INTRODUCTION
Cytotoxicity is one of the most undesirable side effect caused by the antineoplastic chemotherapeutic agents due to it’s diverse action modes. Chemotherapeutic agents induced cardiotoxicity is very common, particularly in the case of alkylating agents [1]. Cyclophosphamide (CYP) is a bifunctional oxazaphosphorine alkylating agent widely used in the treatment of many neoplastic diseases. Breast carcinoma, acute lymphoblastic leukemia, non-Hodgkin’s lymphoma and a variety of bone and soft tissue sarcomas were treated using this nitrogen mustard type alkylating agent [2]. Along with tumor selectivity and a wide spectrum of clinical uses, CYP also causes multiple organ toxicities [3]. An acute type of cardiotoxicity is observed at a high dose of CYP within 10 d of its administration. Fatal complications such as congestive heart failure, arrhythmias, cardiac tamponade and myocardial depression were seen followed by myopericarditis [4].

The precise mechanism of CYP-induced cardiac toxicity has not been established. CYP undergoes extensive metabolism via the cytochrome-P450 enzymatic system to produce active and inactive metabolites, namely phosphoramidate mustard and acrolein respectively [5]. CYP metabolites provoked oxidative stress and direct endothelial capillary damage with resultant extravasation of proteins, erythrocytes, and toxic metabolites. Breakdown of endothelial cells in the presence of toxic metabolites causes direct damage to the myocardium and capillary blood vessels resulting in edema, interstitial hemorrhage and formation of microthrombi. As a result, CYP has developed an acute heart failure and arrhythmias [6].

Traditionally natural products and herbal medicines have been used to avoid side effects caused during treatments of various ailments. Antioxidants such as polyphenols, phenols and flavonoids are produced by the plants in a significant amount. These compounds are potential chemopreventive agents, due to their hydrogen-donating and metal-chelating capacities. Moreover, some studies have shown that plant base natural products could provide protection against cyclophosphamide-induced toxicities [7]. Mentha longifolia is the herb belongs to the family Lamiaceae, which is used for the treatment of bronchitis, flatulence, anorexia, ulcerative colitis and liver complaints due to their anti-inflammatory, carminative, antispasmodic, analgesic, stimulant, emmenagogue and anti-cathedral activities [8].

Flavonoid consumption prevents many cardiovascular diseases [9]. Mentha longifolia contain mainly five flavonoids identified as luteolin-7-D-glycoside, luteolin-7,3-D-glycoside, apigenin, quercetin-3-D-glycoside and kaempferol-3-D-glycoside[10]. Among these molecules, the quercetin-3-D-glycoside had the highest antioxidant activity [9]. Moreover, Mentha longifolia is traditionally claimed for cardioprotective activity since it is used in the treatment of high blood pressure [11]. The other major components present are a cis-piperitone epoxide, pipertenene oxide, carbone, pulegone, menthone, thymol, menthol, β-thione, carvacrol and (E)-caryophyllene [12]. There is paucity for the availability of scientific evidence proving the cardioprotective activity of Mentha longifolia. Hence, the present study was designed to evaluate the cardioprotective activity of ethanol extract of Mentha longifolia.

MATERIALS AND METHODS

Chemicals and reagents
All chemicals and reagents used were of analytical grade and purchased from standard companies. Cyclophosphamide (Cyclus oncosciences, Ahmedabad, India), Sterile water for injection-IP (Ives drugs Pvt. Ltd, India), Ketamine (Neon pharmaceutical Ltd, Mumbai, India), xylazine (Indian immunological, Guntur, India), Sucrose (Merck Specialities Pvt. Ltd, Mumbai, India). Biochemical kits were procured from accucare (Gujarat, India).
Preparation of Mentha longifolia extract

The plant of *Mentha longifolia* (Voucher specimen No. Kacst 00765) was collected from Riyadh, KSA in the month of September 2015. The material was procured and identified by Dr. Shanavakhsh AE, scientist, king abdulaziz city for science and technology, Riyadh, KSA. The leaves were cut into small pieces and shade dried. It was then coarsely powdered and extracted using ethanol (80%) by maceration process. The obtained ethanolic extracts were filtered and evaporated by using a rotary evaporator and freeze dryer, respectively to give the crude dried extract [9]. The yield was found to be 15.38% (W/W). The extract was freshly dissolved in tween 80% before giving each dose to animals.

