

# Systematic Review

## Exploring the Impact of Medicinal Plants on Cardiac Health Compared to Standard Pharmaceuticals Available in Bangladesh: A Systematic Review

### ABSTRACT

**Aims:** Medicinal plants have long been an essential part of healthcare in many cultures with a variety of bioactive chemicals that contribute to different pharmacological effects. Crude medicines, which are frequently refined or utilized in their natural state to treat a variety of ailments, can be derived from these plants. The potential of medicinal plants in medication development and discovery has been acknowledged more and more by contemporary scientific research, especially when it comes to disorders for which there are few available treatments. Numerous chemicals originating from plants have been found to possess antibacterial, anti-inflammatory, antioxidant, and cardioprotective qualities, making them potential candidates for use in pharmaceutical applications.

**Methodology:**

By exploring several published articles in reputed journals, we have listed out the medicinal plants available in Bangladesh that have various pharmacological effects, especially in cardiovascular health, and tried to compare them with the available pharmaceutical drugs in the Bangladesh market that are being widely used as blood thinner and lipid-lowering agents.

**Results:** Twenty-one medicinal plants have been found to have positive effects on cardiovascular health. On the other hand, despite being effective in their intended pharmacological actions, traditional cardiovascular medications including lipid-lowering medicines and anticoagulants are frequently linked to a variety of adverse effects. This presents a significant chance to investigate these medicinal plants' potential as alternative or complementary therapies that might provide comparable therapeutic advantages with fewer side effects.

**Keywords:** Anti-coagulation, hypolipidemic, cardiovascular, lipid-lowering agent.

### INTRODUCTION

The plants that have therapeutic properties or exert helpful medicinal impact on the living body are usually referred to as "Medicinal Plant" (World Health Organization, 2006). According to the World Health Organization, a medicinal plant is any plant that in one or a lot of its organs, contains substances that will be used for therapeutic functions or that could be a precursor for the synthesis of helpful medication (World Health Organization, 1991). Medicinal plants could also be outlined as a bunch of plants that possess some special properties or virtues that qualify them as articles of medicine and therapeutic agents and square measures used for healthful functions (Griggs, 1997). Medicinal plants are used as natural medicinal products. This observation has been occurring since prehistoric times (Li,

2000). Modern approaches to deciding the medicinal properties of plants involve cooperative efforts that will embody anthropologists, pharmacists, pharmaceuticals, chemists, and physicians (Bailey, 2002).

## **HISTORY OF MEDICINAL PLANTS**

Medicinal plants are a large group of plants that are used for therapeutic or prophylactic purposes in medicine or veterinary practice. Literary sources attest to the use of medicinal plants in the early years of the Common Era in ancient Assyria, Egypt, India, and China around 3000 B.C., Iran, Greece, and Rome, and in the Middle Ages in Arab countries, Middle Asia, Georgia, Armenia, and Europe (Griggs, 1997; Li, 2000). After the formation of the Aptekarsky Prikaz (pharmaceutical department) at the beginning of the 17th century, the gathering of medicinal herbs in Russia was organized and their cultivation began (Bailey, 2002; World Health Organization, 2006).

## **AVAILABLE MEDICINAL PLANTS IN BANGLADESH**

Bangladesh has more than 700 medicinal plant species. In Bangladesh, these plants are a habitat and the use of medicinal plants here is growing day by day. More than 80% of people here reside in rural areas and use these medicinal plants as primary healthcare items. Bangladesh is home to nearly 6500 plant species. The Bangladesh National Herbarium (BNH) first drafted a list of 192 species of plants in our country. BNH has published a pictorial book on 700 plant species used by local people living in the Chittagong Hill Tract area. The Bangladesh Agricultural Research Institute (BARI) reports that 722 different types of medicinal plants may be found in Bangladesh. Bangladesh uses 700 plants for medical reasons, compared to 4,000 in India. 255 of these plants are used by producers of Unani and Ayurvedic medications (Alam, 2010; Rahman, 2006).

## **STATUS OF MEDICINAL PLANTS IN WORLD MARKET**

China, France, Germany, Italy, Japan, Spain, the UK, and the US are the main global markets for MAPs. Japan has the highest botanical medicine intake per capita in the world. The International Council for Medicinal and Aromatic Plants predicts global growth to be about 8-10 percent per year in 2001 and 2002. The world demand for herbal medicines was US\$19.4 billion in 1999, with Europe leading the way (US\$6.7 billion), followed by Asia (US\$5.1 billion), North America (US\$4.0 billion), Japan (US\$2.2 billion) and the rest of the world (US\$1.4 billion). Japan's botanical medicine market was estimated at US\$2.4 billion in 1996 (Griggs, 1997; Li, 2000; World Health Organization, 2006).

