Embelin as a Potential Drug Molecule: A Review

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Abstract

*Embelia ribes* are traditionally used medicinal remedies in India. Embelin is a hydroxyl benzoquinone with alkyl substitution and is considered as one of the main constituents. Embelin has various biological activities includes anticonvulsant, antidepressant, antidiabetic, antifertility, antitumor etc. This review is an overall overview of the existing pharmacology of *E. ribes*, helping to fuel further research in the various unexplored activities that are attributable to this plant.

Keywords: *Embelia ribes*, Embelin; Hydroxyl benzoquinone; Anticonvulsant; Antidiabetic; Antifertility; Antitumor

Introduction

Berries of *Embelia ribes* are used for various traditional medicinal remedies in India. Embelin is a natural hydroxy benzoquinone with alkyl substitution and is considered as one of the main constituents of plant *Embelia ribes* (family: Myrsinaceae) [1,2]. Embelin is known to possess various biological activities which mainly include anxiolytic, anticonvulsant, antidepressant, antidiabetic, wound healing, anthelmintic, antimicrobial, antitumor, chemopreventive, antioxidant, antifertility etc. It has also found a valuable role in different diseases like Huntington disease, myocardial infarction, acute respiratory distress syndrome and ulcerative colitis.

*Embelia ribes* Burm., a plant of climber in nature, which belongs to the Myrsinaceae family. A large, scandent climber with long slender, flexible, terete branches; bark studded with lenticels. Leaves are simple, alternate, elliptic-lanceolate, gland-dotted, short and obtusely acuminate, entire, shiny above. Flowers are small, white or greenish, in both terminal and axillary panicles. Fruits are globose, wrinkled or warty, dull red to nearly black, a short pedicel often present usually one-seeded and globose (Figure 1).

The main parts of plants used include fruits (berries), roots and leaf, to cure various diseases. Chemically, embelin is C_{17}H_{26}O_{4} and is written as 2, 5-dihydroxy-3-undecylcyclohexa-2, 5-diene-1, 4-dione according to IUPAC nomenclature. It has molecular weight equivalent to 294.38594 g/mol (Figure 2).

![Chemical structure of Embelin](image)

Figure 2: Chemical structure of Embelin.

Biological activities

Embelin as anxiolytic: Among various psychiatric disorders, anxiety is one of the most common disorder [3,4]. It constitutes a considerable public health burden. Recent epidemiological evidence suggests that it is much more widespread than traditional belief, influencing up to half of the general population [5]. Gupta et al. described that embelin expressed significant activity in light and dark model, elevated plus maze and open field test induced anxiety models, thereby proving its anxiolytic potential in Figure 3.
Anticonvulsant activity

The prevalence of epilepsy in developed countries is about 50 per 100,000 while that of developing countries is 100 per 100,000. Currently available antiepileptic drugs are synthetic molecules related with numerous side effects and about 30% of the patients continue to have seizures with this treatment [6,7]. Safety against Hind Limb Tonic Extension (HLTE) in MES (Maximal Electro Shock) predicts the potential of a sample to check the spread of seizure release from the epileptic focus in brain and suppressing generalized tonic-clonic and partial seizures [8]. The dose dependent blockage of HLTE, decrease in the time taken for stupor, a complete eradication of extension and fast onset of stupor has been observed by embelin which indicates its antiepileptic activity. Embelin has also been found to exhibit significant and dose dependent delayed onset of clonic- tonic actions and protection from PTZ induced mortality. However, it may be interfering with GABAergic mechanism to exert its anticonvulsant effect [9].

Effect of embelin against acute respiratory distress syndrome

Pulmonary neutrophilia is a firm finding in human Adult Respiratory Distress Syndrome (ARDS) and in Chronic Obstructive Pulmonary Disease (COPD) which is mainly distinguished by airway hyper-reactivity, poorly reversible airflow obstruction, presence of pulmonary fibrosis, mucus hyper secretion, tachypnea and dyspnea on exertion [10,11]. Lipopolysaccharide (LPS) is a cell wall constituent of gram-negative bacteria which is ubiquitous in the environment and is commonly used to set up the experimental model of ARDS [12,13]. LPS administration through intratracheal to rats leads to an increase in pulmonary vascular permeability leading to neutrophil migration into the air spaces [14]. In spite of advancement in treatment approaches ARDS is still integrated with substantial mortality with an approximates ranging from 26% to 58% [15,16]. Consequently, the need remains for medications that are safer and equally or more effective. The present study revealed that pre-treatment with embelin significantly attenuated LPS-induced acute respiratory distress syndrome in rats, which is possibly linked with inhibition of albumin, protein level, MPO activity and inflammatory cell recruitment in BAL fluid. In addition embelin also prevented TNF, NO over production and improved lung histology decreasing infiltration of inflammatory cells. The protective effect might be related with the inhibition of overproduction of pro-inflammatory mediators. These experimental results suggest that embelin is a potential therapeutic anti-inflammatory compound for ARDS therapy.

