Effect of Ethanol Extract of Chayote (*Sechiumedule.Jacq.Swartz*) on the Activity of Glutathione Peroxide (GPx) in House Mice (*Musmusculus* L) Strain DD Webster Hyperglycemia Induced by Streptozotocin (STZ)

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ABSTRACT

Background: Streptozotocin can cause hyperglycemia in guinea pig through the mechanism of oxidative stress which damages pancreatic β cells. Ethanol extract of chayote can decrease oxidative stress. This study aimed to determine the effect of Ethanol Extract of Chayote ((EEBLS) Ekstrak Etanol Buah Labu Siam) on decreasing blood sugar content and increasing the activity of glutathione peroxide enzyme.

Subjects and Method: This was an experimental study. The samples were using random sampling technique. The control group was using male white mice (*Musmusculus* L) Strain DD Webster which randomized into four groups: negative control group, positive control group, group which got EEBLS of 100 mg/kgBB, and group which got EEBLS of 200 mg/kgBB.

Results: The result of the research showed that there was significant decrease in blood sugar, compared with the control group. There was insignificant decrease in the activity of glutathione peroxide enzyme, compared with the control group.

Conclusion: The conclusion of the research was that EEBLS of 200 mg/kgBB decreased blood sugar content of mice significantly, but there was no significant change in the activity of glutathione peroxide enzyme when EEBLS was given to the mice.

Keywords: streptozotocin, oxidative stress, antioxidant, flavonoid

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BACKGROUND

Diabetes Mellitus (DM) is a group of metabolic disease with hyperglycemia characteristics as the result of insulin excretion disorder, the work of insulin, or both of them. Prolonged hyperglycemia is related to long term damage, dysfunction, and failure of various organs (American Diabetes Association, 2014).

Persistent hyperglycemia in DM patients will cause the increase in oxidative stress as the result of imbalance between free radical and natural antioxidant formed by body (Sheikhpour, 2013). Hyperglycemia in guinea pig can be induced by using streptozotocin (STZ).

STZ or deoxy-2- [3-(methyl-3-nitro-nitrosoureido)-D-glucopiranoside] is obtained from Streptomyces achromogenes and structurally, it is nitrosourea derivative (Nugroho, 2006; Srinivasan dan Ramarao, 2007). The mechanism of STZ is particularly mediated by the forming of free radical Reactive Oxygen Species (ROS) and Reactive Nitrogen Species (RNS) which cause the damage in pancreas β cells (Srinivasan dan Ramarao, 2007). ROS includes superoxide (•O₂⁻), hydroxide (•OH), peroxide (•RO₂⁻), hydroperoxide (•HRO₂⁻), non-ra-
dical hydrogen peroxide (H$_2$O$_2$), and hydro-
cholorous acid (HOC1). RNS includes nitric oxide (\textit{NO}), nitrogen dioxide (\textit{NO$_2$}), non-
radical peroxyxinitrite (ONO), nitrous oxide (HNO$_2$), and alkyl peroxynitrates (RONOO).

Glutathione peroxide (GPx) enzyme is one of enzymatic antioxidant (Moussa, 2008; 
Johansen et al., 2005; Rahman, 2007). Compared with superoxide dismutase-1 (SOD-1) and superoxide dismutase-2 (SOD-2) content and catalase, GPx content is still higher in islet pancreas cells so that the accurate protection for pancreas $\beta$ cells is GPx (Robertson dan Harmon, 2007). In animal cells, antioxidant enzyme which is prioritized for H$_2$O$_2$ detoxification is GPx, while catalase has much lower affinity for H$_2$O$_2$, compared with GPx (Jurcovic et al., 2008). Low GPx enzyme is caused by the disorder which is related to free radical (Judge et al., 2005). High activity of GPx enzyme is able to protect isle pancreas cells which are caused by glucose toxicity that appears through the process of oxidative stress (Tanaka et al., 2002). The increase in the activity of GPx enzyme in the pancreas of mice is caused by the mechanism of detoxification compensation of peroxide hydrogen. In this research, it is found that the activity of glutathione reductase (GR) and glutathione-S-transferase (GST) does not change in the pancreas of diabetes mice which indicates that GR and GST enzymes do not play a significant role in oxidative stress protection in pancreas beta cells (Erejuwa et al., 2010).

Antioxidant is molecule which can forestall cell damage caused by free radical by stabilizing it. It can be grouped based on its origin to become natural and synthetic antioxidant. The effectiveness of natural antioxidant is lower than that of synthetic antioxidant, but it is safer because it is not contaminated by other chemical substances and it is easy to obtain, while the use of Butylated Hydroxy Toluena (BHT) is limited since it is carcinogenic (Rahman, 2007). The best natural antioxidant comes from fruit and vegetables (Dimitrios, 2006), and one of the sources of natural antioxidant is Chayote (Sechium edule jacq. Swartz) (Firdous, 2011). It has a lot of benefits. Many Indonesian people use it as vegetable besides food industry, and its pectin content can be used in pharmacy and cosmetics (Daryono, 2002). Besides that, its pectin content has the effect of anti-cholesterol, anti-colon cancer and DM (Agustini, 2006).