## **STATUS OF CARDIOVASCULAR DISEASES IN BANGLADESH:**

In Bangladesh, cardiovascular diseases are an extremely prevalent contributor to death. Understanding this fact, both the government and private sectors are working hard to tackle these diseases. In recent decades, important progress has been made in the management of cardiac diseases in the world. New medical and therapeutic methods are routinely implemented to keep pace with the environment. In addition to these pharmacological treatments, almost all kinds of interventional management are being practiced in the region. In Bangladesh, except for a few complex surgeries, major cardiac and vascular surgeries are now possible. In addition, cardiac care centers, previously confined to the capital city, are now being decentralized all over the world. The recommended population of cardiologists by various authorities is 2.6-4.2/100,000; the ratio in Bangladesh in 2007 was around 0.27/100,000, which is very inadequate. The average population ratio of beds in the coronary care unit is 4-5/100,000; the ratio in Bangladesh was about 0.13/100,000 in 2007, which is again insufficient. However, the number of qualified specialists, hospitals, and cardiovascular treatment institutions is rising rapidly in Bangladesh. Hopefully, we will achieve a global standard soon. The primary cause of death has been non-

communicable diseases (NCDs) in Bangladesh. NCD caused 74 percent of the deaths in 2014. The overall prevalence of smoking is 26.2 percent. 18% of adults suffer from hypertension and 4% have diabetes mellitus (Rahman et al., 2010; WHO, 2014; Bangladesh Heart Foundation, 2015).

## RESULT:

A systematic review of some Medicinal plants of Bangladesh which have beneficial activity on Cardiovascular Health. 21 medicinal plants have been chosen for the systematic review from reputed journals that have bioactive compounds and pharmacological activity on Cardiovascular Health problems. I have also listed available anticoagulation and hypolipidemic in the market. These data have been collected by visiting pharmacy shops in various regions.

**Table 1: Plants that have cardiovascular effects:**

| Plant Scientific Name        | Local Name                           | Parts Used | Bioactive compounds  | Common Pharmacological Activity  | Specific Cardiac Activity   | Ref.              |
|------------------------------|--------------------------------------|------------|--|--|---|-------------------|
| 1. <i>Cynara scolymus</i>    | Globe artichoke and Garden artichoke | Leaves     | Inulin, minerals, vitamins, fiber, polyphenols   | Hypolipidemic, antiatherogenic, hepatoprotective, antioxidant  | Hypolipidemic activity  | (Morazzoni, 2006) |
| 2. <i>Allium sativum</i>     | Garlic                               | Fruit      | Allicin, Flavonoid, Alkaloid   | Anti-microbial, antioxidant, anti-inflammatory, anticancer, anti-Alzheimer's, anti-obesity, anti-hypertensive, Antiatherosclerosis, Antithrombotic | Anti-hypertensive, Antiatherosclerosis, Antithrombotic            | (Ryu, 2012)       |
| 3. <i>Allium hirtifolium</i> | Wild garlic                          | Fruit      | Dimethyl trisulfide, methylthioether disulfide, thymol, carvacrol, tris methane, chloroorcylaldehyde | Hypotensive, Anti-Atherosclerotic, Antithrombotic or Anti-aggregatory, Hypoglycemic, Anti-inflammatory, Antidiabetic, antibacterial, antifungal    | Atherosclerotic, Antithrombotic or Anti-aggregatory, Hypoglycemic | (Gachkar, 2010)   |
| 4. <i>Crocus sativus L</i>   | Saffron                              | Stigma     | Ceocetin, safranal, picrocrocin  | Anti-Alzheimer's, antitussive, hypolipidemic, anticonvulsant, antinociceptive, anti-inflammatory   | Hypolipidemic activity  | (Moshiri, 2010)   |

