**Apium graveolens Prevents Intrauterine Growth Restriction via Suppression of Antiangiogenic Factor Production**

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**ABSTRACT**

Preeclampsia is the worldwide leading cause of fetomaternal morbidity and mortality which involves the placental dysfunction. A poor placenta and formation of non-dilated spiral artery caused uteroplacental circulation insufficiency, resulted in inadequate supply of nutrients and oxygen to support normal aerobic growth of the fetus. Apium graveolens or celery has been widely known as antioxidant, antiinflammation and antihypertensive with flavonoid-apigenin as main active compound. Apigenin can inhibit TNF-α, HIF-1α and nitric oxide blocking as major pathophysiological pathway of preeclampsia. This study was aimed to find how the Apium graveolens can improve intrauterine growth and its correlation with suppression of anti-angiogenic factor sFlt-1 in anti-Qa2 preeclampsia animal model. Twenty female BALB/c Mus musculus were divided into 4 groups: control, anti-Qa2 and anti-Qa2 with 500 and 1000 mg/kgBW celery herbs extract. The fetal weights and lengths, placental weights and serum sFlt-1 levels were measured and analyzed with One Way ANOVA and further tested with Least Significance Difference in 95% confidence interval. The result showed a difference between control and treatments group (p≤0.05) with 1000 mg/kgBW significantly increase intrauterine growth and decrease sFlt-1, then there is a negative correlation between intrauterine weight and serum sFlt-1. This study suggests that celery herbs extract (CHE) has an apigenin-flavonoid compound which can prevent intrauterine growth restriction (IUGR) via suppression of antiangiogenic factor production in preeclampsia mice model.

**Keywords**: Apium graveolens, preeclampsia, IUGR, sFlt-1

**INTRODUCTION**

Preeclampsia is present in 2%-8% pregnancies and one of the world leading cause of perinatal mortality and morbidity. In developing countries, almost ten percent maternal deaths incident is related to pregnancy hypertension with preeclampsia-eclampsia as the most major cause. Preeclampsia is referred to the disease of theories because the pathophysiology is not fully understood as many hypotheses have developed: placental vascular abnormalities, placental ischemia, free radicals, endothelial dysfunction, immunological intolerance between mother and fetus, abnormal genetic cardiovascular adaptation, nutritional deficiency, and inflammatory theories (Ahmed, 2015; Powe, 2011; Steegers, 2010). Based on the high incidence mentioned above, World Health Organization in 2011 stated that optimizing health care to prevent and treat preeclampsia is still a necessary step toward development goals. Although the hypertension and proteinuria as the clinical manifestations have occurred in late second or third trimester of pregnancy, the pathogenesis began in early pregnancy as formed of non-dilated spiral artery (Powe, 2011).

Normal pregnancy requires immune-maternal adaptation in NK cell activity for the trophoblast invasion (Lockwood, 2008). Imbalance between T helper CD4+ cell differentiation into Treg cells that induce immunologic tolerance and Th17 cells that cause inflammation or rejection reactions caused increasing inflammation cytokines; TNF-α and IL-6. This immune-maladaptation caused inadequate tolerance, chronic inflammation and poor angiogenesis then disturbed trophoblast invasion and reduced placental blood flow which supply nutrients and oxygen to fetus (Saito, 2010; Santner-Nanan, 2009). Furthermore, placental development requires vasculogenesis and angiogenesis, then a progression imbalance of the vascular endothelial growth factor (VEGF), placenta growth factor (PIGF) and angiopoietin during placental development resulted in adaptive angiogenesis in IUGR placental villi. VEGF also has VEGF receptors VEGFR-1 and VEGFR-2, which soluble form of VEGFR-1 (soluble Fms-tyrosine like/sFlt-1) has a strong antagonistic activity and
neutralizes the effects mediated by VEGF and PlGF (Ahmed, 2000). The increase in sFlt-1 in preeclampsia is also caused by a response to hypoxia that also involves genetic and immunological factors. The increasing expression of sFlt-1 increases in response to hypoxia is mediated by hypoxia inducible factors 1-α (HIF-1 α) and Angiotsensin II type1-receptor autiantibodies (AT1-AA) (Herse, 2013; Maynard, 2011).