**Experimental animals**

Healthy adult *Albino* wistar rats of either sex weighing 170-250 g were housed in standard polypropylene cages and maintained under controlled room temperature (25±5 °C) and relative humidity (55±5%) in a well-ventilated animal house under 12:12 h light and dark cycle at the central animal house, shree devi college of pharmacy, Mangaluru. All the rats were provided with commercially available standard pellet diet, water ad lubricum. The animals were maintained under standard conditions in the animal house as per the guidelines of Committee for the Purpose of Control and Supervision on Experiments on Animals (CPCSEA). The Institutional Animal Ethics Committee approved the experimental protocol (SDCP/AEC-01/2014-15).

**Experimental protocol**

The *Albino* wistar rats of either sex were divided into five groups of six animals each. Group I served as normal control (1 ml/kg of 0.5% carboxymethyl cellulose) for 10 d. Group 2 animals were administered with a single dose of CYP (200 mg/kg, i. p) on the first day of the experimental period and served as cyclophosphamide control group. Group 3 animals were treated with ethanolic extract of *Mentha longifolia* (500 mg/kg, p. o) alone for 10 d. Group 4 and group 5 animals received a single dose of CYP (200 mg/kg, i. p) on first day of the experimental period followed by the administration of ethanolic extract of *Mentha longifolia* 250 mg/kg and 500 mg/kg continuously for 10 d by oral route respectively [13].

**Oxidative marker enzymes assay**

Twenty-four hour after the last treatment, blood was collected by retro-orbital puncture and the serum was separated by centrifugation at 5000 rpm for 10 min. The isolated serum was subjected for assay of marker enzyme namely creatine kinase-MB (CK-MB), creatinine kinase-NAC (CK-NAC) and lactate dehydrogenase (LDH). Estimation of marker enzymes was done by using commercial kits with the help of semi auto analyzer (model: Prietest touch, Robonik India PVT LTD.). Then the animals were sacrificed by mild ether anesthesia. The heart tissue was homogenized with succrose solution (0.25 M) for estimations of superoxide dismutase (SOD), catalase (CAT) and lipid peroxidation (LPO) [13, 14].

**Electrocardiographic studies**

Twenty-four hour after the last treatment, the animals were anesthetized with the combination of ketamine (75 mg/kg, i. p) and xylazine (8 mg/kg, i. p). Leads were attached to the dermal layer of both the semilunar front paws and slightly extended hind legs of a rat lying straight on the back and recordings were made with the help of digital physiograph (model no- DI 2- INCO, Ambala city, India). The changes in heart rate, QRS interval, QT interval, PR interval and RR interval were determined [13].

**Histopathological evaluation**

The heart tissue was dissected and fixed in 10% neutral buffered formalin for approximately 48h. The paraffin sections were prepared on a glass slide and stained with hematoxylin and eosin for subsequent microscopic evaluation [1-4].

**Statistical analysis**

Results are expressed as mean±SEM. Statistical significance was assessed using one-way analysis of variance (ANOVA) followed by Tukey-Karmer multiple comparison tests. *P*<0.05 was considered significant.

**RESULTS**

**Serum enzyme biomarkers**

CYP control group demonstrated a significant increase in serum CK-MB, CK-NAC, LDH values compared to normal control. *Mentha longifolia* (ML, 250 mg/kg and 500 mg/kg) treated rats significantly decreased the CK-MB, CK-NAC, LDH values in dose-dependent fashion compared to CYP control group [table 1].

**Effect on SOD and catalase**

SOD and catalase activity were reduced significantly in CYP control compared to normal control. Experimental groups such as *Mentha longifolia* (250 mg/kg and 500 mg/kg) treated groups resulted in significant improvement in SOD and catalase activity compared to the CYP-treated group [table 2].

**Effect on lipid peroxidation**

CYP control group exhibited a significant increase in LPO levels compared to normal control. *Mentha longifolia* treatment in a dose-dependent manner demonstrated a significant reduction in LPO levels compared to CYP control group [table 2].

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**Table 1: Effect of *M. longifolia* on serum level of CK-MB, CK-NAC and LDH in CYP-induced cardiotoxicity in rats**

<table>
<thead>
<tr>
<th>Treatment</th>
<th>CK-MB (IU/L)</th>
<th>CK-NAC (IU/L)</th>
<th>LDH (IU/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal control</td>
<td>39.6±1.28</td>
<td>69.99±0.67</td>
<td>11.38±1.22</td>
</tr>
<tr>
<td>Cyclophosphamide (CYP) control</td>
<td>251.87±2.07***</td>
<td>366.5±0.79***</td>
<td>381.18±2.16***</td>
</tr>
<tr>
<td><em>Mentha longifolia</em> (ML)</td>
<td>34.59±2.68###</td>
<td>54.96±0.21###</td>
<td>108.5±6.66###</td>
</tr>
<tr>
<td>CYP+ML, 250 mg/kg</td>
<td>90.73±0.66###</td>
<td>192.59±1.85###</td>
<td>30.26±3.84###</td>
</tr>
<tr>
<td>CYP+ML, 500 mg/kg</td>
<td>31.28±0.70###</td>
<td>154.10±2.38###</td>
<td>199.16±1.33###</td>
</tr>
</tbody>
</table>