Based on the study on phytochemical screening of ethanol extract and the analysis on KLT (thin layer chromatography) of chayote extract conducted by Marliana et al., (2005), it is found that it has alkaloid, saponin, kardenolin/ bufadienol, and flavonoid (Marliana et al., 2005). Besides that, EEBSL (ekstrak etanol buah labu siam) can also function as anti-proliferation (Iniguez, 2012), anti-epilepsy, depressant power of central nerve system (Firdous, 2012). The study on the effect EEBSL as an anti-diabetes has also been conducted (Maity, 2013). Its extract water also has neuroprotective effect (Mumtaz et al., 2013). Chayote extract is useful as anti-microbe on bacteria and fungi in digestive tract. Its leaves and fruit have the effect of diuresis, cardio-protective, and anti-inflammation, and its leaves have been used as an anti-arteriosclerosis, hypertension, and can dissolve kidney stones (Kamble et al., 2008; Gordon et al., 2000). Active substance in chayote can be extracted by using certain solution. Generally, solvent which is used to extract natural substances is ethanol because it is better than water (Lukiati, 2012). Ethanol extract in chayote is also used as hepatoprotector probably because of its antioxidant activity (Firdous et al., 2012). There are a lot of benefits of chayote; but unfortunately, not many people get scien-
tific information related to antioxidant content and the benefits of chayote, especially for diseases caused by free radical. Besides that, previous researches still dealt with the effect of antioxidant on hyper in mice; therefore, the researcher is interested in analyzing the effect of EEBLS on the activity of glutathione peroxide in white male mice which hyperglycemia is induced by STZ.

### SUBJECTS AND METHOD

This study used experimental. The samples were taken by using simple random sampling technique, using white male mice (*Mus musculus* L.) Strain DD Webster which was randomized into 4 groups: negative control group, positive control group STZ 60 mg/kgBB, group which got STZ 60 mg/kgBB and EEBLS of 100 mg/kgBB, and group which got STZ 60 mg/kgBB and EEBLS of 200 mg/kgBB. Mice (*Mus musculus* L.) strain DD Webster DM when, in the fourth day after STZ is given, blood glucose content is > 250 mg/dl. When blood sugar content increases, EEBLS is given until the 28th day.

### RESULTS

The guinea pig used in this research is healthy mice with their normal blood sugar content.

Glucometer is used for KGD measurement in order to make sure that the mice are normal. The result of KGD has been observed until the 28 day of the experiment as seen in Picture 1.

From the result of the study, it is found that the data of measuring the activity of peroxide glutathione enzyme from the blood of white male mice is found after the treatment within 28 days, as it is seen in Table 1.
Table 1. The result of the activity of peroxide glutathione of white male mice after the treatment within 28 days

<table>
<thead>
<tr>
<th>Group</th>
<th>Enzyme Outcome</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>P1</td>
<td>420.04</td>
<td>78.95</td>
</tr>
<tr>
<td>P2</td>
<td>425.98</td>
<td>22.50</td>
</tr>
<tr>
<td>P3</td>
<td>427.57</td>
<td>73.82</td>
</tr>
<tr>
<td>P4</td>
<td>418.46</td>
<td>54.90</td>
</tr>
</tbody>
</table>

Information:
P1 Negative control (mg/dl)
P2 Positive control (mg/dl)
P3 EEBLS 100 mg (mg/dl)
P4 EEBLS 200 mg (mg/dl)

DISCUSSION

Based on Picture 1 above, it is found that STZ damages pancreas β cells so that KGD increases. KGD groups of P2, P3, and P4 which have been induced by STZ is higher than that of P1 group. There is the decrease in KGD in P2 group starting from the 21st day while the decrease in KGPD in P3 and P4 groups has been known has been known in the 7th day. The group which gets the dosage of 200 mg undergoes the lowest decrease in KGD which is in line with the result of the research conducted by Maity et al., (2013). The decrease in KGD in the treatment group is caused by giving EEBLS which contains flavonoid (Siciliano et al., 2004; Marliana et al., 2005). Flavonoid is anti-diabetes by disturbing glucose absorption in intestines, improving glucose tolerance, disturbing carbohydrate metabolism through enzyme obstruction α amylase and enzyme α glucosidase, stimulating glucose excretion in peripheral tissues; besides that, flavonoid also stimulates insulin secretagogues and acts like insulin in stimulating glycogen synthesis (insulin mimetics) (Brahmachari, 2011; Piparo, 2008; Sahgal et al., 2010). The decrease in KGD in P2 group can occur because of self-recovery by the body through the improvement of pancreas β cells and new cell splitting (mitosis) gradually. The decrease in the number of pancreas β cells in hyperglycemia animals is known in the 7th day and continuously decreases until the 28th day (Erwin et al., 2012). The decrease in KGD in P3 and P4 groups begins to be known after EEBLS is given in the 7th day and begins to be normal in the 28th day. This indicates that the decrease in KGD in P3 and P4 groups is due to the influence of flavonoid in EEBLS, while the improvement of pancreas β cells is indicated by the improvement of KGD in the mice.

The study shows significant difference in the activity of enzyme glutathione peroxide in the white male mice. The lowest activity of glutathione peroxide enzyme is found in the treatment 4 group which indicates that oxidative stress in P4 group is lower than that in the P1 negative control group which can be seen from the enzyme activity in P4 group which is lower than those in P1, P2, and P3 groups. The decrease in the activity of glutathione peroxide enzyme is in line with the result of the research conducted by Bhatt et al., (2011). C-glycosyl in methanol Enicostem malitto-
rsle Blume extracts decreases the activity of GPx in nephrotoxic mice that are induced by gentamicin (Bhatt et al., 2011). The result of the study conducted by Gargari et al., in which extract of the grape nuts which contains proantisioncan decrease the activity of GPx although the decrease is not significant (Pourhassem et al., 2011)

EEBLS 200 mg/kg BB can significantly decrease blood glucose content of mice and there is no significant change in the activity of glutathione peroxide with giving EEBLS.

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