The effect of castration towards the microstructure of Hippocampus in Wistar rats

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ABSTRACT

Background: Male ageing is mostly referred to as the declining of the capacity of a man due to age. Even though the physical changes cause by ageing are noticeable, there are several effects of ageing in a male that are concealed by the physical appearance, such as a lower level of testosterone. This study aims to determine the effect of lower testosterone towards the neuron, neuroglia and vasculature of Hippocampus.

Methods: An analytic observational study with two-group posttest-only randomized experiment with treatment was conducted among 10 Wistar rats (Rattus norvegicus). Variables assessed in this study were the number of neurons, neuroglia and vasculature of Hippocampus. Data were analyzed using SPSS version 17 for Windows.

Results: The average number of neuron cells of Hippocampus was significantly lower in the treatment group compared with control (p=0.014 and 0.004, respectively). The significantly lower in the number of neuroglia and vasculature of Hippocampus was also found in the treatment group compared with the control group (p=0.014 and 0.004, respectively) following castration.

Conclusion: Hence, we can conclude that the castration has a significant effect in lowering the level of testosterone, number of neurons, neuroglia and vasculature in the Hippocampus of Wistar rats.

Key Words: Testosterone, Hippocampus, Neurons, Neuroglia, Vasculature.


INTRODUCTION

Male ageing mostly refers to the declining of the capacity of a man due to age.1 Even though the physical changes cause by ageing are noticeable, there are several effects of ageing in a male that are concealed by the physical appearance such as a lower level of testosterone.2

Testosterone is a male steroid hormone that plays a significant role in the reproductive system of a male to make sure the male can be sexually stimulated.3 The production of testosterone takes place in the testes of a male.3 The production first starts in the hypothalamus sends a message to the pituitary gland, and from there the instruction given to the testes about the amount of testosterone should be produced.4

Lowered level of testosterone in a male causes effects to a male in various ways. Apart from being sexually dysfunction, a male is capable of developing other symptoms as the consequences of a lower level of testosterone.5 For instance, a male with a low level of testosterone will be having mood problems or might be in depression.6 Besides, a lower level of testosterone also causes the libido and muscle mass of the male to decrease as well.5,6

Hippocampus of human brain functions in storing memories and aid in thinking and learning skills.7,8 The organ located in the medial temporal lobe of the brain is where the neurogenesis takes place.7 This regeneration new cells every time and inter-connects with each other to interpret and store memories.9 Both the spatial and long term memories will be saved as the cells generate and functions normally.9

Based on those mentioned above, this study aims to determine the effect of castration towards the microstructure of Hippocampus in Wistar rats through the evaluation of the number of neurons, neuroglia, and vasculature.

METHODS

This is analytic observation research with two-group posttest-only randomized experiment with treatment which conducted among 10 Wistar rats. This research will be carried out at the Animal Laboratory of Faculty of Medicine, Udayana University with several preparations will be done at distinctive places. This research takes about two months to be completed, which is from April 2019 to June 2019. The population target for this study is male rat species of Rattus Norvegicus age four months old. All the rats selected are to be in two months to be completed, which is from April 2019 to June 2019. The population target for this study is male rat species of Rattus Norvegicus age four months old. All the rats selected are to be in two months to be completed, which is from April 2019 to June 2019. The population target for this study is male rat species of Rattus Norvegicus age four months old. All the rats selected are to be in

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into two groups, with five rats in each of the groups, respectively. The rats in the control group (P0) were kept natural. Whereas, the treatment group (P1) were castrated to reduce the level of testosterone in the male rats. