

Original Research Article

Cardioprotective effect of melatonin-standardized ethanol extract of *Ulva lactuca* L against acute myocardial infarction in rats induced by isoproterenol

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Abstract

Purpose: To investigate the cardioprotective effect of standardized melatonin ethanol extract of *Ulva lactuca* (smEEUL) against acute myocardial infarction induced by isoproterenol (ISO).

Methods: Male Wistar rats were divided into 6 groups of 6 rats each. Group A was control group given carboxymethylcellulose (CMC Na) while Group B was given ISO 85 mg/kg BW, subcutaneously. Groups C, D and E were given ISO plus smEEUL 250, 500 and 750 mg/kg orally, respectively. Group F was given melatonin 10 mg/kg. Both smEEUL and melatonin were administered for 28 days. ISO was injected twice, i.e., on days 29 and 30. At the end of the study, electrocardiogram (ECG) was recorded, and blood samples collected for myocardial necrosis enzyme assay. Macroscopic features of left ventricle of the heart were also examined using triphenyl tetrazolium chloride (TTC) staining

Results: ISO-treated group demonstrated a significant ($p < 0.05$) elevation in the level of creatine kinase (CK), lactate dehydrogenase (LDH) and aspartate aminotransferase (AST), with increase of 64.01, 65.66 and 99.40 %, respectively, compared with control group. ECG pattern from ISO also showed ST segment elevation. The groups given smEEUL reduced significantly the level of CK, LDH and ST segment elevation ($p < 0.05$). The highest dose showed a reduction ($p < 0.05$ of 62.52 and 77.70 % in the levels of CK and LDH, respectively, compared with ISO treated group. Macroscopic examination using TTC staining supported the above results.

Conclusion: The results show that administration of smEEUL produces cardioprotective effect by restoring ST segment elevation, and reducing both the area of myocardial infarction and myocardial necrosis enzyme level.

Keywords: Cardioprotective, *Ulva lactuca* L, Isoproterenol, Melatonin, Myocardial necrosis and infarction, ST segment

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INTRODUCTION

Cardiovascular diseases remain a problem in Indonesia, one of which is acute myocardial infarction (AMI) [1]. AMI can be triggered by

certain chemical compound resulting in the damage of myocardium cells [2]. In animal model, AMI may also induced by chemical agent. Isoproterenol is a nonselective β -adrenergic agonist for the induction of myocardial infarction in rats [3,4], which can modulate cellular and

functional responses in the heart by its reactive oxygen species [5].

Marine organisms are potential sources of highly bioactive secondary metabolites that are important in the development of new pharmaceutical agents [6]. *Ulva* sp is an algae which are numerous in coastal intertidal habitats worldwide [7]. Some studies have identified the melatonin on algae [8,9]. Melatonin also has been reported as the main compound of *U. lactuca* [9]. Pape and Luning, (2006) showed that *U. lactuca* contain melatonin approximately 12 pg/g (fresh weight). Antioxidant activities and cardioprotective effect of melatonin have been reported in previous studies [10–12]. Administration of melatonin with the dose of 10 mg/kg/day for 7 days in Sprague Dawley male rats has shown cardioprotective effect [13].

Compounds of natural origin can play important roles in prevention and treatment of various diseases. The main compound of *U. lactuca* (melatonin) has been reported to have antioxidant and cardioprotective effect. Therefore, in this study, melatonin standardized in the ethanol extract of *U. lactuca* (smEEUL) was used to investigate the cardioprotective effect against AMI induced by isoproterenol in Wistar rat.

EXPERIMENTAL

Chemicals

Isoproterenol hydrochloride and melatonin were purchased from Sigma Aldrich Co. St Louis, MO., USA. All other biochemical and chemical reagents used were of analytical grade.

Collection and identification of plant materials

Ulva lactuca L were collected at low tide in the afternoon in January 2015 from Drini Beach, Yogyakarta, Indonesia. Plant species was authenticated by Sujadmiko from Systematical Plant Laboratory, Faculty of Biology, Gadjah Mada University. A voucher specimen (no. 0626/S.Tb/I/2015) was deposited in the herbarium of Faculty of Pharmacy, Ahmad Dahlan University Yogyakarta, Indonesia

Plant extraction

Plant extraction was performed as previously described [14]. Briefly, algae were washed with water, dried and were grinded to make powder. The powder was extracted with 96 % ethanol by maceration method. Ethanol extract then was

evaporated with a vacuum rotary evaporator at 4 °C. The extract was filtered and dried at high vacuum and then stored in the refrigerator.

