Ficus deltoidea aqueous leaves extract abrogates enhanced-oxidative damage in ovariectomized rat models

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ABSTRACT
The aim of the study is to investigate the potential of Ficus deltoidea aqueous extract in protecting the vital organs of female rats against ovariectomy-induced oxidative stress. The rats were grouped into control intact (normal saline), ovariectomized control (OVX), and ovariectomized rats fed orally with 500 mg/kg F. deltoidea extract (OVX+T). Upon treatment completion, the collected organs were homogenized and measured for malondialdehyde (MDA), superoxide dismutase (SOD), glutathione (GSH), catalase (CAT), and glutathione peroxidase (GPx). Serum total cholesterol (TC) was also measured. MDA levels were found to increase in the heart and brain of OVX rats. Ovariectomy caused a decrease in the GSH levels and reduced CAT, SOD, and GPx enzyme activities in the organs of the OVX rats. Ovariectomy also caused an increase in serum TC level. The F. deltoidea leaves extract supplementation not only reduced the MDA levels and increased GSH levels and CAT activities in the selected organs of the OVX+T rats but also decreased the TC level in serum. Glutathione (GPx) activity was found to be increased only in the heart and kidney of the OVX+T rats. In conclusion, F. deltoidea aqueous extract has the potential to be developed as an antioxidant supplement to protect the vital organs against oxidative damages during the postmenopausal condition.

INTRODUCTION
Enhanced oxidative stress and decreased antioxidant defense in postmenopausal women as compared to premenopausal women play an important role in the pathogenesis of various diseases related to menopause (Aroka et al., 2009). For example, the incidence of cardiovascular morbidity and mortality in women after menopause was more prominent compared to premenopausal women (Skafar et al., 1997). These might be due to the loss of protective effects provided by estrogen as it is widely documented that this hormone has protective actions in stimulating the oxidative defense system and inhibits oxidative stress (Kerstin et al., 2003). In addition, estrogen exerts its protective effects on several organs such as the brain, liver, kidney, and heart. Previous studies have shown that estrogen was neuroprotective by preventing intracellular peroxide accumulation in neurons (Behl et al., 1997), hepatoprotective against inflammation and oxidative stress (Shimizu et al., 2007), nephroprotective by attenuating glomerulosclerosis and tubulointerstitial fibrosis (Gluhovschi et al., 2012), and protective against atherosclerosis by modulating low-density lipoprotein (LDL) oxidation, binding of free radicals, and lowering plasma cholesterol (Barp et al., 2009). However, these protective abilities are declined as the women undergo the menopausal process.

Oxidative stress is the term used to explain the oxidative damages in cells, tissues, or organs due to reactive oxygen species (ROS). Oxidative stress occurs when there is an imbalance between the oxidant and the antioxidant system. Superoxide anion (O2−•), hydrogen peroxide (H2O2), and hydroxyl radical are ROS species that target proteins, nucleic acids, or lipid structures and eventually cause tissue damage.

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The addition of oxygen radical to the polyunsaturated fatty acid (PUFA) of lipids causes oxidative damage that resulted in the production of malondialdehyde (MDA). MDA is the end product of lipid peroxidation (LPO) and thus is used as a marker of oxidative stress. To protect the body against the scavenging activity of the ROS, the body has a built-in defense antioxidant mechanism that consists of superoxide dismutase (SOD), catalase (CAT), and glutathione peroxidase (GPx). SOD neutralizes the \( \text{O}_2^- \) radicals to \( \text{H}_2\text{O}_2 \) while GPx reduces \( \text{H}_2\text{O}_2 \) with reduced glutathione (GSH) to water and oxygen. Meanwhile, CAT will disassociate \( \text{H}_2\text{O}_2 \) to water and oxygen (Salvi et al., 2007).

Postmenopausal women are more vulnerable to estrogen-deficiency-induced oxidative stress than their premenopausal counterparts. To date, researchers have been searching for an alternative treatment to reduce the impact of oxidative stress in menopausal women. Foods rich in antioxidants have been shown to be of great benefit in women experiencing menopausal symptoms because they help to eliminate oxidative stress within the body (Doshi and Agarwal, 2013). The natural product has been favorably used for health and medicinal purposes long before drugs ever being invented. A wide variety of natural products are being used to treat ailments, known as traditional medicine. Current estimation states that 80% of the overall population still depends on traditional medicines to fulfill their health care needs as an alternative to regular medication (WHO, 2008). Previous studies on the effectiveness of herbal preparations to manage the symptoms of menopause have been carried out on Labisia pumila, Cimicifuga racemosa, and Trifolium pratense (Berul et al., 2002; Stacie et al., 2009). These herbs contain phytoestrogens, which have a similar chemical structure to human estrogen, thus able to bind to estrogen receptors in human and elicit downstream signaling (Glazier and Marjorie, 2001).

Interestingly, in Malaysia, a traditional plant known as Ficus deltoidea or locally known as Mas Cotek, is popular among traditional medical practitioners. Ficus deltoidea that grows in the Peninsular of Malaysia has different varieties (Fatihah et al., 2012). All parts of the plant including the roots, bark, leaves, and fruits are reported to have medicinal benefits with therapeutic potentials (Abdulla et al., 2010; Adam et al., 2007; Azizan et al., 2017; Bunawan et al., 2014; Kamran et al., 2020; Misbah et al., 2013). The aqueous preparation of F. deltoidea leaves is traditionally used to assist in parturition and after birth, as it has the ability to stimulate uterine contraction (Ibrahim et al., 2018; Sulaiman et al., 2008; Zainudin et al., 2014). The leaves also can be used in tea beverages with a good source of minerals (Nihayah et al., 2012). Besides, it has been reported to possess antihypertensive effects (Aminudin et al., 2007) as well as antioxidant and anti-inflammatory properties (Abdullah et al., 2009). Ficus deltoidea is also able to reduce both blood glucose level (Adam et al., 2007) and cancer risk (Aris et al., 2009). Findings by Hakiman and Maziah (2009) found that F. deltoidea contains phytochemical properties that are related to flavonoid and phenolic compounds. These two antioxidants are capable of delaying or inhibiting the oxidation of lipids or other biomolecules by inhibiting the initiation or propagation of oxidizing chain reactions (Sirissha et al., 2010). However, the benefit of this traditional herb in alleviating oxidative stress in the brain, liver, heart, and kidney due to estrogen deficiency has not been proven to the best of our knowledge. This study aims to investigate and scientifically prove that the reported antioxidant property of F. deltoidea aqueous extract has the potential to protect against ovariectomy-induced oxidative stress in the brain, liver, heart, and kidney of rats.

**METHODOLOGY**

**Animals and study design**

Twenty-four virgin female Sprague-Dawley rats, aged three months old and weighing between 200 and 250 g, were divided into three groups, with \( n = 8 \) rats/group. The animals were housed paired and kept at 24°C with a 12 : 12-hours light-dark cycle. Only virgin female rats were used in this study to minimize hormonal influence variations between animals. The rats were fed with food and water made available ad libitum. The groups were (1) sham-operated (control); (2) ovariectomized rats with orally fed vehicle (distilled water) ovariectomized control (OVX), and (3) ovariectomized rats and orally fed with 500 mg/kg F. deltoidea (OVX+T) via oral route for 28 days. The rats were anesthetized and the ovaries were bilaterally removed via dorsal approach. Ovariectomized female rats were used to ensure the relative homogeneity of the animals’ endogenous hormonal background and to eliminate potential interference from ovarian factors (Schirman-Hildesheim et al., 2006). Here, in this study, ovariectomized rats were used as a menopausal animal model as it creates intrinsic changes in the body biochemistry and physiological function in these animals similar to those of menopausal women (Bosse and Di Paolo, 1995). The dorsal part of the lumbar region was cleared off and disinfected with 70% ethanol. Only one incision of 1.5 cm was made on the cleared skin and another 1 cm incision on the muscle wall on each side of the backbone. The ovaries were located and removed after the uterine horns were tied with sterile sutures. The incision on the skin was closed with skin staples. After surgery, the animals were housed individually for one day. Then, the animals were pair-housed and allowed to recover for 6–7 days before further manipulations. All postoperative animals were monitored closely for any indicator of pain or distress according to the Animal Ethical Committee (AEC) guidelines. Animals that were affected were excluded from the experiment and euthanized appropriately. Rats were anesthetized using a cocktail of Ketamine-Zoletil-Xylazine (KTX) with a dose of 0.4 ml/kg intraperitoneally. For sham-operated rats, the ovaries were left intact. All rats were given a two-week recovery period before the F. deltoidea treatment protocol started. This study was granted approval from the Universiti Kebangsaan Malaysia Animal Ethics Committee (UKMAEC): FSK/BIOMED/2012/FARAH/12-Dec/483-DEC-2012-JUNE-2013.