CK-MB-creatinine kinase-MB, CK-NAC-creatinine kinase-NAC, LDH-lactate dehydrogenase, IU/L-international units per litre. All the values are in mean±SEM, n=6. ns: not significant, ***P<0.001 when compared to normal, **P<0.01 when compared to CYP control.

**Table 2: Effect of *M. longifolia* on antioxidants SOD, Catalase and Lipid peroxidation in CYP-induced cardiotoxicity in rats**

<table>
<thead>
<tr>
<th>Treatment</th>
<th>SOD (Unit/mg protein)</th>
<th>Catalase (Unit/mg protein)</th>
<th>Lipid peroxidation (Unit/mg protein)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal control</td>
<td>25.5±0.81</td>
<td>6.04±0.26</td>
<td>2.74±0.11</td>
</tr>
<tr>
<td>Cyclophosphamide (CYP) control</td>
<td>14.7±0.19***</td>
<td>3.41±0.42***</td>
<td>7.79±0.06***</td>
</tr>
<tr>
<td><em>Mentha longifolia</em> (ML)</td>
<td>26.9±1.51***</td>
<td>62.18±0.17***</td>
<td>2.52±0.20***</td>
</tr>
<tr>
<td>CYP+ML, 250 mg/kg</td>
<td>20.5±0.31***</td>
<td>40.16±0.20***</td>
<td>6.79±0.06***</td>
</tr>
<tr>
<td>CYP+ML, 500 mg/kg</td>
<td>22.4±0.30***</td>
<td>49.24±0.23***</td>
<td>4.79±0.09***</td>
</tr>
</tbody>
</table>

SOD-superoxide dismutase, Unit/mg protein-Unit per milligram protein. All the values are in mean±SEM, n=6. ns: not significant, ***P<0.001 when compared to normal, **P<0.01 when compared to CYP control.
Table 3: Effect of M. longifolia on heart rate and electrocardiographic parameters in CYP-induced cardiotoxicity in rats

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Heart rate (Beats/min.)</th>
<th>QRS duration (ms)</th>
<th>QT segment (ms)</th>
<th>RR interval (ms)</th>
<th>PR interval (ms)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal control</td>
<td>361.33±1.76</td>
<td>28.33±0.09</td>
<td>70.88±0.36</td>
<td>28.89±0.07</td>
<td>62.45±0.18</td>
</tr>
<tr>
<td>Cyclophosphamide (CYP) control</td>
<td>272.33±2.02</td>
<td>31.58±0.26</td>
<td>106.73±0.26</td>
<td>39.22±0.15</td>
<td>85.72±0.02</td>
</tr>
<tr>
<td>Mentha longifolia (ML)</td>
<td>363.00±2.08</td>
<td>28.14±0.08</td>
<td>70.84±0.15</td>
<td>28.82±0.12</td>
<td>62.28±0.16</td>
</tr>
<tr>
<td>CYP+ML 250 mg/kg</td>
<td>295.93±2.26</td>
<td>30.14±0.08</td>
<td>96.41±2.22</td>
<td>37.88±0.10</td>
<td>79.56±0.04</td>
</tr>
<tr>
<td>CYP+ML 500 mg/kg</td>
<td>343.69±2.22</td>
<td>29.87±0.05</td>
<td>79.45±0.18</td>
<td>31.85±0.07</td>
<td>63.66±0.12</td>
</tr>
</tbody>
</table>

Beats/min - beats per minute, ms - millisecond. All the values are in mean±SEM, n=6. ns = not significant, ***P<0.001 when compared to normal, +++P<0.001 when compared to CYP control.

Effect on electrocardiographic parameters

CYP control group reported a significant decrease in heart rate and a significant increase in QRS duration, QT segment, PR and RR interval compared to normal control. ML treated group rectified CYP-induced changes in ECG to normal in dose-dependent manner [table 3].

Histopathological observations

CYP control group showed the distorted structure of heart tissue by necrosis of the cells with degeneration of myofibril, increased interstitial space and diffused inflammation compared to normal control group. The treatment groups treated with low dose of ML (250 mg/kg) showed moderate myocardial damage with less interstitial space and myofibrillar degeneration and high dose of ML (500 mg/kg) showed effective inhibition of CYP-induced cardiac damage by reversal of infiltration of inflammatory cells and fragmentation of myofibrils when compared to CYP control group [fig 1].

