Anti-Diabetic Activity of *Murraya koenigii* – A Comprehensive Review

U. Vidhya Rekha a*, S. Bhuminathan b and P. Ravi Shankar c*

a Department of Public Health Dentistry, Sree Balaji Dental College and Hospital, Chennai, India. 
b Department of Prosthodontics, Sree Balaji Dental College and Hospital, Chennai, India. 
c Department of Public Health Dentistry, Rajas Dental College and Hospital, Kavalkinaru, Tirunelveli Dist, India.

Authors’ contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/JPRI/2021/v33i58B3422

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: https://www.sdiarticle5.com/review-history/78628

Received 07 October 2021 
Accepted 14 December 2021 
Published 16 December 2021

ABSTRACT

Diabetes mellitus, one of the noncommunicable illnesses, is a severe problem worldwide as one of the leading causes of death. Because existing synthetic medications have various drawbacks, researchers are still looking for better anti-hyperglycemic treatments. Plants have been used in ancient medicine for thousands of years. India is the biggest producer of medicinal plants and is aptly regarded as the "World's Botanical Garden." *Murraya koenigii* Linn, also known as Meethi neem, is a Rutaceae plant. Curry trees are unique to India and likely found almost everywhere else on the subcontinent, except in the Himalayan highlands. For centuries, curry leaves were used as an antiemetic, diarrhea remedy, febrifuge, and blood purifier. Curry leaves are useful as an antioxidant, anti-diabetic, antibacterial, antihypertensive, cytotoxic, and in treating bronchial respiratory problems. Traditionally, the leaves were utilized as a spice in curries as well as other dishes. It includes coumarins and derivatives, alkaloids, flavonoids, phenolic compounds, and essential oil. Numerous studies have found that these phytochemicals have a significant effect on type 2 diabetes. This review focuses on this plant's anti-diabetic action and concludes that it has the potential to be evaluated as a candidate for developing a new diabetes mellitus medication.
Keywords: Medicinal plants; Murraya koenigii; diabetes mellitus.

1. INTRODUCTION

Diabetes mellitus, among the most frequent endocrine and metabolic illnesses, has caused enormous morbidity and death as a result of microvascular (retinopathy, neuropathy, and nephropathy) and macrovascular (heart attack, stroke, and peripheral vascular disease) effects. In human bodies, anti-oxidative processes, both enzymatic and non-enzymatic, contribute to the reduction of reactive oxygen species, which are linked to a number of degenerative disorders, including diabetes [1]. The sickness is spreading swiftly over the world and is impacting people in every country. Diabetes patients have elevated blood glucose levels due to insulin insufficiency [2]. Type 2 diabetes, also called the non diabetic Mellitus, has to be the most common type of diabetes, accounting for 90 percent to 95 percent of instances in which the body fails to make sufficient insulin or use it properly [3]. According to the World Health Organization, the diabetes population will reach 300 million or more by [4]. Insulin and various oral anti-diabetic medications, including sulfonylureas, biguanides, and glinides, are currently available for diabetes treatment. Many of them have a variety of significant side effects; as a result, one of the most critical areas of research is the search for more effective and safer hypoglycemic agents [5].

2. COST OF DIABETES

Numerous research on the cost of sickness have been conducted as a result of the economic burden of diabetes mellitus. Diabetes costs may be divided into three categories: direct costs, indirect costs, and intangible costs [6,7]. Both direct health care expenditures (diagnostic, treatment, care, and preventive) and direct non-health care costs are included in direct costs (transport, housekeeping, social service and legal cost). With complications, the total direct cost is ₹ 28,888/- per year [8]. Absenteeism, lost output, and incapacity are all examples of indirect costs. With complexity, the total indirect cost is ₹1746/- per annum [9]. Finally, intangible costs include costs associated with social isolation and dependency, low socioeconomic status, mental health and behavioural disorders, and a reduction in quality of life. [10].

3. PLANT AS ALTERNATIVE SOURCE

The hypoglycemic impact of various plants used as anti-diabetic agents has been established, and the mechanisms underlying this effect are being researched. This article discusses natural compounds with anti-diabetic characteristics that operate as insulin-mimetic or secretagogues. Traditional remedies derived from commonly available medicinal plants hold considerable promise for developing new anti-diabetic medications [11].