Role of embelin in Huntington’s disease (HD)

Huntington’s disease is a progressive neurological disorder which is mainly characterized by clinical triad movement disorder, dementia and psychiatric disturbance due to striatal-specific neuronal degeneration [17,18]. Although the exact cause of neuronal death in HD remains unknown, it has been postulated that the abnormal aggregation of the mutant huntingtin protein may cause toxic effects in neurons leading to a cascade of pathogenic mechanisms associated with transcriptional dysfunction, oxidative stress, mitochondrial alterations, apoptosis, bioenergetic defects and subsequent excitotoxicity [19]. HD patients often exhibit deficits in executive tasks which mainly require planning, cognitive flexibility and problem solving. HD possesses challenges for health and social care professionals due to its complexity and unpredictability. With an incidence of 2-10 per 100,000, HD afflicts 30, 000 people in USA and another 250, 000 persons are genetically at risk [18].

Several animal models exists for HD such as stereotactic injection of kainic, quinolinic and ibotenic acids into specific region of the brain, but systemic administration of 3-nitropropionic acid (3-NP) is the recent and most popularly used [20,21]. 3-NP crosses the blood-brain barrier and at cellular level it is an irreversible inhibitor of the electron transport enzyme succinate dehydrogenase (SDH). It is a mitochondrial complex II enzyme responsible for the oxidation of succinate to fumarate in Kreb’s cycle. Subsequently it blocks the transport of electrons in oxidative phosphorylation causing decreased ATP levels in brain. It affects normal brain electrical activity and oxidative stress has been suggested to play a role in 3-NP toxicity. Neurons are considered to be highly active metabolically and therefore processes that affect the mitochondrial function invariably leads to neuronal death [22,23]. Thus, a major factor involving 3-NP toxicity could be mainly because of cellular and mitochondrial stress [24]. Accumulating data indicates that 3-NP produces free radicals and consequent disturbance of glutathione reduct cycle [25]. The inflammation associated with 3-NP, also acts as a contributing factor for neuronal damage and further free radical generation [25]. In addition to striatum which is the major site of toxicity, 3-NP also causes severe damage to the neurons present in other regions of brain like hippocampus, cortex and hypothalamus [26]. Dhadde et al. demonstrated that embelin normalized the altered behavioral, biochemical and histopathological parameters in 3-NP treated rats, suggesting its neuroprotective efficacy. The possible mode of protection may be due its antioxidant properties. However, further research is warranted to elucidate the specific mechanisms involved.

Anti-depressant action of embelin

According to WHO, 1999, Depression is a common disorder that is projected to become the second biggest contributor to the global health problem and disability by the year 2020. There is now an ever increasing rate of depression-related death both by suicide and due to the associated physical/physiological disorders [27]. The primary symptoms of depression are unhappiness, low mood or reduced interest. Most of the today's antidepressant therapies have limitations due to either their inadequate efficacy over prolonged usage or unwanted side effects.
The major bioactive constituent of *Embelia ribes* is embelin, was found to possess significant activity in mice TST and FST experimental models. The observed potent activity at doses lower than the standard antidepressant drug, imipramine, suggests the potential of embelin and *Embelia ribes* for treating mental depression. Future studies are required to ascertain the exact mechanism(s) of action of embelin’s antidepressant-like effects.

**Anti-diabetic potential**

Diabetes is a major endocrine disorder causing morbidity and mortality worldwide. The problem of diabetes is mostly concentrated to India, as several studies have clearly documented an increased ethnic susceptibility to diabetes in migrant Asian Indians [28,29]. Recent epidemiological studies have pointed to the growing epidemic of diabetes in India [30]. Indeed, according to the recent Diabetes Atlas produced by the International Diabetes Federation (IDF), India is home to the largest number of people with diabetes in the world, 40.9 million diabetic subjects in 2007 and these numbers are predicted to increase to 69.9 million by 2025 [31]. Diabetes mellitus is a heterogeneous group of disorders characterized by chronic hyperglycemia, polyurea, polydipsia, polyphagia, emaciation and weakness due to disturbance in carbohydrate, fat and protein metabolism and directly related to absolute or relative deficiency in insulin secretion and/or insulin action [32,33]. Diabetes is fast turning out to be the epidemic of the 21st century. Type 2 Diabetes Mellitus (T2DM), non-insulin dependent diabetes, is more prevalent (more than 90% of all diabetes cases) and the main driver of the diabetes epidemic, now affects 5.9% of the world’s adult population in developing countries [34].

Diabetes is induced by various models in rats to study the effect of embelin. The models which are used to induce diabetes are alloxan induced diabetes, streptozotocin-induced diabetes, lithium-induced nephrogenic diabetes insipidus. It is a type of diabetes insipidus, manifested by a lack of response of the collecting duct to circulating Anti-Diuretic Hormone (ADH) and cause frequent and excessive urination.

The present study demonstrated that the treatment of diabetic rats with embelin have exerted a considerable hypoglycemic effect. In addition, embelin could ameliorate the impaired renal function, inhibition of liver damage and resemble of islets of pancreas associated with alloxan diabetes. It is an ideal lead molecule for further antidiabetic drug development due to its safety and efficacy. It has also been found that embelin treatment reduced the elevated plasma glucose, lipid profile, ameliorated oxidative stress and inhibited intracellular pro inflammatory mediators in HFD+STZ induced diabetic rats. Moreover, further results indicate that pro-inflammatory mediators and oxidative stress may be major triggering factors in type 2 diabetes mellitus. The therapeutic properties of embelin may be useful as bio-modulator or adjuvant and can be combined with clinically effective anti-diabetic agents. With the use of such adjuvants in clinical medicine, some of the side effects observed on chronic treatment of anti-diabetic drugs can be markedly reduce. Ethanolic extract of *E. ribes* has potential for its evaluation as a protective agent against toxicity induced by streptozotocin.

Embelin exhibited insulin sensitizing effect through adipose tissue specific partial agonism of PPARγ and activated glucose transport through translocation and activation of GLUT4 mediated by insulin dependent PI3k/p-Akt pathway in epididymal adipose tissue. It also protected β-cells by scavenging free radicals and alleviated dyslipidemia in insulin resistant animal model. This drug can be considered useful in the prevention and treatment of obesity related T2DM. However, it seemed to be effective in NDI by its predominant effect on promoting antioxidant status and decrease the urine excretion may be due to the blocking of sodium channels.

**Ulcerative colitis (UC)**

Inflammatory bowel diseases (IBD), Crohn’s disease and Ulcerative Colitis (UC) are chronic, relapsing, remitting gastrointestinal (GI) diseases characterized by chronic inflammation of the intestine [35-38]. UC and Crohn’s disease are associated with intestinal and extra intestinal clinical manifestations of disease, which include weight loss, diarrhea accompanied by blood and/or mucus, fever, gastric dysmotility and shortening of the colon [39,40]. The pathogenesis of IBD remains unclear, but is mainly related to the imbalances between various pro-inflammatory cytokines such as, tumor necrosis factor (TNF-α), interferongamma (IFN-γ), interleukin-1 (IL-1), IL-6 and IL-12 and anti-inflammatory cytokines, such as IL-4, IL-10, IL-11, are believed to play a central role in modulating inflammation [41]. Experimental colitis was induced by giving mice drinking water ad libitum containing 5% (w/v) DSS for 7 days. Results of this study suggested the potent anti-inflammatory effects of embelin in DSS-induced colitis model. It is mediated via the inhibition of pro-inflammatory mediator production. Furthermore, embelin was found to have better therapeutic effects than 5-ASA, which is currently used to treat IBD. However, further studies are required to elucidate the mechanism responsible for therapeutic effects of embelin.