Identification and determination of melatonin in ethanol extract of *U. lactuca* L

Identification of melatonin was performed by thin layer chromatography (TLC) method [15]. TLC elution was carried out on Silica F254 eluted with a mobile phase of n-butanol-acetic acid-water (12: 3: 5) with melatonin as standard and then observed under UV light at 254 nm. Determination of melatonin content was carried out using TLC densitometry and quantitated using the calibration curve of melatonin (0-2.0 mg/ml).

Treatment of animals

Male albino Wistar rats aged 7 - 8 weeks were used in this study. The animals were housed in air conditioned room and were kept in standard laboratory condition, which included a 12-h light-dark cycle and temperature (25 ± 2) ° C. The rats were fed with standard pellet diet AD-2 (Comfeed Industries Ltd) and water *ad libitum*. All procedures related with animal experiments were approved by the Animal Ethical Committee of Integrated Laboratory of Research and Testing (LPPT), Gadjah Mada University (no. 205/KEC-LPPT/XII/2014). Equipment including handling and sacrificing of the animals were in accordance with the Guide for the Care and Use of Laboratory Animals of US National Institutes of Health [16].

Experimental design

The procedure used followed that described previously [17]. The animals were randomly divided into 6 groups of 6 rats. Group A was control group and received carboxymethyl-cellulose (CMC-Na). Group B was given isoproterenol 85 mg/kg BW- (ISO). Groups C, D, and E were given ISO plus smEEUL 250, 500 and 750 mg/kg BW, respectively. Group F was given melatonin 10 mg/kg BW. Both smEEUL and melatonin were orally administrated for 28 days. ISO was injected subcutaneously twice, at days 29 and 30. At the end of the study (day 31), all the animals were anesthetized, electrocardiogram was recorded, and blood samples were collected from the infraorbital sinus for cardiac necrosis enzyme determination. Macroscopic features from left ventricle of the heart were also examined by using TTC staining to determine the area of myocardial infarction.

Electrocardiographic studies

Rats were anesthetized with intraperitoneal injection of pentobarbital sodium (50 mg/kg BW). Needle electrodes were inserted under the skin (lead II position) and then ECG was recorded using animal electrocardiograph Nihon Kohden [2]. Changes in P-wave, QRS Complex and ST segment were determined from electrocardiogram result [17].

Triphenyl tetrazolium chloride (TTC) staining

Myocardial infarction staining was done as mentioned by Lie *et al* [17]. In brief, the heart was transversely cut across the left ventricle, and sections of 2 to 3 mm thick were incubated in 2 % TTC solution prepared in phosphate buffer (pH 7.4) for 60 min at 37 °C, following which they were fixed with 10 % formalin. The slices subsequently were photographed using a digital camera, and the % infarction was analyzed using the computerized Image J (1.40g Wayne Rasband National Institutes of Health, USA).

Biochemical analysis of cardiac necrosis enzyme marker

Animals were anesthetized (intraperitoneal injection of pentobarbital sodium 50 mg/kg BW), and blood was collected via infraorbital sinus. Plasma was separated by centrifugation 5000 xg at 4 °C for 15 min and then kept at -80 °C for enzyme analysis. The markers of cardiac necrosis, such as creatine kinase (CK), lactate dehydrogenase (LDH) aspartate aminotransferase (AST) and alanine aminotransferase (ALT) in the plasma were then determined by commercial diagnostic kits (Biovision, Milpitas, CA 95035 USA and DiaSys, Holzheim, Germany).

Statistical analysis

Electrocardiogram results (duration of basal amplitude of P wave, QRS complex, ST-segment), the area of myocardial infarction and the level of cardiac necrosis enzyme were expressed as mean \pm SEM and were analysed by one-way ANOVA followed by Dunnett's test for multiple comparison using SPSS version 16.0 for Windows. The results were considered significant at $p < 0.05$.

RESULTS

Presence of melatonin in smEEUL

The identification of smEEUL by TLC densitometer method showed the presence of melatonin with retardation factor (Rf) 0.78 co-

chromatographed with melatonin standard. The content of melatonin in smEEUL is 0.52 ± 0.04 % (0.52 mg melatonin/100 mg EEUL).

Effect of smEEUL on electrocardiogram parameters

Figure 1 shows the effect of smEEUL administration on the electrocardiogram parameter. Isoproterenol administration of 85 mg/kg BW for 2 consecutive days led to acute myocardial infarction. ISO group demonstrated a decrease ($p < 0.05$) of the P wave and the QRS complex and elevation in the ST segment. However administration of various dose of smEEUL restored electrocardiogram parameter, such as, an increased in the P wave and the QRS complex and a decreased ST segment compared with ISO ($p < 0.10$).