Certain medicinal herbs have recently been described to be effective in the treatment of diabetes throughout the world, and they have been used empirically in anti-diabetic and anti-hyperlipidemic drugs. Plants' anti-hyperglycemic action is mainly related to their capacity to restore pancreatic tissue function by increasing insulin secretion, inhibiting glucose absorption in the intestine, or facilitating metabolites in insulin-dependent activities. Even though literature lists over 400 plant species with hypoglycemic action, hunting for new anti-diabetic medications from natural plants remains appealing since they contain chemicals that have alternative and harmless effects on diabetes mellitus.

Plant-derived active components that have shown action as in treatment of diabetes include alkaloids, glycosides, galactomannan gun, polysaccharides, peptidoglycans, hypoglycin, guanidine, steroids, sugars, glycopeptides, terpenoids, amino acids, and inorganic ions [12].

Curry leaves are Murraya koenigii (M. koenigii) (L) Spreng of the Rutaceae family. M. koenigii is found all across the world's tropical and subtropical regions. [13]. Only two Murraya species, M. koenigii and M. paniculata are found in India, out of 14 worldwide. M. koenigii is more important because of the wide range of traditional therapeutic characteristics it possesses. This plant has been utilized in various ways for millennia and is known as "krishnamimba" in Indian Ayurvedic medicine [14]. M. koenigii's leaves, roots, bark, and fruit have been demonstrated to support a range of biological activities. Even after drying, aromatic bioactive components in M. koenigii leaves retain their flavor and other properties [15]. The leaves of M. koenigii have a slightly bitter taste, a pungent odor, and are somewhat acidic. They are used as antihelminthics, analgesics, digestive, and appetizers in Indian cuisine [16]. Piles, inflammation, itching, fresh cuts, diarrhea, bruising, and edema are treated using M.
Koenig's green leaves. To some extent, the roots are purgative. They are stimulating and are used to treat aches and pains in the body. The bark can be used to treat snakebites [17]. The essential oil obtained from M. koenigii leaves has been proven in animal models to have antioxidative and hepatoprotective properties [18] antibacterial, antifungal, anti-inflammatory, and nephroprotective effects [19]. Several chemical elements of distinct carbazole alkaloids and other significant metabolites, such as terpenoids, flavonoids, phenolics, carbohydrates, carotenoids, vitamins, and nicotinic acid, have been attributed to the therapeutic qualities of M. koenigii from various regions of the plant.

M. koenigii has received increased interest in traditional medicines and home cures in recent years. On the other hand, few research have been undertaken to assess M. koenigii’s pharmacological and therapeutic usefulness in improving health and healing sickness [20]. This review aims to present previous and existing major studies on M. koenigii activity in diabetics.

4. TRADITIONAL AND MEDICINAL USES OF M. KOENIGII

M. koenigii essential oils and fresh leaf powder can be used to season foods and make ready-to-eat meals. The essential oil from leaf extracts can also be employed as a perfume, and taste agent in traditional practise due to its increased antibacterial activity [21]. Fresh curry leaves are cooked with coconut oil until they have been crushed to a black residue to make a great hair tonic for restoring normal hair tone and promoting hair development. Curry leaves have long been used as an antidiarrheal, antifungal, blood purifying, anti-inflammatory, and anti-depressant medication, either whole or in little amounts. [22].

Different plant parts, like the leaves, roots, and bark, could be used as tonics to induce digestion and flatulence or as antiemetics [23]. The leaves turn unpleasant to the taste after infusion and are useful in lowering fever. The root's juice is used to treat kidney pain [24]. The leaves and roots can be used as an antihistamine, analgesic, piles treatment, body heat reducer, and thirst quencher, as well as to relieve inflammation and irritation. They're also suitable for treating leucoderma and blood problems. The green leaves can be consumed raw to treat diarrhea, and the paste made from boiling the leaves in milk can be used to treat toxic bites and eruptions. [25].