**Wound healing activity**

Healing of wound is a complex cellular event by which injured tissue repaired as steadily as possible to its normal stage. The healing process depends upon the curable abilities of the tissue, the type and degree of injury and natural environment of health of the tissue [42]. It is a physiological feedback to the tissue injury that results in the restoration of injured tissue by living tissue and thus restoration of tissue integrity. Wound repair mechanism occurs by four basic processes such as inflammation, wound contraction, epithelialization and granulation tissue formation. Inflammation outset promptly after the interruption of tissue integrity. The platelets became adherent with clotting factors and form haemostatic plug to prevent bleeding from the vessels. The prostaglandins (PGE1 and PGE2) are discharged in the inflammatory area and suggest to be the final mediators of acute inflammation and may play a haemostatic role for leukocytes and fibroblasts. The active ambulatory leukocytes migrate into the wound and start engulfing cellular debris, at the introductory stages contraction of wound begin slowly and turn into speedy after 3 to 4 days. The myofibroblasts which is present in the margin of wound, emerge to establish the machinery for the wound contraction. These are liable for overlaying debris. The epithelialization of wound is mostly carried out by proliferating and migrating marginal basal cells which are lying close to the wound margin. The hematoma present inside the wound can be replaced by granulation tissue consisting of new capillaries and fibroblasts. The fibroblasts are most likely responsible for production of the mucopolysaccharide ground substance. The lymphatics develop new nerve fibres along with the formation of scar tissue in which collagen turn over increases. The ethanolic extract of *Embelia ribes* tends to augment the formation of collagen fibres. At last, the breaking strength of the wound increases according to the increase of collagen content. Morton and Malone
explained wound closure and time of epithelialization in a way that rats were inflicted with excision wound [43] under light ether anesthesia. The skin of the impressed area was excised to full thickness on the dorsal thoracic region of the rats to obtain a wound area of about 500 mm². The drugs were applied topically on a daily basis till the complete epithelialisation has resulted. The wound closure was calculated at regular intervals to see the percentage of wound closure and epithelialization time. The results indicated the formation of new epithelial tissue which was covering the wound. The period of epithelialisation was given by the number of days required for deteriorating of the scar without any residual of the raw wound. Ehrlich and Hunt described incision wound model [44] consisting of 6 cm long paravertebral incisions which were made through full thickness of the skin on either side of the vertebral column of the rat. The incision was carefully done with at least 1 cm lateral to vertebral column. The wounds were left undressed and drugs were applied topically to the wound once a day, till complete healing had resulted. The skin breaking strength of the 10-day-old wound was measured by continuous constant water technique as explained by Lee and Tong. The skin breaking strength is expressed as the minimum weight (in grams) of water required to bring about the gapping of the wound. Significant wound healing activity was most commonly observed in animals treated with Ethanolic extract and embelin. The histology of granulation tissue for animals treated with embelin revealed complete healing with more of fibroblasts within marked increase of collagen tissue and increased number of blood vessels.

Anthemintic activity

Helminthes infections is one of the most widespread infections in humans, covering a huge population around the world. Although the majority of infections caused due to helminthes are generally restricted to tropical regions and results in enormous health hazard contributing to the prevalence of undernourishment, anemia, eosinophilia and pneumonia among various individuals [45,46]. Parasitic diseases affects population prominently in endemic areas causing brutal morbidity [47]. The foremost problem in treatment of helminthes diseases is that the gastro-intestinal helminthes generally becomes resistant to currently available anthelmintic drugs [48].

Anthelmintics are basically the drugs that tend to eliminate parasitic worms (helminths) from the body, by either inhibiting or killing them. They are also referred as vermifuges or vermicides [49]. Honiberger [50] along with other scientist described Embelia ribes as an essential vermifuge. Watt [51] explained the additional power of powdered seeds (E. ribes) in curdled milk in combination with castor oil which tends to fasten the eradication/expel of tapeworms. According to Ved Prakash and Mehrrota [52], E. ribes is considered as one of the important herbs among the fifty-two Indian traditional herbs responsible for anthemintic activity. Furthermore, herbal monographs (of M/s Himalayan Drugs Company, Bengaluru) suggested that embelin is primarily effective against tapeworm and not roundworm or hookworm. The nematocidal activity of embelin has been explained by Mojumder and Mishra [53-56] against root-knot nematode (Meloidogyne incognita) with 90% mortality after exposure for 48 h (at 100 ppm concentration level). E. ribes seed oil demonstrated effective activity against Pheretima posthuma compared to standard piperazine citrate (10 mg/ml) was reported by Jalalpure et al.