|                                      |                              |              |  |  |  |                      |
|--------------------------------------|------------------------------|--------------|--|--|--|----------------------|
| 5. <i>Cinnamomum cassia</i> (L.)     | Chinese cinnamon             | Bark         | Terpenes, phenylpropanoids, lignans, flavonoids, aromatic compounds, aliphatic compounds, coumarins, steroids  | Anticoagulant, anti-obesity, immunoregulation, Anticoagulant, insecticidal and cardiovascular acaricidal, protective cardiovascular protective, cytoprotective | Anti-obesity, Anticoagulant, cardiovascular protective | (Khan,2012)          |
| 6. <i>Artemisia dracuncululus</i> L. | Tarragon                     | Leave        | 1,8-Cineole, camphor, camphene, borneol, thymene, terpinene-4-ol, $\gamma$ -terpinene, $\alpha$ -terpineol, caryophyllene oxide, $\beta$ -pinene, $\alpha$ -pinene, $\beta$ -myrcene, limonene | Anti-inflammatory, antipyretic, antiseptic, antispasmodic, antiparasitic, anticoagulant, antimicrobial, anthelmintic, fungicidal                               | Anticoagulant  | (Mistry,2013)        |
| 7. <i>Cyamopsis tetragonoloba</i>    | Guar                         | Seeds        | Gum residues, saponins, polyphenols  | Anticoagulant, anti-diabetic, hemolytic, Anti-ulcer, Cytoprotective, Anticholinergic, Hypoglycemic, Hypolipidemic, Antimicrobial, anti-inflammatory            | Anticoagulant, Hypolipidemic                           | (Prakash,2015)       |
| 8. <i>Pulmonaria officinalis</i>     | Lungwort                     | aerial parts | Flavonoids, vitamin C, B-complex; minerals-iron, copper, silver, manganese and nickel  | Anticoagulant, astringent, Antioxidant activity, acetylcholinesterase, tyrosinase inhibitory   | Anticoagulant  | (Galić,2011)         |
| 9. <i>Petroselinum crispum</i>       | Cambodia                     | aerial parts | Flavonoids, dihydroxycoumarin, apiin, apigenin   | Anticoagulant, carminative, antioxidant, immunomodulant, anti-spasmodic, diuretic, antirheumatic, antimicrobial, laxative                                      | Anticoagulant  | (Emami,2012)         |
| 10. <i>Tridax procumbens</i>         | Coatbuttons and tridax daisy | Leave        | Bis-bithio-phene, oleonic acids, lupeol, beta-amyrone and taraxasteryl acetate   | Anticoagulant, anti-inflammatory, antiviral, anti-oxidant, antibiotic.   | Anticoagulant  | (Chattopadhyay,2014) |

|                                     |   |                          |   |  |  |                    |
|-------------------------------------|---|--------------------------|---|--|--|--------------------|
| 11. <i>Filipendula ulmaria</i>      | meadowsweet                                     | leaves, roots and flower | flavonoids, vitamins, tannins, polyphenols, ellagitannins, phenolic acids, methyl gallate 3-O-beta-glucoside and rugosins     | Anticoagulant and anti-inflammatory.   | Anticoagulant  | (Esmaeili,2016)    |
| 12. <i>Paeonia anomala</i>          | Anomalous peony                                 | Fruits and roots         | acetophenone, ellagic acid, tert-butylhydroperoxide, methyl gallate, ethyl gallate, fischeroside B and quercetin derivatives. | Anticoagulant, antithrombotic, thrombolytic  | Anticoagulant  | (Zhang,2017)       |
| 13. <i>Ferula communis</i>          | Giant fennel                                    | Leave                    | ferulenol, fert-din, anisate, oxajaeskeanadiol and costic acid  | Anticoagulant, antioxidant and anti-hemolytic  | Anticoagulant  | (El-Sherif,2019)   |
| 14. <i>Panax notoginseng</i>        | Chinese ginseng                                 | Roots                    | ginsenoside Rh4, ginsenoside Rh1, notoginsenoside S, notoginsenoside T ginsenoside Re, notoginsenoside R1 and ginsenoside Rd  | Anticoagulant, anti-hemorrhagic, anti-hypertensive, anti hypercholestrolemic                               | Anticoagulant, anti-hypertensive and hypercholestrolemic | (Zhang,2017)       |
| 15. <i>Thymas vulgaris</i>          | German thyme                                    | Leaves and flower        | thymol, b-caryophyllene, linalool, a-terpinene  | Anticoagulant, antiseptic, antifungal  | Anticoagulant  | (Rahman,2019)      |
| 16. <i>Ginkgo biloba</i>            | Ginkgo  | Leave                    | Phenolic acids, proanthocyanidins, flavonoid, glycosides,   | Anticoagulant, neuroprotective   | Anticoagulant  | (Lee,2017)         |
| 17. <i>Harpagophytum procumbens</i> | grapple plant, wood spider, devil's claw        | Leave                    | Iridoid glycosides, phytosterols, aromatic acids and flavonoids   | Hypolipidemic activity, analgesic, anti-oxidant, anti-diabetic, antiepileptic, antimicrobial, antimalarial | Hypolipidemic activity                                   | (Glombitza, 1999)  |
| 18. <i>Amaranthus spinosus</i>      | spiny amaranth, spiny pigweed, prickly amaranth | Leaves                   | Alkaloids, carbohydrates, cardiac glycosides, flavonoids, phenol, aminoacid, proteins, saponins, tannins                      | Hypolipidemic activity, hepatoprotective, antioxidant, anti-diabetic, anti-inflammatory                    | Hypolipidemic activity                                   | (Akindahunsi,2010) |