**Apium graveolens or Celery is a common plant which has active compounds with flavonoids apigenin and apiin as main contents. Apigenin in Celery has ability as an antioxidant, anti-inflammatory (Kooti, 2017; Yao, 2010), antihypertensive (Jin, 2009; Zhang, 2002), antiischemic and antiarrhythmia (Occhiuto et al, 1991). Apigenin extracted from celery herbs has been shown as endothelial nitric oxide synthase (eNOS) stimulator and able to inhibit HIF-1α expression as a factor causing placental hypoxia by blocking its interactions with HIF-1α-Hsp90 receptors (Manolescu, 2009; Olszanecki, 2002). Yamagata (2011) also stated that the flavon structure and double bonding on C ring in apigenin content of herb celery extracts can cause binding of ROS through inhibition of Tumor Necrosis Factor (TNF) -α and lectinlike oxidized LDL receptor-1 (LOX-1).

In recent few years, many studies reported the prevention and treatment with the compound which able to bind inhibit sFlt-1 production and restore the vasculogenesis and angiogenesis function, such as VEGF121 recombinant, xanthone in mangosteen peel and thymoquinone in black cumin extracts (Rahma, 2017; Sulistyowati, 2017; Hidayati, 2015). The objective of the present study was to investigate whether the actions of Apium graveolens compound can prevent intrauterine growth restriction with reducing sFlt-1 production in preeclampsia animal model.

**METHODOLOGY**

**Experimental Design and Sampling**

This study was an experimental study using completely randomized design with twenty healthy female BALB/c mice (Mus musculus) which bred and cared in the Faculty of Veterinary Medicine, Airlangga University. All samples were randomly divided into 4 groups: control group, anti-Qa2 group, anti-Qa2+CHE 500 mg/kgBW and anti-Qa2+1000 mg/kgBW. The mice were housed in plastic cages with wooden chip bed in a temperature-controlled room (23° C), given food and water *ad libitum*, and exposed to a light-dark cycle of 12:12 hours as its circadian rhythm. All protocols were approved by the Animal Care and Use Committee (ACUC), Faculty of Veterinary Medicine, Airlangga University No. 2.KE.001.01.2018.

**Mating Procedure**

All samples were injected with 5 IU Pregnant Mare’s Serum Gonadotropin (PMSG PG 600, MSD Animal Health, Intervet Inc.), followed by human Chorionic Gonadotropin (hCG Chorulon, MSD Animal Health, Intervet Inc.) forty-eight hours later. This method was done for oestrus cycle synchronization and prepared for mating following the procedures by The Practice Committee of the American Society for Reproductive Medicine in 2008. After the injection of hCG, all samples were mated one-on-one with mature male mice (aged 7 months and weighed ± 40 g). The presence of sperm plug seventeen hours later considered as day 0 of gestations.

**Anti-Qa2 Injection**

The injection group was injected with anti-Qa2 10 ng i.p. on 1-4 days of gestations which has known that the implantation occurred approximately in a first quarter of murine pregnancy (Sones, 2016). Injection of anti Qa2 use the previous protocols by Sulistyowati (2010) that Qa2 was expressed in fifth day of gestations in normal mice, then the absence with anti-Qa2 injection i.v. has led to increasing of Heat Shock Protein (HSP-70) and Vascular Cell Adhesion Molecule-1 (VCAM-1) in preeclampsia. The previous study used the intravenous injection then we modify to intraperitoneal injection as it has similar efficient bioavailability as i.v. which is impractical and at risk of mice perivascular trauma. Turner, et al (2011) in Journal of the American Association for Laboratory Animal Science also stated that intravenous injection routinely and has potential for severe complications, including blindness, cerebrovascular stroke, permanent motor deficits, and limb gangrene. Most previous preeclampsia animal model studies also used the intraperitoneal injection, such as TNF-α injection and human preeclampsia serum injection (Wicaksono, 2015; Kalkunte, 2010).