5. OTHER USES

Using essential oils Murraya koenigii is utilised in formulations as a sun protection and erythema agent [26]. Curry leaf oil with your usual skin care cream or lotion helps to heal skin problems such as pimples, athlete's foot, ringworm, itches, acne, boils, and septic wounds and burns by applying it to the afflicted region [27]. Due to the active constituents -pinene and Caryophyllene, essential oils of Murraya koenigii were tested for toxicity and repellent action against Callosobruchus maculatus[28]. Murraya koenigii structure-function investigations suggest that the protein has a trypsin inhibitory effect as a compact structure with helical content as temperature rises. In Dalton's Ascitic Lymphoma, the effects of Murraya koenigii column extract demonstrate a protective effect [29]. The ability to include dried curry leaf powder into everyday foods boosts micronutrient sources[30]. Murraya koenigii aqueous extract has larvicidal, pupicidal repellent, and anti-vector activities against larvae and pupae. [31] leaf oil's high vitamin A and calcium content is used to treat osteoporosis, calcium shortage, and cancer radiation and chemotherapy treatments [32]. Murraya koenigii is used to cure or prevent orofacial dyskinesia (OD), which is caused by neuroleptics [33]. The anti-Trichomonas gallinae action of carbazole alkaloids derived from Murraya koenigii extract and its derivatives [34]. Curry leaves and essential oil are used both orally and topically for healthy, long, strong, and glossy hair. A balanced diet with an equal percentage of vitamins, minerals, iron, and other nutrients is needed to keep hair healthy. [35]. Curry leaf oil aids in muscle and tissue contraction. Curry leaf extract aids in the reduction of white areas on the body and helps with pigmentation. Curry leaf fresh leaves, dried leaf powder, and essential oil are commonly used to flavour soups, curries, fish, meat, and egg meals, as well as classic curry powder mixes, seasoning, and ready-to-use other culinary preparations. Murraya koenigii essential oil is used in the soap and cosmetic sector for aromatherapy. Murraya koenigii is used to treat bruises, eruptions, and deadly animal attacks. [36].

6. PHYTOCHEMISTRY OF M. KOENIGII

The leaves, roots and stem bark of M. koenigii have been used to isolate a variety of phytochemicals. Alkaloids, flavonoids, terpenoids, and polyphenols have been found in M. koenigii extracts of leaves, roots, stem bark,
fruits, and seeds. Moisture is 63.2 percent, protein is 8.8%, carbohydrate is 39.4%, total nitrogen is 1.15 percent, fat is 6.15 percent, total sugars are 18.92 percent, starch is 14.6 percent, and crude fiber is 6.8 percent in the plant leaves. Vitamin A (B-carotene): 6.04 0.02 mg/100 g; vitamin B3, (niacin): 2.73 0.02 mg/100 g; vitamin B1 (thiamin): 0.89 0.01 mg/100 g; calcium: 19.73 0.02 mg/100 g; magnesium: 49.06 0.02 mg/100 g; sodium: 16.50 0.21 mg/100 g [37] The alcohol-soluble extract has a value of 1.82 percent, ash has a value of 13.06 percent, acid-insoluble ash has a value of 1.35 percent, cold water (20 °C) extractive has a value of 27.33 percent, and maximum of hot-water-soluble extractive has a value of 33.45 percent. Carbazole alkaloids, essential oils, terpenoids, and flavonoids all play important functions in the human body. List of main phyto compounds present in M. koenigii was listed in Table 1.

7. ANTIDIABETIC ACTIVITY OF M. Koenigii

Because of their low cost, medicinal plants are particularly useful in managing diabetes mellitus in developing countries. Diabetes mellitus, a metabolic disorder, is rapidly being a major public health concern. In recent years, numerous phytochemicals with anti-diabetic properties have been found in plants. M. koenigii leaf alkaloids were examined and found to inhibit the aldose reductase enzyme, glucose consumption, and other enzyme systems, potentially extending anti-diabetic benefits[38]. The -glucosidase inhibitory property of M. koenigii was examined, and it was discovered to inhibit glycosidase. Patients with type 2 diabetes are commonly treated with alpha-glucosidase inhibitors [39]. According to one study, an ethanolic extract of M. koenigii reduced blood glucose levels significantly, and this action of M. koenigii reducing blood glucose is mediated by antioxidant properties and insulin-mimetic effects. M. koenigii also demonstrated a high antioxidant effect, lowering MDA levels, increasing GSH levels, and significantly lowering the homeostatic model assessment (HOMA)-insulin resistance index. Overall, M. koenigii appears to have anti-diabetic and antioxidant properties in rats. [40].

8. HYPOGLYCEMIC ACTIVITY

Feeding the leaves to rats resulted in hypoglycemia because it enhanced hepatic glycogenesis, as evidenced by increased glycogen synthetase activity. The activity of glycogen phosphorylase and gluconogenic enzymes reduced, indicating a reduction in glycogenolysis and gluconeogenesis. [41].

The hypoglycemic effect can be achieved by increasing insulin production from [beta]-cells of Langerhans islets in the pancreas or emancipating insulin from its bound state [42]. The antioxidant defence system of plasma and pancreas, as well as the probable protective impact of M. koenigii leaf extract against -cell damage, were examined in streptozotocin-induced diabetic rats. It was determined that M. koenigii therapy protects against diabetes by lowering oxidative stress and pancreatic-cell damage.

The effects of M. koenigii leaves were researched by Arulselven and Subramanian [43]. Streptozotocin-induced diabetic rats were used in the experiment, and they were given 200 mg [kg.sup.-1] M. koenigii leaves for 30 days. M. koenigii dramatically reduced blood glucose and glycosylated hemoglobin levels while significantly increasing insulin and liver glycogen levels, according to the findings. It also decreased lactate dehydrogenase, glucose-6-phosphatase, fructose-1,6-diphosphatase, and glycogen phosphorylase activities while increasing hexokinase and pyruvate kinase activities. The effects of M. koenigii fruit juice were researched by Tembhurne and Sakarkar [44]. They used alloxan-induced diabetic mice treated for 15 days with 2.5 and 5.0 ml/kg M. koenigii fruit juice.

The hypoglycaemic effect of M. koenigii leaf extracts, as well as the number of spices employed, was investigated, proving that they can be used as an effective anti-diabetic diet [45].

M. koenigii leaf extract reduced blood glucose levels by 13.1, 16.3, and 21.4 percent and 3.2, 5.58, and 8.21 percent for mild and moderate diabetes produced by alloxan in rats fed the extract as a meal, demonstrating its potential as an antihyperglycaemic agent[46].

The effect of an aqueous extract of M. koenigii leaves on hypoglycaemic activity in normal and alloxan-induced diabetic rabbits was compared to the impact of a common hypoglycaemic medication, tolbutamide. In both normal and diabetic rats, a single treatment of varied dose levels (200, 300, and 400 mg/kg) of the aqueous extract resulted in a reduction in blood glucose levels [47].
Table 1. The pharmacological activity of *M. koenigii*'s key bioactive substances

| S. No | Compound Name       | Activity                                                                 |
|-------|---------------------|--------------------------------------------------------------------------|
| 1     | Mahanine            | Cytotoxicity, anti-microbial, and anti-cancer                            |
| 2     | Mahanimbine         | Cytotoxicity, anti-oxidant, anti-microbial, anti-diabetic, and hyperlipidemic |
| 3     | Isomahanine         | Cytotoxicity, anti-oxidant, anti-microbial, anti-diabetic, and hyperlipidemic |
| 4     | koenimbine          | Cytotoxicity and anti-diarrhea                                           |
| 5     | Girinimbine         | Anti-tumor                                                               |
| 6     | Isolongifolene      | Anti-oxidant and neuroprotective                                         |
| 7     | Pyrayafoline D      | Anti-cancer and anti-bacterial                                          |
| 8     | Murrayafoline       | Cytotoxicity and anti-inflammatory                                      |
| 9     | Murrayazoline       | Cytotoxicity and anti-tumor                                              |
| 10    | Koenoline           | Cytotoxicity                                                             |
| 11    | 9-formyl-3-methyl carbazole | Anti-oxidant          |
| 12    | O-Methylmurrayamine | Anti-oxidant and neuroprotective                                         |
| 13    | Koenine             | Anti-oxidant                                                             |
| 14    | Koenigine           | Anti-oxidant                                                             |
| 15    | Mukonicine          | Anti-oxidant                                                             |
| 16    | Mahanimbinine       | Anti-oxidant, anti-microbial, anti-diabetic, and hyperlipidemic          |
| 17    | Murrayacinine       | Anti-oxidant, anti-microbial, anti-diabetic, and hyperlipidemic          |
| 18    | Mahanimboline       | Cytotoxicity, anti-oxidant, anti-microbial, anti-diabetic, and hyperlipidemic |
| 19    | Mukoeic acid        | Anti-oxidant                                                             |
| 20    | Murrayanine         | Anti-oxidant                                                             |
Curry leaf extract has been shown to lower blood cholesterol and blood glucose levels in diabetic rats, as well as reduce body weight after therapy [48].

For 30 days, oral administration of an ethanolic extract of *M. koenigii* to Streptozotocin-induced diabetic rats dramatically reduced blood glucose, glycosylated hemoglobin, urea, uric acid, and creatinine levels in the diabetic treatment group of animals [49].

For a brief period of 6 hours, the aqueous extract of *M. koenigii* had a favorable effect in reducing the severity of diabetes in alloxan and normal induced diabetic rabbits 48. *M. koenigii* considerably reduced blood glucose levels, according to the findings. Many types of research [50] examined flavonoids, quercin, metformin, quinolizidine, anthocyanin, catechin, flavone, phenylpropanoids, lipoic acid, and coumarin as the most phytochemical compounds having anti-diabetes activity.

Traditional or alternative therapy, in addition to mainstream pharmaceuticals, plays a crucial part in the treatment of diabetes mellitus. It must understand how to use it and what phytochemical ingredients are present. The goal of this review study was to compile the new medicinal plant, *M. paniculata*, as the therapy of choice. All of this data will aid researchers in their investigation of the scientific evidence.

Alkaloids found in the leaves of *M. koenigii* have been studied and shown to inhibit the aldose reductase enzyme, glucose consumption, and other enzyme systems, potentially prolonging antidiabetic effects[51]. *M. koenigii* was investigated for its ability to inhibit glycosidase, and it was discovered to do so. Alphaglucosidase inhibitors are commonly used to treat type 2 diabetic patients. [52]. An ethanolic extract of *M. koenigii* exhibited a considerable reduction in blood glucose levels, according to one research, and this action of *M. koenigii* decreasing blood glucose is mediated by antioxidant qualities and insulin mimetic effects. Furthermore, *M. koenigii* had a strong antioxidant impact, lowering malondialdehyde (MDA), boosting GSH, and dramatically lowering the homeostatic model assessment (HOMA) insulin resistance score. Overall, *M. koenigii* appears to have anti-diabetic and antioxidant properties in rats. [53].

9. ADVANTAGES OF *M. Koenigii*

The best health advantages of Curry Leaves are listed here:

- Curry Leaves Aid in Cholesterol Reduction
- Curry Leaves Aids Digestion
- Curry Leaves Benefits the Liver
- Curry Leaves Promotes Hair Growth
- Curry Leaves Improves Eye Health
- Curry Leaves Eradicates Bacteria
- Curry Leaves Promotes Weight Loss
- Curry Leaves Prevents Anemia

10. DISADVANTAGES OF *M. Koenigii*

Although no harm has been recorded from curry leaves, it may be detrimental in some instances or at excessive quantities.

- Some individuals may experience allergic reactions. Its usage should be ceased in such a case.
- Pregnant and breastfeeding women should see a doctor before using it, as some of its negative effects are common in this condition.
- At the same time, in some situations, the use of its oil can weaken the hair roots and cause them to fall out.

11. CONCLUSION

The present review discusses *M. koenigii*s medicinal uses, phytochemical constituents, and pharmacological qualities, with a focus on its anti-diabetic properties. *M. koenigii* contains alkaloids, polyphenols, terpenoids, and flavonoids, among other bioactive substances. *M. koenigii* and its substances appear to have anticarcinogenic, proapoptotic, antiangiogenic, antimitastatic, immunomodulatory, and antioxidant properties. The broad activity of *M. koenigii* and its derivatives in cell signalling pathways at multiple levels in various illnesses illustrates the molecular processes behind these activities. *M. koenigii* and its derivatives reduce oxidative stress, neurotoxicity, neuroinflammation, neuronal loss, and cognitive dysfunctions. However, like other polyphenols, *M. koenigii*s actions are restricted to some extent by its bioavailability, and in such cases, increased efficiency should be pursued. As a result, future research should involve additional experimental studies on improving bioavailability and efficiency in clinical trials.

CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.
COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Ponnulakshmi R, Shyamaladevi B, Vijayalakshmi P, Selvaraj J. In silico and in vivo analysis to identify the antidiabetic activity of beta sitosterol in adipose tissue of high fat diet and sucrose induced type-2 diabetic experimental rats. Toxicol Mech Methods. 2019;29(4):276-290. DOI: 10.1080/15376516.2018.1545815. Epub 2019 Jan 15. PMID: 30461321.

2. Ponnusamy S, Ravindran R, Zinnarde S, Bhargava S, Kumar AR. Evaluation of traditional Indian antidiabetic medicinal plants for human pancreatic amylase inhibitory effect in vitro. Evid Based Complement Alternat Med. 2011;2011:515647.

3. Li WL, Zheng HC, Bukuru J, De Kimpe N. Natural medicines used in the traditional Chinese medical system for therapy of diabetes mellitus. J Ethnopharmacol. 2004;92(1):1–21.

4. Sy GY, Cissé A, Nongonierma RB, Sarr M, Mboj NA, Faye B. Hypoglycaemic and antidiabetic activity of acetonic extract of Vernonia colorata leaves in normoglycaemic and alloxan-induced diabetic rats. J Ethnopharmacol. 2005;98(1–2):171–175.

5. Saxena A, Vikram NK. Role of selected Indian plants in management of type 2 diabetes: a review. J Altern Complement Med. 2004;10(2):369–378.

6. Pagano E, Brunetti M, Tediosi F, Garattini L. Costs of diabetes. A methodological analysis of the literature. Pharmaco economics. 1999;15(6):583-95.

7. Sobocki P, Lekander I, Borgström F, Ström O, Runeson B. The economic burden of depression in Sweden from 1997 to 2005. Eur Psychiatry. 2007;22(3):146-52.

8. American Diabetes Association. Diagnosis and classification of diabetes mellitus. Diabetes Care. 2013;36 Suppl 1(Suppl 1):S67-74.

9. Glynn LG, Valderas JM, Healy P, Burke E, Newell J, Gillespie P, Murphy AW. The prevalence of multi-morbidity in primary care and its effect on health care utilization and cost. Fam Pract. 2011;28(5):516–523.

10. Rodwin BA, Spruill TM, Ladapo JA. Economics of psychosocial factors in patients with cardiovascular disease. Prog Cardiovasc Dis. 2013;55(6):563–573.

11. Jung M, Park M, Lee HC, Kang YH, Kang ES, Kim SK. Antidiabetic agents from medicinal plants. Curr Med Chem. 2006;13(10):1203–1218.

12. Grover JK, Yadav S, Vats V. Medicinal plants of India with anti-diabetic potential. J Ethnopharmacol. 2002;81(1):81–100.

13. Wojdylo A., Oszmiański J., Czemerys R. Antioxidant activity and phenolic compounds in 32 selected herbs. Food Chem. 2007;105:140–149.

14. Ahluwalia V., Sisodia R., Wala S., Sati O.P., Kumar J., Kundu A. Chemical analysis of essential oils of Eupatorium adenophorum and their antimicrobial, antioxidant and phytotoxic properties. J. Pest Sci. 2014;87:341–349

15. Amna U., Halimatussakdiah P.W., Saidi N., Nasution R. Evaluation of cytotoxic activity from Temurui (Murraya koenigii [Linn.] Spreng) leaf extracts against HeLa cell line using MTT assay. J. Adv. Pharm. Technol. Res. 2019;10:51–55.

16. Desai SN, Patel DK, Devkar RV, Patel PV, Ramachandran AV. Hepatoprotective potential of polyphenol rich extract of Murraya koenigii L.: An In vivo study. Food Chem. Toxicol. 2012;50:310–314.