|   |                      |       |   |   |                        |                  |
|---|----------------------|-------|---|---|------------------------|------------------|
| 19.<br><i>Glycyrrhiza glabra</i>        | Liquorice            | Roots | alkaloids, glycosides, flavonoids, phenolics, saponins, tannins, terpenes, anthraquinones, essential oils, steroids         | Hypolipidemic, neuroprotective, antioxidant, sedative, anti-depressive, antimicrobial, antiviral, skin effects, anti-inflammatory, anti-ulcer, anti-tumor, hepatoprotective | Hypolipidemic activity | (Mohamed,2014)   |
| 20.<br><i>Withaniasomnifera</i>         | Poisonous gooseberry | Roots | steroidal lactones, alkaloids, flavonoids, tannin   | Hypolipidemic, anti-inflammatory, anticancer, chemoprotective, immunomodulatory, anti-oxidant   | Hypolipidemic activity | (Choudhary,2008) |
| 21.<br><i>Chlorophytum borivilianum</i> | Safed Musli          | Roots | Carbohydrates, gums, mucilage, glycosides, phenolic compounds, proteins, amino acids, saponins, steroids, sterols, terpenes | Hypolipidemic, immunomodulatory, anticancer, anti-inflammatory, antioxidant, hypolipidemic activity   | Hypolipidemic activity | (Tiwari, 2010)   |

**Table 2: Available anti-coagulant drugs in the market**

| Agents        | Class                                       | Route of Administration | Mechanisms   | Major side effects  |
|---------------|---|-------------------------|--|---|
| 1.Ticlopidine | Thienopyridine                              | Oral                    | Irreversibly blocks the P2Y12 component of the ADP receptor on the surface of platelets        | Diarrhea, stomach upset, nausea, vomiting, dizziness              |
| 2.Clopidogrel | Thienopyridine                              | Oral                    | The active metabolite specifically and irreversibly inhibits the P2Y12 subtype of ADP receptor | Stomach ache or abdomen pain, nosebleed, increased bleeding       |
| 3.Prasugrel   | Thienopyridine                              | Oral                    | Reduce the aggregation ("clumping") of platelets by irreversibly binding to P2Y12 receptors    | Bleeding, anemia, atrial fibrillation, back pain, slow heart rate |
| 4.Aspirin     | Nonsteroidal anti-inflammatory drug (NSAID) | Oral                    | Decreased production of prostaglandins and TXA2  | Rash, gastrointestinal ulcerations, abdominal pain,               |

|               |                               |      |   |   |
|---------------|-------------------------------|------|---|---|
|               |                               |      |   | upset stomach   |
| 5.Cangrelor   | ATP analogue                  | IV   | Reversibly inhibit P2Y12 receptors  | Allergic reactions, severe bleeding, anaphylaxis  |
| 6.Ticagrelor  | Cyclopentyltriazolopyrimidine | Oral | Blocks adenosine diphosphate (ADP) receptors of subtype P2Y12   | Bruising, bleeding more easily, nosebleeds, headache, dizziness   |
| 7.Dabigatran  | Direct thrombin inhibitors    | Oral | Binds to the active site on the thrombin molecule, preventing thrombin-mediated activation of coagulation factors | Nausea, abdominal or stomach pain, stomach upset, indigestion, heartburn  |
| 8.Apixaban    | Factor Xa inhibitors          | Oral | Reversibly direct inhibition of free and clot-bound factor Xa   | Headache, dizziness, weakness   |
| 9.Rivaroxaban | Factor Xa inhibitors          | Oral | Inhibits both free Factor Xa and Factor Xa bound in the prothrombinase complex                                    | Back pain, bloody stools, bowel or bladder dysfunction, burning, crawling, itching  |
| 10.Edoxaban   | Factor Xa inhibitors          | Oral | Selective, reversible and competitive inhibitor of human factor Xa  | Bleeding that takes longer to stop, bruising more easily, skin rash, reduced liver function   |
| 11.Warfarin   | Vitamin K antagonist          | Oral | Inhibits the vitamin K-dependent synthesis of biologically active forms of the clotting factors II, VII, IX and X | Severe bleeding, including heavier than normal menstrual bleeding, red or brown urine, black or bloody stool, severe headache or stomach pain |
| 12. Heparin   | Antithrombotic                | Oral | Binds to the enzyme inhibitor antithrombin III  | Bleeding and swelling, discomfort, redness, warmth, irritation, or changes in the skin where the medication was administered                  |