**Samples collection**

At the end of the 28-day treatment regime, the animals were anesthetized and blood samples were taken via cardiac puncture and collected in the plain tubes. The blood samples were left for 1–3 hours at room temperature and were centrifuged at 3,000 rpm at 4°C for 10 minutes. Serum was collected and stored at −20°C until further investigation. Brain, heart, liver, and kidney were harvested and washed with ice-cold 1.15% (w/v) potassium chloride (KCl). Each of the tissue samples was homogenized by adding 1.15% KCl with a ratio of 3 ml/g (v/w). The tissue
homogenates were centrifuged at 8,000 rpm at 4°C for 20 minutes and stored at −20°C until further use.

**Plant extract preparation**

The leaves were collected from Juaseh Tengah, Negeri Sembilan, Malaysia, and were brought to Herbarium UKMB for identification (voucher specimen UKMB29781). The leaves were extracted as described by Zainudin *et al.* (2014). Briefly, 300 g of the dried leaves were cut into smaller pieces and extracted with distilled water using Soxhlet apparatus. The resultant product was filtered and later freeze-dried before being diluted with distilled water to the desired concentration

**MDA level and total antioxidant status**

The LPO level via MDA quantification was adapted from the Ledwozyw *et al.* (1986) method and measured at 532–535 nm, whereas the total antioxidant status was measured via SOD, CAT, and GPx activities. SOD was detected via Beyer and Fridovich (1967) method at 560 nm, while CAT activity was measured by the Aebi (1984) method at 240 nm. The activity of GPx activity was quantitated based on Paglia and Valentine (1972) method at 340 nm. These tests were carried out on tissue homogenates by measuring the designated optical density using a spectrophotometer (Secomam, France). The MDA level was measured in nmol/mg, the SOD level in UE/minute/µg, the CAT level in UE/mg, and GPx in UE/µg.

**GSH level**

GSH level was measured as described by Ellman (1959). The GSH in the sample reacted with the Ellman reagent to produce a red-colored product that can be measured at 415 nm by a spectrophotometer. The GSH level was measured in nmol/µg.

**Total serum cholesterol level**

The serum cholesterol level was determined using cholesterol high-density lipoprotein (HDL) Direct (Biosystems, Spain). Ten µl serum was added to a 1.0 ml cholesterol reagent. Next, the mixture was incubated at 37°C for 5 minutes and the optical density was measured at 500 nm using Bioanalyzer (Biosystem, Spain). The cholesterol level was measured in mmol/L.

**Statistical analysis**

Data analysis was performed using IBM Statistical Package for the Social Sciences Statistics for Windows, Version 19.0 (Armonk, New York). Statistical significance was determined via one-way analysis of variance followed by Tukey’s honestly significant difference (HSD) for normally distributed data with p < 0.05. The results were presented as means ± standard error of the mean (SEM).

**RESULTS AND DISCUSSION**

Estrogen deficiency in postmenopausal women often leads to oxidative stress (Doshi and Agarwal, 2013). A low level of estrogen has prooxidant effects as it can lead to oxidation of bases, the formation of DNA adduct, and breakage of genetic material (Doshi and Agarwal, 2013). These, in turn, resulted in increased ROS-induced tissue damages that are the basis for a number of diseases and biochemical disorders (Murugananthan and Sathy, 2012). The antioxidant is widely used as a therapeutic approach to overcome oxidative stress, as it can remove free radicals and protects the body from detrimental oxidative damages. As plants are one of the best sources of antioxidants (Sen *et al.*, 2010), thus, the current study was conducted to investigate the antioxidant properties of *F. deltoidea* aequous leaves extract on ovariectomized rats as an estrogen-deficiency model to recapitulate postmenopausal women.

Organs in the body are constantly exposed to the threat of oxidative stress. In women, ovarian estrogen deficiency during menopause accelerates aging by decreasing the body’s defense against oxidative stress. Oxidative stress activates multiple intracellular signalings that often lead to organ dysfunctions and damages (Ogura and Shimosawa, 2014). The brain is more susceptible to oxidative damages since it has a high content of PUFA (Peet, 2002). Since estrogen has dual functions as an anti-inflammatory as well as antioxidant, it has been known to exert a protective role in organs. Estrogen deficiency affects the brain in such a way that it has been reported to be closely related to the pathogenesis of various neurodegenerative diseases such as Parkinson’s and Alzheimer’s disease (Norbury *et al.*, 2003). Besides the brain, liver is one of the sources of oxidative stress as liver damage increases ROS production (Jaeschke *et al.*, 2002). The liver of ovariectomized mice showed high LPO and decreased antioxidant enzymes (Oliveira *et al.*, 2018). The heart, on the other hand, showed deterioration in function following the loss of ovarian estrogen which is reversible through estrogen supplementation (Firth *et al.*, 2020). Furthermore, the presence of estrogen was reported to diminish angiotensin II-induced free radical production in vascular smooth muscle cells and upregulated manganese SOD and extracellular SOD expressions and activities in the cardiovascular system (Kerstine *et al.*, 2003). Another organ susceptible to oxidative stress is the kidney. Estrogen exposure inhibits the mitogenic effect of free radicals on human glomerular mesangial cells (Dubey and Jackson, 2001), thus implicating estrogen’s impact on the kidney in its absence.

**LPO status**

In this study, bilateral removal of the ovaries in the female rats induced oxidative stress in the vital organs. This was evidenced by the significant increase in the MDA level in the heart (1.048 ± 0.003 nmol/mg) and liver (3.08 ± 0.365 nmol/mg) of OVX rats compared to the control groups (1.022 ± 0.005 nmol/mg and 1.561 ± 0.211 nmol/mg, respectively) (Fig. 1). MDA is an indicator of LPO of PUFA of the cell membrane. Estrogen deficiency can cause the excessive formation of ROS. This often leads to LPO as less estrogen is able to bind to the estrogen receptors in order to upregulate the expression of the antioxidant enzyme via the intracellular signaling pathways (Borras *et al.*, 2010; Sridhar *et al.*, 2005). The levels of MDA in the OVX+T groups were significantly lower in the heart (1.026 ± 0.006 nmol/mg), liver (2.053 ± 0.235 nmol/mg), kidney (1.042 ± 0.004 nmol/mg), and brain (4.513 ± 0.005 nmol/mg) compared to the OVX groups (1.048 ± 0.003 nmol/mg, 3.08 ± 0.365 nmol/mg, 1.056 ± 0.004 nmol/mg, and 4.513 ± 0.906 nmol/mg, respectively) (p < 0.05). These observations suggested that *F. deltoidea* leaves aequous extract has the potential as an exogenous antioxidant source to complement endogenous antioxidant capacity. Furthermore, the aequous extract of *Ficus deltoidea* leaves also has been known to contain flavonoids (Omar suggestion).
et al., 2011; Zainudin et al., 2014), phytochemicals that also have been shown to possess estrogenic activities, phytochemicals that also have been shown to possess estrogenic activities.

**Antioxidant status**

All antioxidant enzymes were found to be reduced in ovariectomized rats as demonstrated in the previous study by Ha (2004). SOD is an enzyme used to convert superoxide ion to H$_2$O$_2$, which is then converted by GPx and CAT to oxygen and water (Nazrun et al., 2008). In the current study, ovariectomy had caused the level of SOD enzyme activity (Fig. 2) to be significantly lower in the heart (0.167 ± 0.025 U/mg), kidney (0.083 ± 0.016 U/mg), and brain (1.457 ± 0.329 U/mg) of OVX groups compared to the control groups (brain: 5.901 ± 1.664 U/mg, heart: 0.255 ± 0.020 U/mg, and kidney: 1.457 ± 0.081 U/mg) (p < 0.05). Likewise, the CAT enzyme activity (Fig. 3) in the OVX groups was also significantly lower in the heart (0.666 ± 0.071 U/mg), liver (0.564 ± 0.054 U/mg), kidney (0.85 ± 0.169 U/mg), and brain (0.948 ± 0.089 U/mg) when compared to the control groups (1.484 ± 0.26 U/mg, 1.812 ± 0.237 U/mg, 1.976 ± 0.169 U/mg, and 3.548 ± 0.232 U/mg, respectively) (p < 0.05).