17. Gajaria TK, Patel DK, Devkar RV, Ramachandran AV. Flavonoid rich extract of Murraya koenigii alleviates in-vitro LDL oxidation and oxidized LDL induced apoptosis in raw 264.7 Murine macrophage cells. J. Food Sci. Technol. 2015;52:3367–3375.

18. Ma QQ, Xu K, Sang ZP, Wei RR, Liu WM, Su YL, Yang JB, Wang AG, Ji TF, Li LJ. Alkenes with antioxidative activities from Murraya koenigii (L.) Spreng. Bioorg. Med. Chem. Lett. 2016;26:799–803.

19. Tripathi Y., Anjum N., Rana A. Chemical Composition and In vitro Antifungal and Antioxidant Activities of Essential Oil from Murraya koenigii (L.) Spreng. Leaves. Asian J. Biomed. Pharm. Sci. 2018;8:6–13

20. Dar RA, Shahnawaz M, Qazi PH, Qazi H. General overview of medicinal plants: A review. J. Phytopharm. 2017;6:349–351.

21. Erkan N, Tao Z, Vasantha Rupasinghe HP, Uysal B, Oksal BS. Antibacterial activities of essential oils extracted from leaves of Murraya koenigii by solvent-free microwave extraction and hydro-distillation.
24. Tembhurne SV, Sakarkar DM. Hypoglycemic effects of fruit juice of Murraya koenigii (L) in alloxan induced diabetic mice. Int. J. PharmTech Res. 2009;1:1589–1593

25. Sim KM, Teh HM. A new carbazole alkaloid from the leaves of Malayan Murraya koenigii. J. Asian Nat. Prod. Res. 2011;13:972–975

26. Prakash V, Natarajan CP. Studies on Curry Leaf. Food Sci and Technol. 1974;11(6):284–286.

27. Dasgupta T, Rao AR, Yadava PK. Chemomodulatory action of curry leaf (Murraya koenigii) extract on hepatic and extrahepatic xenobiotic metabolising enzymes, antioxidant levels, lipid peroxidation, skin and forestomach papillomagenesis. Nutrition Research. 2003;23:1427

28. Jamil R, Nasir NN, Ramli H, et al. Extraction of essential oil from Murraya koenigii leaves: potential study for application as natural-based insect repellent. Journal of Engineering and Applied Sciences. 2016;11(4):1–5.

29. Gupta V, Sharma M. Protective effect of Murraya koenigii on lipid peroxide formation in isolated rat liver homogenate. Int J Pharma Bio Sci. 2010;1(3):1–6.

30. Harve G, Kamath V. Larvicidal activity of plant extracts used alone and in combination with known synthetic larvicidal agents against Aedes aegypti. Ind J Exp Biol. 2004;42(12):1216–1229.

31. Arulselvach P, Subramanian SP. Beneficial effects of Murraya koenigii leaves on antioxidant defence system and ultra structural changes of pancreatic beta cells in experimental diabetes in rats. Chem Biol Interact. 2007;165(2):155–164.

32. Srinivasan K. Plant foods in the management of diabetes mellitus, spices as beneficial antidiabetic food adjuncts. Int J Food Sci Nutr. 2005;56(6):399–414.

33. Xie JT, Chang CZ, Mehendale SR, et al. Curry leaf reduces blood glucose and blood cholesterol level in ob/ob mice. Am J Chin Med. 2006;34(2):279–284.

34. Shree C, Islam A, Ahmad F, et al. Structure function studies of Murraya koenigii trypsin inhibitor revealed a stable core beta sheet structure surrounded by alpha helices with a possible role for alpha helix in inhibitory function. Int J Biol Macromol. 2007;41(4):410–414.

35. Debosree G, Syed BF, Elina M, et al. Protective effect of aqueous leaf extract of murraya koenigii against lead induced oxidative tress in rat liver, heart and kidney: a dose response study. Asian J Pharm Clin. 2012;5:54–59.

36. Mittal J, Sharma MM, Batra A. Tinospora cordifolia: a multipurpose medicinal plant- A review. Journal of Medicinal Plants Studies. 2014;2(2):32–47.