**Table 3: Available lipid-lowering drugs in the market**

| <b>Agents</b>   | <b>Class</b> | <b>Route of Administration</b> | <b>Mechanisms</b>   | <b>Major side effects</b>                                  |
|-----------------|--------------|--------------------------------|---|--|
| 1. Atorvastatin | Statin       | Oral                           | A competitive inhibitor of HMG-CoA reductase  | Joint pain, insomnia, urinary tract infection, nausea.     |
| 2. Fluvastatin  | Statin       | Oral                           | Blocks the liver enzyme HMG-CoA reductase   | stomach upset or pain, indigestion, nausea, muscle pain    |
| 3. Lovastatin   | Statin       | Oral                           | Inhibits 3-hydroxy-3-methyl glutaryl-coenzyme A reductase (HMG-CoA reductase)   | Pain in the stomach area, nausea, heartburn, constipation. |
| 4. Pitavastatin | Statin       | Oral                           | Inhibitor of HMG-CoA reductase, the enzyme that catalyzes the first step of cholesterol synthesis                     | Muscle pain, back pain, joint pain, constipation.          |
| 5. Pravastatin  | Statin       | Oral                           | Inhibits the function of hydroxy methyl glutaryl-CoA (HMG-CoA) reductase  | Muscle pain, nausea, vomiting, diarrhea.                   |
| 6. Rosuvastatin | Statin       | Oral                           | Works by inhibiting HMG-CoA reductase   | Headache, muscle pain.                                     |
| 7. Simvastatin  | Statin       | Oral                           | Acts by inhibiting 3-hydroxy-3-methylglutaryl (HMG) coenzyme A reductase.   | Headache, difficulty sleeping, flushing of the skin.       |
| 8. Bezafibrate  | Fibrates     | Oral                           | An agonist of PPAR $\alpha$   | Stomach upset, gas, nausea, itchy skin, redness.           |
| 9. Clofibrate   | Fibrates     | Oral                           | Increases the activity of extrahepatic lipoprotein lipase (LL), thereby increasing lipoprotein triglyceride lipolysis | Headache, muscle aches and gastrointestinal upset.         |
| 10. Fenofibrate | Fibrates     | Oral                           | Enhanced catabolism of triglyceride-rich particles and reduced secretion of VLDL                                      | Headache, back pain, nausea, indigestion.                  |
| 11. Gemfibrozil | Fibrates     | Oral                           | Inhibits lipolysis and decrease subsequent hepatic fatty acid uptake  | Stomach pain, indigestion, diarrhea.                       |

|            |                          |      |  |  |
|------------|--------------------------|------|--|--|
| 12. Niacin | Pyridine carboxylic acid | Oral | Reduces the synthesis of low-density lipoprotein cholesterol (LDL-C), very low-density lipoprotein cholesterol (VLDL-C), lipoprotein(a), and triglycerides, and increases high-density lipoprotein cholesterol | Severe skin flushing combined with dizziness, rapid heartbeat, Itching, nausea and vomiting. |
|------------|--------------------------|------|--|--|

## Discussion

This systematic analysis of local Bangladesh medicinal plant data shows 21 plants are available in this country which is beneficial for cardiac health and several pharmaceutical agents are available for the treatment of blood coagulation and obesity. Judging from this plant source we can assume that plants that are available in South Asia have more beneficial effects on cardiovascular health. Various drugs are available for anticoagulation and lipid-lowering activity which have various side effects. To avoid these side effects natural sources can be used in this case. Further study is needed to isolate more cardiovascular drugs from plant sources.

## Conclusion:

Plants have played an important role in the world of medicine since ancient times. A significant potential source of therapeutics or curative aids has been described as medicinal herbs or plants. My systemic data shows that 21 medicinal plants that benefit cardiac health are available in our country. On the other hand, available drugs for the treatment of these diseases have so many side effects which can be ignored by using medicinal plants.

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