Antimicrobial activity

Anti-microbial is a compound that either kills or inhibits the growth of microbes such as bacteria, fungi, or protozoans. Chitra et al. studied antimicrobial activity of embelin against 12 pathogenic bacteria using disk diffusion method. It was observed that inhibition occurred for all the twelve at a concentration of 100 microgram/disk, but two of the bacterium namely E. coli and K. pneumoniae revealed significant level of inhibition. Embelin inhibition against methicillin-sensitive and methicillin-resistant strains of Staphylococcus aureus with minimal inhibitory concentration (MIC) value of 250, 62 μg/ml respectively has also been reported by Feresin et al. [57]. It was also examined against E. coli which was found to have MIC value of 50 μg/ml. Embelin has also been reported to inhibit various dermatophytes namely Epidermophtyton floccosum, Microsporum canis, Trichophyton mentagrophytes and Trichophyton rubrum with MIC of 50 μg/ml, however 100 μg/ml was observed against Microsporum gypseum. Rani and Khullar [57] explained the moderate antibacterial activity of aqueous and methanol extracts of E. ribes. Tambekar et al. [58] reported that acetone fraction of E. ribes berries possessed mild antibacterial property against Enterobacter aerogenes, Klebsiella pneumoniae, compared to the standard drug Amoxicillin. Rathi et al. [59] examined petroleum ether extract of E. ribes which had lowest MIC value of 250 μg/ml against Candida parapsilosis (MTCC 1744) and 360 μg/ml against Candida laurintis (MTCC 2898) while water extract of E. ribes had higher MIC value for all organisms. Suthar et al. [60] reported embelin with antifungal potential against Aspergillus flavus 871 and Aspergillus fumigates 2550 with MIC50 values of 470, 1015 mg/L respectively. In our study, we observed bactericidal activity of embelin against gram positive organisms and bacteriostatic activity against gram negative organism [61].

Antitumor activity

Antitumor/anticancer activity compounds are drug or agent that inhibit, delay or reverse tumor. Chitra et al. [62] attempted to describe antitumor activity of embelin in methylcholanthrene-induced fibrosarcoma in albino rats along with their enhanced survival time. Nikolovska-Coleska et al. [63] reported embelin as a fairly potent, non-peptidic, cell-permeable, small-molecule inhibitor of XIAP. It represents a promising lead compound for entirely new class of anticancer agents that target the BIR3 domain of XIAP. Dai et al. [64] explained the embelin potential to inhibit chemical carcinogen-induced colon carcinogenesis.

Chemopreventive activity

The uses of a drug or compound which tend to interfere with a disease process are called chemopreventive agents. Sreepriya and Bali [65] revealed that embelin prevents the induction of hepatic hyper plastic nodules, bodyweight loss, increase in the levels of hepatic diagnostic markers and hypoproteinemia against DENA/PB-induced hepatocarcinogenesis in wistar rats.

Antioxidant activity

An antioxidant is a molecule or agent capable of inhibiting the oxidation of molecules. Oxidation reactions generate free radicals which trigger the biochemical chain reactions and finally damaging cells. Antioxidants tend to terminate these chain reactions by neutralizing free radical intermediates and also inhibiting other oxidation reactions. Both plants and animals owns complex systems of
multiple types of antioxidants defense systems endogenously, such as glutathione, vitamin C and vitamin E (classical examples of non-enzymatic antioxidants) and catalase, superoxide dismutase and various peroxidases (enzymatic antioxidants). With regard to antioxidant activity of embelin few reports are available in public domain. Sumino et al. [66] reported that embelin exhibited free radical scavenging activities towards diphenyl-pircyldihydrayl (DPPH) radicals inhibitory concentration (IC50) of 23.3 ± 0.5 μM. Joshi et al. [67] explained the inhibition of lipid peroxidation and restoration of impaired Mn-superoxide dismutase in rat liver mitochondria by embelin. Siddharthan Sureswaran et al. [68] illustrated crude E. ribes displaying free radical scavenging activities when tested using Diphenyl-2-pircyl Hydrazyl (DPPH). Uma et al. [69] described antioxidant activity against streptozotocin induced diabetic rats using aqueous extract of E. ribes administered orally (100 and 200 mg/kg body weight) and whereas Venkateshwar Rao et al. [70-79] substantiates the antioxidant activity of embelin. Dharmendra Singh et al. reported antioxidant activity of embelin against hepatotoxicity induced rats (at concentration of 25 mg/kg body weight). Radhakrishnan et al. [72] observed embelin lipid peroxide inhibition and restoration of impaired Superoxide dismutase in UV B-induced lymphocytes and fibroblasts by embelin.

Conclusion
Conclusively, embelin is an efficient drug molecule with a potential to act against different diseases. It has widespread biological activities such as antioxidant, anticancer, antimicrobial and has minimum side effects. Thus, embelin can be considered as potential drug molecule as well as advaut for various biological implications and thereby serving the human society.

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