Effect of smEEUL on the area of myocardial infarction

It can be seen in Figure 2, by using TTC staining that the ISO-induced group showed a large infarct area (white to yellow colour area), while the control group demonstrated a red colour area. ISO-induced group showed a significant ($p < 0.05$) increase of the myocardial infarction area (100 %), compared with the control-group. Administration of smEEUL at doses of 250, 500 and 750 mg/kg BW reduced ($p < 0.05$) the myocardial infarction area state as shown by 48.55 %, 81.82 %, and 57.43 %, respectively compared with ISO treated group. The highest reduction of the area infarction can be found in the group treated with smEEUL 500 mg/kg BW.

Effect of smEEUL on the level of myocardial necrosis enzyme markers

As shown in Figure 3, ISO treated group demonstrated a significant ($p < 0.05$) elevation in the level of creatine kinase (CK), laktat dehydrogenase (LDH) and aspartate aminotransferase (AST); an increase of 64.01 %, 65.66 % and 99.40 % respectively compared with the control group. Treatment with smEEUL 250, 500, 750 mg/kg BW and melatonin 10 mg/kg BW reduce the level of CK by 26.48 %, 41.21 %, 62.52 % and 61.54 %, respectively, compared with ISO group. Compared with ISO treated group, the level of LDH was significantly ($p < 0.05$) reduced by 24.07 %, 64.28 %, 77.70 % and 59.34 % in group treated with smEEUL 250, 500, 750 mg/kg BW and melatonin 10 mg/kg BW, respectively.

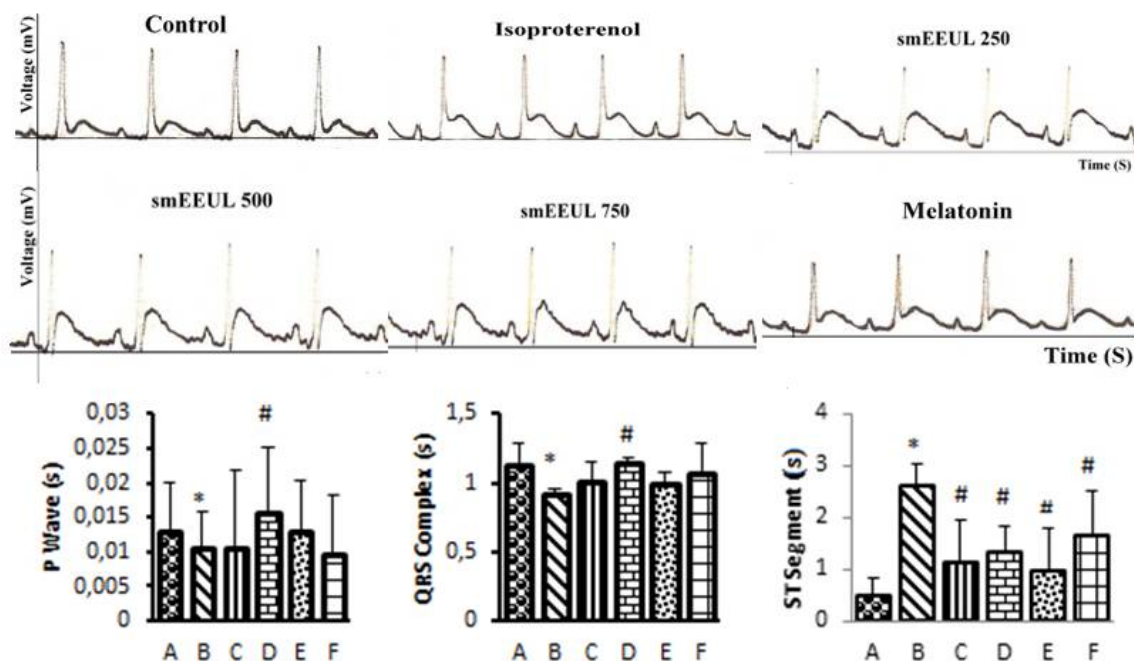


Figure 1: Effect of smEEUL on electrocardiogram parameters. P wave, QRS complex and ST segment are expressed as mean \pm SEM. * $p < 0.05$ compared with control; and # $p < 0.05$ compared with ISO. A = control group; B = ISO group; C = smEEUL 250 group; D = smEEUL 500 group; E = smEEUL 750 group and F = melatonin 10 mg/kg