**Celery Herbs Extraction and Treatment**

Sterilized and shade air-dried Apium graveolens herbs were extracted with 70% ethanol maceration methods in the Phytochemical Laboratory of Materia Medica, Batu, Indonesia. Powdered celery herbs were soaked and shaken in ethanol 70%, then filtered with filtration cloth. Filtrate were extracted for second and third time with additional solvent. The combined filtrate then evaporated and suspended in Na CMC 0.5%
suspension agent to a final concentration before use. The extracted celery herbs injected in treatment group in 6-15th days of gestation as known in mid-first trimester until third trimester.

**Measurement of Results**

**Samples Termination**

All samples were terminated on the 16th day of gestation as the third trimester of pregnancy which reflects the time preeclampsia onset (Sulistyowati, 2017). The surgical procedure was performed under general anesthesia using intramuscular injection of ketamine. Then, the number of fetus, any fetus with resorption and placental weight were measured. Each newborn fetus was measured using analytical instruments and the lengths were also measured and documented.

**Serum sFlt-1**

Mice blood was directly taken from the right heart then centrifuged at 6000 rpm for 10 minutes. Serum was taken and measured the sFlt-1 levels using ELISA kit (Bioassay Technology Laboratory, China, catalog number E0611Mo) and ELISA reader 450 nm wavelength.

**Statistical Analysis**

Collected data were tested with Shapiro-Wilk normality test and Lavene homogeneity test. If the data were normal and homogeneous (P≥0.05), these were continued with 5% One Way ANOVA analysis and further tested with LSD (Least Significance Difference) 5% if there are significant differences (P≤0.05). Both intrauterine growth and serum sFlt-1 also tested with Pearson correlation test.

**RESULT AND DISCUSSION**

**A graveolens in Fetal and Placental Growth**

Anti Qa-2 group injection had lower the number of fetus born compared to normal group. Anti Qa-2 also increased the number of fetal resorption and reduced other characteristics as fetal weight, fetal length and placental weight. Anti Qa-2 caused growth restriction as normal fetuses (0.233 ± 0.18 g) which bigger than anti Qa-2 injection group (0.792 ± 0.32 g) (see Table 1.), then celery herbs extract injection in both 500 mg/kgBW and 1000 mg/kgBW show increasing of number of fetus, fetal weight (0.432 ± 0.19 g and 0.472 ± 0.07 g), fetus length and placental weight (0.114 ± 0.030 and 0.129 ± 0.11), followed by reducing the number of number of fetus with resorption with P=0.00 as significant difference in all of groups. In this study, we assumed that fetal growth restriction was symmetrically indicated by shorter fetal length in anti Qa-2 group which was 11.129 mm compared to normal group which was 23.038 mm (P-value=0.0005), then celery herbs extract increasing fetus length as 13.830 mm in 500 mg/kgBB CHE and 15.818 mm in 1000 mg/kgBB CHE.

