaraj C, Singh SK, Chaube R, Garg N, Tripathi YB. Targeting COVID-19 (SARS-CoV-2) main protease through active phytochemicals of ayurvedic medicinal plants - *Withania somnifera* (Ashwagandha), *Tinospora cordifolia* (Giloy) and *Ocimum sanctum* (Tulsi) - a molecular docking study. *J Biomol Struct Dyn*. 2020 Aug 27;1-14. doi: 10.1080/07391102.2020.1810778. Epub ahead of print. PMID: 32851919; PMCID: PMC7484581.
  106. Singh, S., and Majumdar, D. K. (1995). Analgesic activity of *Ocimum sanctum* and its possible mechanism of action. *International journal of Pharmacognosy*, 33(3), 188-192.
  107. Mediratta, P. K., Sharma, K. K., and Singh, S. (2002). Evaluation of immunomodulatory potential of *Ocimum sanctum* seed oil and its possible mechanism of action. *Journal of Ethnopharmacology*, 80(1), 15-20.
  108. Mondal, S., Varma, S., Bamola, V. D., Naik, S. N., Mirdha, B. R., Padhi, M. M., ... and Mahapatra, S. C. (2011). Double-blinded randomized controlled trial for immunomodulatory effects of Tulsi (*Ocimum sanctum* Linn.) leaf extract on healthy volunteers. *Journal of ethnopharmacology*, 136(3), 452-456.
  109. Mediratta PK, Dewan V, Bhattacharya SK, Gupta VS, Maiti PC, Sen P. Effect of *Ocimum sanctum* Linn. on humoral immune responses. *Indian J Med Res*. 1988 Apr;87:384-6. PMID: 3169894.
  110. Jeba, C. R., Vaidyanathan, R., and Rameshkumar, G. (2011). Immunomodulatory activity of aqueous extract of *Ocimum sanctum* in rat. *Int J Pharm Biomed Res*, 2(1), 33- 8.
  111. Sharma, P., Kulshreshtha, S., and Sharma, A. L. (1998). Anti-cataract activity of *Ocimum sanctum* on experimental cataract. *Indian journal of pharmacology*, 30(1), 16.
  112. Kadian, R., and Parle, M. (2012). Therapeutic potential and phytopharmacology of tulsi. *International Journal of Pharmacy and Life Sciences*, 3(7).
  113. Singh, S., and Majumdar, D. K. (1996). Effect of fixed oil of *Ocimum sanctum* against experimentally induced arthritis and joint edema in laboratory animals. *International Journal of Pharmacognosy*, 34(3), 218-222.



114. Kadian, R., and Parle, M. (2012). Therapeutic potential and phytopharmacology of tulsi. *International Journal of Pharmacy and Life Sciences*, 3(7).
115. Upadhyay, R. K. (2017). Tulsi: A holy plant with high medicinal and therapeutic value. *International Journal of Green Pharmacy (IJGP)*, 11(01).
116. Singh, M., Hamid, A. A., Maurya, A. K., Prakash, O., Khan, F., Kumar, A., ... and Bawankule, D. U. (2014). Synthesis of diosgenin analogues as potential anti-inflammatory agents. *The Journal of steroid biochemistry and molecular biology*, 143, 323-333.