Supplementation with *F. deltoidea* for 28 days in the OVX+T group did not show any significant elevation in the SOD activity in the vital organs although the increasing trends were observed (Fig. 2). There were, however, a significant increase in CAT level in the heart (1.435 ± 0.214 U/mg), kidney (1.75 ± 0.220 U/mg), liver (2.294 ± 0.338 U/mg), and brain (1.75 ± 0.220 U/mg) compared to OVX groups (p < 0.05) (Fig. 3). In all, CAT and SOD levels were observed to decrease following ovariectomy but supplementation with *F. deltoidea* prevented the reduction by increasing the endogenous enzyme activities instead.

The glutathione peroxidase (GPx) enzyme activity in the OVX groups was significantly lower in both the heart (2.428 ± 0.072 U/mg) and kidney (1.456 ± 0.081 U/mg) organs compared to the control groups (heart: 2.711 ± 0.038 U/mg and kidney: 2.600 ± 0.060 U/mg) (p < 0.05). On the other hand, ovariectomy had caused the rats from the OVX groups to exhibit lower GSH levels in the heart (8.780 ± 0.815 nmol/mg protein), kidney (9.710 ± 0.792 nmol/mg protein), and brain (8.480 ± 0.671 nmol/mg protein) compared to the rats from the control groups (heart: 12.872 ± 0.651 nmol/mg, kidney: 13.435 ± 0.972 nmol/mg, and brain: 11.942 nmol/mg ± 0.350) (p < 0.05) (Fig. 5). GSH is the cofactor of GPx used for the completion of the antioxidant system.
(Li et al., 2000). According to Lu (2009), this intracellular peptide has diverse functions that include detoxifications, modulation of cell proliferation, and antioxidant defense. A large reserve of GSH is present in hepatocytes and erythrocytes for the detoxification of free radicals (Ha, 2004). Thus, the dysregulation of GSH synthesis may contribute to oxidative stress and later development for the pathological condition.

Interestingly, supplementation with *F. deltoidea* significantly increased the GPx level only in the heart (2.651 ± 0.027 U/mg) and kidney (2.012 ± U/mg) of OVX+T groups when compared to the negative OVX control group, but not in the other two organs (heart and liver). The data for the level of GPx in the liver, however, cannot be determined due to technical issues (Fig 4). Similarly, GSH levels in the heart (11.635 ± 0.704 nmol/mg protein), liver (17.646 ± 0.0.459 nmol/mg), kidney (13. ± 0.182 nmol/mg protein), and brain (10.430 ± 0.219 nmol/mg protein) all showed significant elevation upon *F. deltoidea* aqueous extract supplementation (p < 0.05). Interestingly, supplementation with *F. deltoidea* increases GSH level in all organs of ovariectomized rats suggesting that *F. deltoidea* has the potential in regulating GSH level (Fig 5).

Previously, *F. deltoidea* is documented to contain flavonoids, tannins, proanthocyanins, and phenols, which are known to be responsible for strong antioxidant properties (Rosmani et al., 2012). In addition, Omar et al. (2011) has identified flavan-3-ol monomer and proanthocyanidins as the main compounds for this property. Meanwhile, another study demonstrated that *F. deltoidea* was a potent antioxidant as shown via its scavenging free radical activities of more than 50% in 2,2-diphenyl-1-picrylhydrazyl assay (Ruzaina et al., 2009). In contrast, the findings from this study suggested that *F. deltoidea* exerts its antioxidant potential via increasing the level of the antioxidant enzymes in the organs of ovariectomized rats, thus highlighting the potential of *F. deltoidea* in combating ROS.

