Herbs have been a great source of natural substances used to prevent many cardiovascular diseases (CVD). Blood lipid levels and coagulation parameters are probably the major determinant for producing CVDs. The present study was undertaken to evaluate the hypo-lipidemic and anti-thrombotic activities of aqueous extract of Myrtus communis L. fruit (AEM) in cholesterol-fed rabbits. Hyperlipidemia was induced following administration of cholesterol for 45 days. Animals of treated group were administered AEM daily for 30 and 45 days at dose of 50 mg/kg. Biochemical tests were performed at the end of dosing that is, on 31st day and again on 46th day. The administration of AEM (50 mg/kg/day) revealed reduction in serum triglycerides and low density lipoprotein; there was also an increase in thrombin and fibrinogen time. Results of present study indicate that the extract exhibited hypolipidemic effects and has also an effect on blood coagulation parameters which may be of value in CVDs. However, further studies are necessary to explore the precise mechanism of action of these effects.

**Key words:** Herbs, lipid profile, blood coagulation, rabbits.
deep-green leaves, white flowers and small dark fruits (Asif et al., 2011). In folk medicine, the decoction of leaves and fruits of myrtle are used orally for stomach aches, hypoglycemia, cough, constipation as well as externally for wound healing (Serçe et al., 2010). The volatile oil (Myrtii Oleum) obtained from leaves are used to lower blood glucose (Jung et al., 2006). Similarly, in Italian folk medicine, the fruit of this plant is used in the treatment of many types of infectious disease, including diarrhea and dysentery (Gortzi et al., 2008).

Lipoprotein abnormalities are considered as a highly modifiable risk factor for CVD (Allen et al., 1996). Hyperlipidemia constitutes a foremost etiopathological factor for atherosclerosis (Banerjee and Maulik, 2002) and is often associated with myocardial infarction and cerebrovascular disorders (Ross, 1999). Hence, alteration in plasmalipid levels, coagulation proteins, platelets and fibrinolytic factors may reduce the chance of atherosclerosis (Eitzman et al., 2000). The use of herbal drugs is well-established in many countries. However, despite the increasing scientific interest in this field, there is a lack of summarized data on composition of herbal medicines and therapeutic applications (Aleksic and Knezevic, 2013). Hence, present study has been planned to evaluate the effects of aqueous extract of Myrtus communis (AEM) on lipid profile and blood coagulation in hyperlipidemia induced rabbits.

MATERIALS AND METHODS

Myrtus communis extract

Myrtus communis L. is a medicinal plant endemic to the Mediterranean area and has been used by locals for its food and medicinal properties since ancient times (Atzei, 2003). Myrtus species have been reported to be very rich in volatile oils (Satrani et al., 2006; Shikhiev et al., 1978), phenolic acids and flavonoids (Romani et al., 1999), tannins (Diaz and Abeger, 1986), anthocyanin pigments (Martin et al., 1990) and fatty acids (Cakir, 2004). The fruit of M. communis was grinded and aqueous extract was prepared by decoction. The grinded fruit was boiled in water (10 parts of water and 1 part of herb) for 3 h followed by filtration. The filtrate was further concentrated by boiling to obtain the aqueous extract. The extract so obtained was kept at -20°C until further use.

Animal's treatment protocols

The aqueous extract of Myrtus communis L. (AEM) was initially tested in the dose of 20 mg/kg for 30 days in a group of 5 animals, but no significant response was observed. New rabbits were then uniformly divided into 2 groups of 7 animals each that is, control and treated. Animals of either group received high cholesterol diet (HCD) regularly for 45 days, 0.125 g/kg cholesterol supplied by Merck in 0.5% corn oil. After 45 days, animals of treated group were administered AEM at the dose of 50 mg/kg (Jung et al., 2006) for 30 days during the first phase of study. Animals of control group were given saline equal to the volume of respective doses according to their body weight. During the second phase of study, animals of treated group were further administered AEM for more 15 days making a total period of 45 days and compared with control for the same period. All substances were administered through oral route. Blood samples were collected thrice from the ear vein of animals, first after 45 days of HCD then again after 30 and 45 days dosing of AEM.

Estimation of lipid profile

Blood sample of about 5 ml were collected in gel tube. Serum was immediately separated out by centrifuging blood samples on 14K Humax centrifuge at 3000 rpm for 15 min. Lipid profile were analyzed on Humalyzer 3000 (semi-automatic chemistry analyzer, Model #16700) (Human Germany) using standard kits supplied by Human. Total cholesterol (TC) and low density lipoprotein cholesterol (LDL-C) was estimated by CHOD-PAP method; triglyceride (TG) by GPO-PAP methods (Trinder, 1969), and high density lipoprotein cholesterol (HDL-C) by the method of Friedwald et al. (1972).