The growth restriction due to anti Qa-2 injection as compared to preeclampsia might be caused by hypoxia while ischemia was caused by reduced placental blood flow. Previous studies using the same haplotype of mice showed that Qa-2 is a murine homolog of Human Leukocyte Antigen-G (HLA-G) as non-classical Class Ib Major Histocompatibility Complex (MHC) located in Q region (Q6, Q7, Q8, and Q9) (Da Silva, 2017). HLA-G regulation has been widely reported as the pre-implantation embryonic development gene which plays an important role as Killer Inhibitory Receptor (KIR) ligand for maternal NK cell. The dysregulation of human leukocyte antigen (HLA)-G was found in both placentas and maternal sera from preeclampsia patients (Tang, 2015; Djurisic, 2014; Durmanova, 2013). The disruption of placental development led to limited cytrophoblast invasion of the arteries in the superficial decidua and the myometrial segments remained narrow and undilated (Maynard, 2011). The intrauterine growth restriction (IUGR) defined as is defined as a rate of fetal growth that is less than normal for the growth potential of a specific infant as per the race and gender of the fetus. The IUGR in preeclampsia is caused by maternal factor as preeclampsia is the disease with the effect on blood circulation, resulted in decrease in uteroplacental blood flow and lead to IUGR (Sharma, 2016). The placental factor in preeclampsia IUGR involved the decrease of terminal villi number and surface area which shows a malfunction of vascularization in pregnancy. As mentioned before, the adequate trophoblast invasion and an increase in uteroplacental blood flow is required with increase in gestation to meet the demand of growing fetus (Zygmunt, 2003). The intrauterine growth restriction caused by anti Qa-2 injection was also significantly shown (P-value=0.0005) which reduced placental weight (0.085 ± 0.05 g) compared to normal group (0.150 ± 0.03 g).

According to Cunningham (2014), placental dysfunction in preeclampsia is indicated by the miointima cells proliferation and tunica media necrosis. This mechanism started with lipids that accumulate in miointima cells and subsequently in macrophages and caused atherosis and uteroplacental blood flow disruption.
Furthermore, celery has been commonly known to have antihypertensive effects through antioxidant-relaxation dependent pathway, increased nitric oxide (NO) and antiproliferation of vascular smooth muscle (Jorge, 2013; Jin, 2009; Zhang, 2002). This effect was shown to increase placental blood flow, as indicated by the increased placental weight of the anti-Qa2 group 0.085 ± 0.05 g to 0.114 ± 0.30 g and 0.129 ± 0.11 g in the celery group 500 and 1000 mg/kgBW, followed by the increased of fetal weight.

*Apium graveolens* in sFlt-1 Serum Levels

Figure 1 shows that anti-Qa2 group having higher serum sFlt-1 levels than control (p ≤0.05), then the anti-Qa2 group with the treatment of 500 celery herbal extracts and 1000 mg/kgBW show decreased sFlt-1 levels. In addition, at the sFlt-1 level in the group with the CHS treatment 1000 mg/kg BW did not differ statistically with the control group (p≥ 0.05). Celery has been known to inhibit the increase of TNF-α and HIF-1α which are the activation paths of AT1-AA which causes an increase in sFlt-1. This sFlt-1 supression by celery restores the VEGF angiogenesis function, decreases the cloting factor and restores the endothelial function to improve the blood flow to the placenta (Yamagata, 2011; Manolescu, 2009). This effect has been proved with Pearson correlation test among sFlt-1 levels, fetal weights, fetal lengths, placental weights which shows that sFlt-1 has strong negative correlation with each intrauterine growth characteristic (P≤0.05). These results indicated that the decrease of sFlt-1 by celery caused an increase in the growth of fetus and placenta.

Table II shows that sFlt-1 has negative correlation to fetal weight, fetal length and placental weight with high coefficient of Pearson correlation -0.648, -0.610, and -0.621 respectively. Herraiz (2015) has reviewed the role of sFlt-1 as angiogenesis-related biomarkers in placental dysfunction which resulted in inhibit intrauterine growth. sFlt-1 mRNA as alternative splicing of the Flt-1 gene has generated in response to hypoxia, then sFlt-1 is produced and secreted from the placenta to the maternal circulation, causing a reduction of the inhibition of proangiogenic factors. This mechanism will disrupt the remodeling of the spiral arteries in their myometrial segments during normal pregnancy and regulation of the flow pressure to the intervillous space. This led to the appearance of hypoperfusion-reperfusion phenomena,
Apium graveolens Prevents Intrauterine Growth