DISCUSSION

The aim of the present study was to investigate the effect of ML leaf extract against CYP-induced myocardial toxicity.

From the documented results, it can be concluded that ML leaf extract (250 and 500 mg/kg, p.o.) showed beneficial results dose dependently. Mentha longifolia contain different flavonoids, monoterpene ketones, tannins and saponins which are responsible for different pharmacological and biological activity of these plants [15]. Flavonoids are a class of polyphenolic substances with free radical scavenging, inhibition of hydrolytic and oxidative enzymes properties [16]. It also has anti-inflammatory, anti-allergic, antiviral, anti-aging and anti-carcinogenic activity. Antioxidant properties of the flavonoids leads to its broad therapeutic effects. Besides this, flavonoid compounds may exert protection against heart disease through the inhibition of cyclooxygenase and lipoxygenase activities in platelets and macrophages [17].

Cardiotoxicity is caused due to the destructive effect of CYP. CYP mediates production of xanthine oxidase which catalyses the oxidation of hypoxanthine to xanthine and generates superoxide and uric acid. Superoxide is known to generate reactive oxygen species [18]. Intracellular antioxidation systems play an important role in detoxifying the reactive oxygen species (ROS), which is interfered by the toxic metabolites of CYP and the drug itself [19]. Thus, free radical production is increased and anti-oxidant enzymes are decreased in the presence of CYP. Lipid peroxidation is also induced [18].

In our present finding, animals treated with only CYP demonstrated a significant decrease in SOD, Catalase and a significant increase in lipid peroxidation, which indicates the induction of myocardial toxicity. Prophylactic treatment with Mentha longifolia dose-dependently increase SOD and Catalase activity and decrease lipid peroxidation level which justify its protection.

The marker enzymes such as CK-MB, CK-NAC and LDH were leaked from the myocytes to the blood due to CYP-induced myocardial damage. Hence to detect myocardial necrosis, these marker enzymes were estimated [20].

In the present study, experimental animals treated with CYP without any other treatment reflected the remarkable amount of increase in serum marker enzymes levels which confirms the induction of myocardial toxicity. Mentha longifolia dose-dependently restored the marker enzymes levels.

In the present study, CYP showed abnormal changes in an electrocardiographic pattern such as a decrease in heart rate and an increase in RR, PR and QT intervals and prolongation of QRS interval. Abnormal changes in ECG parameters were also found to be extremely significant (p ≤ 0.001) compared to normal control.

The release of a significant amount of acetylcholine is one of the main reason for decreased heart rate, which is also linked with the genesis of myocardial damage [21]. In the present study, QT prolongation in CYP-treated group may be due to increasing in the cellular Na+ content and decrease in K+ content. Change in parasympathetic tone and conduction system deformation can cause AV block, which in turn leads to prolongation of PR interval associated with CYP [22].

Mentha longifolia in both 250 mg/kg and 500 mg/kg doses, in a dose-dependent manner, bring back the ECG parameters towards the normal.

A histological study in CYP-induced cardiotoxicity supported the findings of other parameters analyzed in different treatment groups.
For the normal heart, myocardial fibers were found to be of uniform size, shape, and configurations with no inflammatory cell infiltrates. Treatment with Mentha longifolia doses dependently inhibited CYP induced cardiac damage by decreasing fragmentation of myofibrils and inflammation. Mentha longifolia predominantly in higher dose (500 mg/kg) was able to retrieve the pathological changes associated with CYP in myocardial cell. The phytoconstituents such as flavonoids, tannins, and saponins along with others in ML may be providing the protective effect against CYP-induced cardiotoxicity.

CONCLUSION

It can be concluded from the present study that CYP treatment causes pronounced oxidative stress and tissue damage in the heart. Administration of ML extract protects the CYP-induced cardiotoxicity in dose-dependent manner. Biochemical, ECG and histopathological studies confirm the cardioprotective role of Mentha longifolia. Future studies can be carried out to establish the fact clinically.

ACKNOWLEDGEMENT

Authors would like to thank Dr. Shanavaskhan AE, scientist, king Abdulaziz city for science and technology, Riyadh, KSA for identification of plant Mentha longifolia. The authors are also sincerely thankful to the management of Shree Devi College of Pharmacy, Mangaluru, India for their constant encouragement and support and providing all the necessary facilities for carrying out this study.

CONFLICT OF INTERESTS

Declared none

REFERENCES


How to cite this article