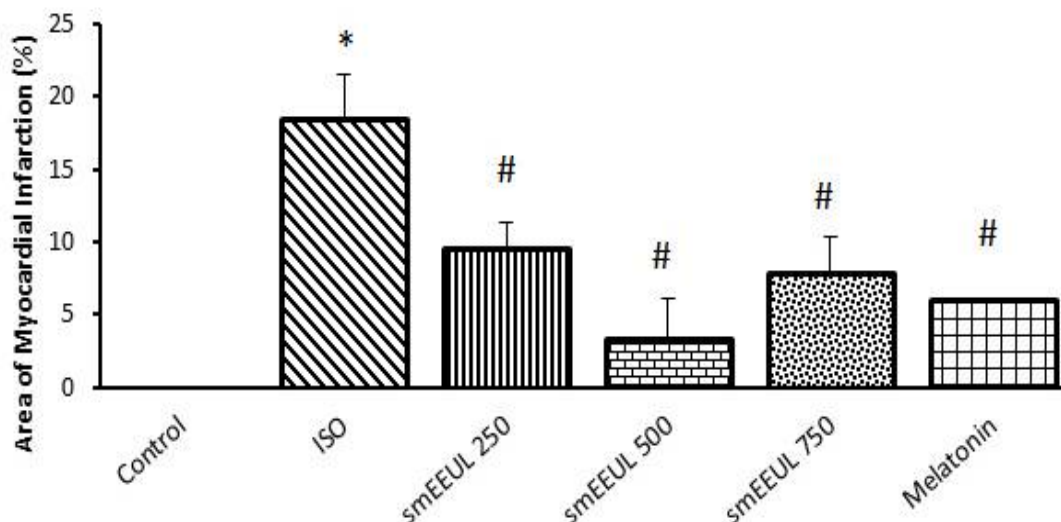
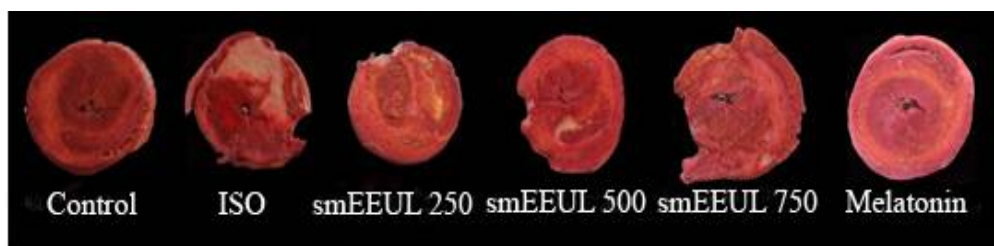


Figure 2: Area of myocardial infarction in the control and experimental animals. Normal myocardium was stained red, while pale white or yellow areas indicate infarct areas. Values are expressed as mean \pm SEM (n = 6); $p < 0.05$ compared with control; # $p < 0.05$ compared with ISO