Estimation of coagulation parameters

Blood sample of about 3 ml were collected in coagulation tubes containing 3.2% sodium citrate. Plasma was separated by centrifuging blood samples on 14K Humax centrifuge at 3000 rpm for 15 min. Thrombin time (TT), prothrombin time (PT), activated partial thromboplastin time (aPTT) and fibrinogen were measured by Humaclot duo (Human Germany), using standard reagent kits supplied by Human (Chan et al., 2007).

Mortality rate

Mortality rates were observed in animals receiving HCD and AEM during the total period of experiment. The number of animals that died during these intervals was also noted.

Statistical analysis

All values were compared with the control by taking mean and standard error to the mean using t-test, values of P < 0.05 were considered as significant and P < 0.01 as highly significant. All statistical procedures were performed according to the method of Alcaraz and Jimenez (1989).

RESULTS

Mortality rate

No death was observed in any group of animals during...
Table 1. Effects of *Myrtus communis* on lipid profile after 30 days.

<table>
<thead>
<tr>
<th>Animal group</th>
<th>Parameter (mg/dl)</th>
<th>Cholesterol</th>
<th>Triglyceride</th>
<th>HDL-C</th>
<th>LDL-C</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>132.4±22</td>
<td>234.0±17</td>
<td>4.49±0.75</td>
<td>176.7±7.4</td>
<td></td>
</tr>
<tr>
<td><em>Myrtus communis</em></td>
<td>76.0±6.5</td>
<td><strong>149.9±7.5</strong></td>
<td><em>2.68±0.26</em></td>
<td>132.7±14.0</td>
<td></td>
</tr>
</tbody>
</table>

n = 7. Average value±SEM *P < 0.05 significant as compared to control. **P < 0.01highly significant as compared to control

Table 2. Effects of *Myrtus communis* on lipid profile after 45 days.

<table>
<thead>
<tr>
<th>Animal group</th>
<th>Parameter (mg/dl)</th>
<th>Cholesterol</th>
<th>Triglyceride</th>
<th>HDL-C</th>
<th>LDL-C</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>85.2±18</td>
<td>115.4±17</td>
<td>1.74±0.16</td>
<td>141.4±12</td>
<td></td>
</tr>
<tr>
<td><em>Myrtus communis</em></td>
<td>44.4±3.4</td>
<td><em>61.2±9.7</em></td>
<td>1.72±0.08</td>
<td><strong>58.11±1.3</strong></td>
<td></td>
</tr>
</tbody>
</table>

n = 7. Average value ± SEM. *P < 0.05 significant as compared to control. **P < 0.01highly significant as compared to control

the total period of experiment.

**Lipid profile**

Table 1 gives the comparison of serum total cholesterol (TC), triglyceride (TG), high density lipoprotein (HDL-C), low density lipoprotein (LDL-C) levels in animals of control and treated groups after 30 days, while a similar comparison between the same groups of animals after 45 days is presented in Table 2. Animals which received AEM for 30 days showed highly significant and significant decrease in the levels of TG and HDL-C that is, 149.9 ± 7.5 mg/dl and 149.9 ± 7.5 mg/dl in comparison to control values that is, 234.0 ± 17.0 mg/dl and 4.49 ± 0.75 mg/dl, respectively. Conversely there was no significant change in the levels of TC and LDL-C levels at the completion of dosing. Animals which received AEM for 45 days showed highly significant and significant decrease in the level of LDL-C and TG that is, 58.11 ± 1.3 mg/dl and 61.2 ± 9.7 mg/dl in comparison to control values that is, 141.4 ± 12.0 mg/dl and 115.4 ± 17 mg/dl. However the other parameters were not altered significantly at the end of dosing.

**Coagulation parameters**

Table 3 discloses the comparison of thrombin time (TT), partial thrombin time (PT), activated partial thrombo- plasin time (aPTT) and fibrinogen time in animals of control and treated animals, while a similar comparison between the same groups of animals after 45 days is presented in Table 4. Animals given AEM for 30 days showed significant increase in fibrinogen time that is, 57.3 ± 13 s as compared to control that is, 26.01 ± 2.9 s. However there were no significant changes in TT, aPTT and PT at the completion of dosing. Animals given AEM for 45 days showed highly significant increase in TT that is, 24.7 ± 1.9 s as compared to control, that is, 16.56 ± 1.4 s. However the other parameters were not altered significantly.