**Serum total cholesterol (TC) level**

The TC level in OVX (2.636 ± 0.036 mmol/l) was significantly (p < 0.05) higher than the control group (2.184 ± 0.120 mmol/l). The concentration of cholesterol was significantly (p < 0.05) decreased in ovariectomized rats after treatment with *F. deltoidea* (2.316 ± 0.058 mmol/l) in comparison to OVX (Fig. 6). Both estrogen and progesterone influence cardiovascular function (Thomas and Pang, 2013; White et al., 2010). Women in the menopause stage lost protective functions against oxidative stress due to the diminishing ovarian hormones and oxidative stress is one of the risk factors for cardiovascular diseases (Brunelli et al., 2017). Estrogen level can affect cholesterol as (LDL-c) level rises and HDL-c level declines after menopause (Guetta and Cannon, 1996). Estrogen induces hepatic LDL-c receptors for LDL-c catabolism; thus, low estrogen levels may increase LDL-c (Bittner, 2001).

Menopausal women with metabolic syndrome who were given ovarian hormonal therapy have shown reduced oxidative stress and improvements in serum triglyceride and high-density lipoprotein cholesterol (HDL-c) levels (Sánchez-Rodríguez et al., 2016).

Supplementation with *F. deltoidea* was found to decrease the serum TC level and this clearly indicates its potential in regulating cholesterol level. This finding is further supported by Taufik et al. (2005) that reported that *F. deltoidea* extracts are able to reduce blood cholesterol levels in rats fed with high cholesterol food and also in humans (Kalman et al., 2013). The observed serum cholesterol-lowering effects by *F. deltoidea* is shown to be mediated by pathways involved in lipid metabolism and transport, inflammation and immune response, antioxidant proteins, acute phase proteins, cholesterol homeostasis, fatty acid catabolism, and cholesterol binding (Chuo et al., 2020). Besides flavonoids, our previous qualitative study on the phytochemical screening of *F. deltoidea* aqueous extract has shown the presence of saponin and tannins (Zainudin et al., 2014), both of which, in combination, are reported to possess antioxidant as well as anticholesterol properties (Hagerman et al., 1998; Marelli et al., 2016). It is not known, however, whether the anticholesterol effects seen in the current study are due to the sequential effects of reduced oxidative stress or vice versa. Although in the current study the animals were first ovariectomized to mimic a menopausal state, therefore, the TC lowering effects observed is most probably due to the dual biological effects of *F. deltoidea* aqueous extract; antioxidant and anticholesterol.

![Figure 5](image5.png) The level of GSH in the heart, liver, kidney, and brain of experimental rats. Data are presented as means ± SEM. (a) Significant difference compared with control group (p < 0.05) and (b) significant difference compared with OVX group (p < 0.05).

![Figure 6](image6.png) The serum cholesterol level in the ovariectomized rats. Data are presented as means ± SEM. (a) Significant difference compared with control group (p < 0.05, and b) significant difference compared with OVX group (p < 0.05).
Additionally, the toxicological evaluation was conducted on the leaves and it was found that it does not contain toxic elements while the number of heavy metals was reported to be within the acceptable limits (Armaghan et al., 2011). Thus, *F. deltoidea* holds a good prospect to be developed as an alternative supplement for postmenopausal women to prevent oxidative stress due to estrogen depletion.

**CONCLUSION**

*Ficus deltoidea* aqueous extract given as oral supplementation at 500 mg/kg was shown to prevent the oxidative stress induced by ovariectomy in the brain, liver, heart, and kidney of rats. *Ficus deltoidea* aqueous extract supplementation also decreased the cholesterol level in the serum of the ovariectomized rats. Taken together, *F. deltoidea* has the potential to be used as a supplement to improve the health status of postmenopausal women.

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**AUTHOR CONTRIBUTIONS**

All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; agreed to submit to the current journal; gave final approval of the version to be published; and agree to be accountable for all aspects of the work. All the authors are eligible to be an author as per the international committee of medical journal editors (ICMJE) requirements/guidelines.

**CONFLICTS OF INTEREST**

The authors report no financial or any other conflicts of interest in this work.

**ETHICAL APPROVAL**

This study was granted approval from the Universiti Kebangsaan Malaysia Animal Ethics Committee (UKMAEC): FSK/BIOMED/2012/FARAH/12- Dec/483.

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