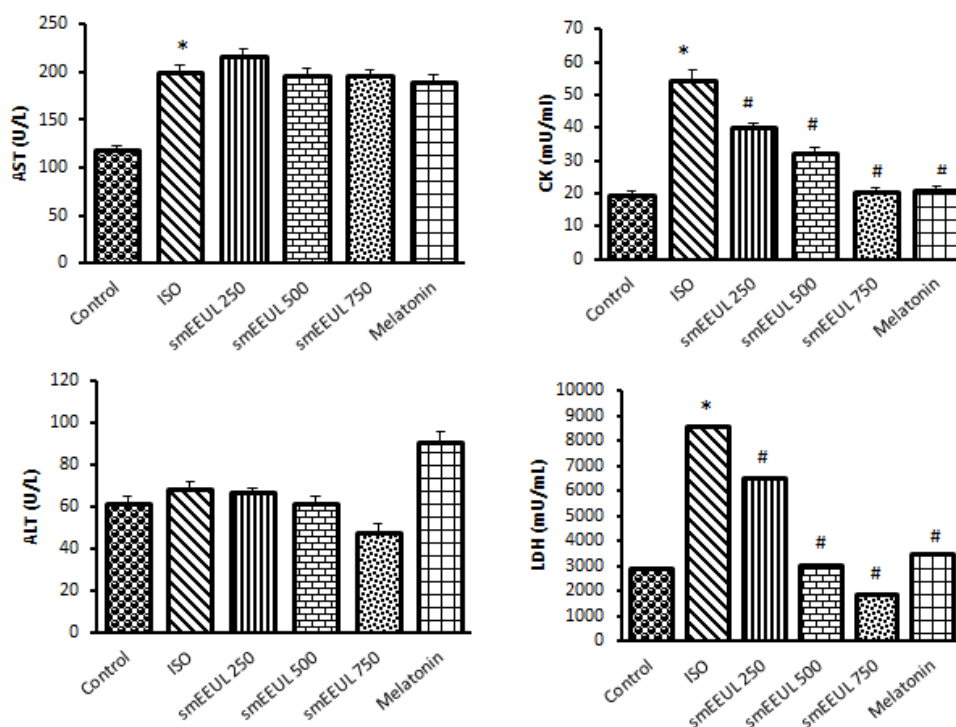


Figure 3: Effect of smEEUL on various cardiac necrosis enzyme marker. (All value are expressed as mean \pm SEM for n = 6. * p < 0.05 compared with control; and # p < 0.05 compared with ISO)

DISCUSSION

Results of this study showed that ISO administration at 85 mg/kg for 2 consecutive days in male Wistar rat led to acute myocardial infarction. The main criteria generally used for the definite diagnosis of MI is evolving pattern of electrocardiograph (ECG)-abnormalities. ISO administration lowered P wave and QRS complex significantly and increased elevation in the ST segment. It has been reported in previous study that administration of ISO 85 mg/kg in two consecutive days in male SD rat increases the segment ST, QT interval and lower P wave, RR interval and QRS complex [17]. Several studies also mentioned that elevation and depression of the ST-segment are typical in the AMI model in rats induced by ISO [17,18]. The change in ECG parameter is due to the changes in potential membrane in the area of infarction [17]. In this study various doses of smEEUL normalized ECG by increasing P wave and QRS complex and decreasing ST segment. It thus seems that the melatonin content of EEUL can modify potential membrane in the area of infarction. However, the mechanism is not clear. Previous study reported that ST-segment elevation resulted in the loss of cell membrane function which reflects the potential difference in the boundary between ischemic and non-ischemic zones [19].

Reactive oxygen species (ROS) are formed in the myocardium due to ISO administration.

Mechanism of ISO-induced AMI involves causing impaired balance between the formation of free radicals and antioxidative defense system, cardiac dysfunction, increased lipid peroxidation and altered enzyme activity which affects the heart muscle [4]. Myocardial infarction increased activities of these marker enzymes (CK, LDH, AST and ALT) in the plasma and this indicates cellular damage and loss of functional permeability of the cell membrane [20]. Administration of smEEUL decreased the level of these enzyme markers of myocardial necrosis (CK and LDH, p < 0.05). This indicates the protective action of smEEUL in preventing myocardial cell and membrane permeability damage by ISO.

Result in this study also demonstrated that smEEUL contains melatonin 0.52 mg in 100 mg EEUL. Therefore, smEEUL 250, 500 and 750 mg/kg BW contains 1.3, 2.6 and 3.9 mg melatonin respectively. The content of melatonin in this study is lower than mentioned in previous study. However, these above doses still have cardioprotective effect as shown from our study. Patel *et al* [] reported that administration of pure melatonin 10 mg/kg BW for 7 days has cardioprotective effects via its ability to restore ST segment elevation, and reduce myocardial necrosis enzyme marker [13].

Previous studies have shown an increase in infarct area due to isoproterenol as indicated with

white to yellow colour by using TTC staining [2,21]. In this study, the level of the enzyme marker of myocardial necrosis (LDH) supported TTC staining results. It has been reported that the area of infarction show loss of membrane integrity that occurs because of the release of lactate dehydrogenase [21,22]. Administration of smEEUL not only significantly decreased the myocardial infarct area by TTC staining but also decreased the level of LDH.

Melatonin is not the only content of *U. lactuca* [23]. Previous research by Abirami *et al* and El-Baky proved that *U. lactuca* have active chemical compound, such as chlorophyll, carotenoids, vitamin C, polyphenols and polysaccharides sulfate which have a role as an antioxidant [24,25]. Thus, in this study, the cardioprotective effect of smEEUL is not only as a result of melatonin, but also possibly by combining effects of melatonin and various compounds mentioned above such as chlorophyll, carotenoids, vitamin C, polyphenols and polysaccharides sulfate.

CONCLUSION

Oral administration of smEEUL for 28 days exhibits cardioprotective effect in rats by restoring ST segment elevation, as well as reducing the area of myocardial infarction and myocardial necrosis enzyme level. smEEUL has a potential to be developed to a cardiopreventive agent for the treatment of myocardial infarction in human.

DECLARATIONS

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Conflict of Interest

No conflict of interest associated with this work.

Contribution of Authors

The authors declare that this work was done by the authors named in this article and all liabilities pertaining to claims relating to the content of this article will be borne by them.

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