37. Igara C, Omoboyowa D, Ahuchaogu A, Orji N, Ndukwe M. Phytochemical and nutritional profile of Murraya koenigii (Linn) Spreng leaf. J. Pharmacogn. Phytochem. 2016;5:7–9

38. Patel DK, Kumar R, Laloo D, Hemalatha S. Natural medicines from plant source used for therapy of diabetes mellitus: An overview of its pharmacological aspects. Asian Pac. J. Trop. Dis. 2012;2:139–150.

39. Gul MZ, Attuluri V, Qureshi IA, Ghazi IA. Antioxidant and α-glucosidase inhibitory activities of Murraya koenigii leaf extracts. Pharmacogn. J. 2012;4:65–72.

40. Husna F., Suyatna F.D., Arozal W., Poerwaningsih E.H. Anti -trypsin inhibitor revealed a stable core beta sheet structure surrounded by alpha helices with a possible role for alpha helix in inhibitory function. Int J Biol Macromol. 2007;41(4):410–414.

41. Fiebig Manfred, Pezzuto John M., Soejarto D., Plant Anticancer Agents. Part 40, Koenoline A Further Cytotoxic Carbazole Alkaloid From Murraya koenigii, Phytochemistry, 1985; 24(12):3041-3043

42. Gautam MK, Gupta A, Rao CV, Goel RK. Antihyperglycemic and antioxidant potential of Murraya paniculata Linn. Leaves: a preclinical study. Journal of Pharmacy Research. 2012c; 5:1334-1337.
43. Arulselven P, Subramanian S. Effect of Murraya koenigii leaf extract on carbohydrate metabolism studied in streptozotocin induced diabetic rats. International Journal of Biological Chemistry. 2007;1:21-28.

44. Tembhrune SV, Sakarkar DM. Hypoglycemic effects of fruit juice of Murraya koenigii (L) in alloxan induced diabetic mice. International Journal of PharmTech Research. 2009;1:1589-1593.

45. Srinivasan K. Plant foods in the management of Diabetes Mellitus, spices as beneficial antidiabetic food adjuncts. Int. J. Food Sci. Nutr. 2005;56(6):399-414

46. Yadav SP, Vats V, Ammini AC, Grover JK. Brassica juncea significantly the development of insulin resistance in rats fed fructose enriched diet. J Ethnopharmacol. 2004;93(1):113-116.

47. Kesari AN, Gupta RK, Watal G. Hypoglycaemic effects of Murraya koenigii on normal and alloxan diabetic rabbits. J. Ethnopharmacol. 2005;97(2):247-51

48. Xie JT, Chang CZ, Mehendale SR, Ambihipahar R, Ambihipahar U, Fong HH et al. Curry leaf reduces blood glucose and blood cholesterol level in ob/ob mice. Am J Chin Med 2006;34(2):279-84.

49. Aruselvan P, Senthil KGP, Satish KD, Subramanian S. Antidiabetic effect of Murraya koenigii leaves on streptozotocin induced diabetic rats. Pharmazie. 2006;61(10): 874-7.

50. Harve G, Kamath V. Larvicidal activity of plant extracts used alone and in combination with known synthetic larvicidal agents against Aedes Aegypti. Ind J Exp. Biol. 2004;42(12):1216-9.

51. Patel DK, Kumar R, Laloo D, Hemalatha S. Natural medicines from plant source used for therapy of diabetes mellitus: An overview of its pharmacological aspects. Asian Pac. J. Trop. Dis. 2012;2:139–150

52. Gul MZ, Attuluri V, Qureshi IA, Ghazi IA. Antioxidant and α-glucosidase inhibitory activities of Murraya koenigii leaf extracts. Pharmacogn. J. 2012;4:65–72.

53. Husna F, Suyatna FD, Arozal W, Poerwaningsih, E.H. Anti-Diabetic Potential of Murraya koenigii (L) and its Antioxidant Capacity in Nicotinamide-Streptozotocin Induced Diabetic Rats. Drug Res. (Stuttg). 2018;68:631–636.

Peer-review history:
The peer review history for this paper can be accessed here:
https://www.sdiarticle5.com/review-history/78628