Table I. The fetal growth in each group, shows the number of fetus born and fetus with resorption. Fetal weights, lengths and placental weights were statistically different *p≤0.05

<table>
<thead>
<tr>
<th>Comparison of Variable</th>
<th>Number of pups</th>
<th>Number of pups with resorption</th>
<th>Pup Weights (g)</th>
<th>Pup lengths (mm)</th>
<th>Placental weights (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control (n=5)</td>
<td>86</td>
<td>1</td>
<td>0.792±0.292</td>
<td>23.038±6.063</td>
<td>0.150±0.034</td>
</tr>
<tr>
<td>anti Qa2 mice (n=5)</td>
<td>78</td>
<td>14</td>
<td>0.233±0.179</td>
<td>11.129±4.272</td>
<td>0.085±0.039</td>
</tr>
<tr>
<td>CHE 500 mg/kgBW (n=5)</td>
<td>75</td>
<td>10</td>
<td>0.432±0.188</td>
<td>13.383±4.022</td>
<td>15.818±2.038</td>
</tr>
<tr>
<td>CHE 1000 mg/kgBW (n=5)</td>
<td>87</td>
<td>11</td>
<td>0.472±0.075</td>
<td>0.114±0.029</td>
<td>0.129±0.109</td>
</tr>
<tr>
<td>P-value</td>
<td>-</td>
<td>-</td>
<td>0.000*</td>
<td>0.000*</td>
<td>0.000*</td>
</tr>
</tbody>
</table>

Table II. Matrix of bivariate correlations among sFlt-1 levels, Fetal Weights, Fetal Lengths, Placental Weights

<table>
<thead>
<tr>
<th>Variable</th>
<th>sFlt-1 levels</th>
<th>Fetal Weights</th>
<th>Fetal Lengths</th>
<th>Placental Weights</th>
</tr>
</thead>
<tbody>
<tr>
<td>sFlt-1 levels</td>
<td>Pearson correlation</td>
<td>1</td>
<td>-0.648**</td>
<td>-0.610**</td>
</tr>
<tr>
<td></td>
<td>P-value</td>
<td></td>
<td>0.002</td>
<td>0.004</td>
</tr>
<tr>
<td>Fetal Weights</td>
<td>Pearson correlation</td>
<td>1</td>
<td>0.907**</td>
<td>0.776**</td>
</tr>
<tr>
<td></td>
<td>P-value</td>
<td></td>
<td>0.000</td>
<td></td>
</tr>
<tr>
<td>Fetal Lengths</td>
<td>Pearson correlation</td>
<td>1</td>
<td>0.823**</td>
<td></td>
</tr>
<tr>
<td></td>
<td>P-value</td>
<td></td>
<td>0.000</td>
<td></td>
</tr>
<tr>
<td>Placental Weights</td>
<td>Pearson correlation</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td></td>
<td>P-value</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**correlation is significant at the 0.01 level (2-tailed)

REFERENCES
Ahmed, A. & Ramma, W., 2015, ‘Unravelling the theories of pre-eclampsia: Are the protective pathways the new paradigm?’, Br. J. Pharmacol. 172, 1574–1586
Herraiz, I. et al., 2015, ‘Angiogenesis-related biomarkers (sFlt-1/PLGF) in the prediction

CONCLUSION
This study suggests that herb celery extract in 1000 mg/kgBW can prevent intrauterine growth restriction via suppression of antiangiogenic factor production in preeclampsia mice model. The possible pathway is a celery apigenin-flavonoid compound which inhibit TNF-α and HIF-1α actions.

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The practice Committee for the American Society for Reproductive Medicine, 2008. "Ovarian stimulation and factors affecting ovarian response, including stimulation protocols, long-acting follicle-stimulating hormone, GnRH analogs, and gonadotropin dose.


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Kinase 1 (sFlt-1) and Blood Pressure of Pregnant Mice, J. Trop. Life. Sci. 5, 8–13