**DISCUSSION**

AEM have been found to reduce LDL-C and increase fibrinogen time that is, decrease fibrinogen level which seems to be of clinical importance since there are studies which shows that increased fibrinogen has been associated with cardiovascular risks. Fibrinogen increases cardiovascular risk by several mechanisms since it specifically binds to activated platelets via glycoprotein IIb/IIIa, contributing to platelet aggregation, promotes fibrin formation and increase plasma viscosity (James et al., 2000).

Hypercholesterolemia is a strong risk factor for producing atherosclerosis. There is a strong relationship between blood lipids and coagulation parameters since acute changes in plasma lipids appear to have significant effects on factors affecting thrombosis (Eitzman et al., 2000). Coagulation abnormalities are usually present in critically ill patients and may contribute to morbidity and mortality; hence requires rapid examination to establish the underlying cause and to initiate corrective and supportive treatment (Marcel and Steven, 2006). Thus in the present study, highly significant decrease in LDL-C after 45 days and significant increase in the fibrinogen time after 30 days seems to be very important from clinical point of view since there has been a connection between augmented LDL-C and atherosclerosis. High plasma LDL-C concentration is one of the major risk factor for
Table 3. Effects of *Myrtus communis* on blood coagulation after 30 days.

<table>
<thead>
<tr>
<th>Animal group</th>
<th>Parameter (seconds)</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>TT</td>
<td>PT</td>
</tr>
<tr>
<td>Control</td>
<td>17.84±1.7</td>
<td>7.11±0.6</td>
</tr>
<tr>
<td><em>Myrtus communis</em></td>
<td>30.5±4.0</td>
<td>6.2±0.26</td>
</tr>
</tbody>
</table>

n = 7. Average value ± SEM. *P < 0.05 significant as compared to control.

Table 4. Effects of *Myrtus communis* on blood coagulation after 45 days.

<table>
<thead>
<tr>
<th>Animal group</th>
<th>Parameter (seconds)</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>TT</td>
<td>PT</td>
</tr>
<tr>
<td>Control</td>
<td>16.56±1.4</td>
<td>5.87±0.22</td>
</tr>
<tr>
<td><em>Myrtus communis</em></td>
<td><strong>24.7±1.9</strong></td>
<td>5.60±0.13</td>
</tr>
</tbody>
</table>

n = 7. Average value±SEM. **P < 0.01 highly significant as compared to control.

atherosclerosis (Ross, 1999).

There are studies which suggest that reducing LDL-C decreases the risk of CVD (Aghasadeghi et al., 2008). The decrease in LDL-C by AEM, may be due to the presence of myrtle oil (Jung et al., 2006), while natural compounds semi myrtucummulone and myrtucummulone A may also be responsible for potential anti-atherogenic effect of *Myrtus communis* (Rosa et al., 2008). Hypercholesterolemia induces oxidative stress, since it increases the formation of reactive oxygen species from membrane phospholipids during prostaglandin synthesis. Thus antioxidants and hypolipidemic agents suppress the development of hypercholesterolemic atherosclerosis and induce regression of atherosclerosis (Paul and Kailash, 2003). Therefore suppression of atherosclerosis is associated with decrease in oxidative stress and serum lipids (Kabiri et al., 2011). Hence it may be concluded that *M. communis* may have some role in the prevention of atherosclerosis due to its hypolipidemic and anti-inflammatory effects.

Present study also revealed significant increase in TT that indicates deficiency of fibrinogen or inhibition of thrombin (Lane et al., 2005). Hence prolonged TT may be the results of reduced activity of coagulation factors because factors IX and X (Di Cera, 2008), XI and XII are essentially required for thrombin generation (Gailani et al., 2007). Present study also revealed significant increase in fibrinogen time that is, decreased fibrinogen level after 30 days at 50 mg/kg dose. There are studies which show that increases in the fibrinogen levels are a strong risk factor for the development of CVD (Barazzoni et al., 2000; Zhao et al., 2011). Hence, AEM reduces the risk of vascular diseases by reducing the fibrinogen level.

**Conclusion**

The present study was conducted to explore the effects of AEM on lipid profile and blood coagulation. The overall results of the study reveal AEM to be effective as hypolipidemic agent in the dose of 50 mg/kg and also have an effect on blood coagulation parameters which may be of value in CVD.

**ACKNOWLEDGEMENT**

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**Conflict of Interests**

The author(s) have not declared any conflict of interests.

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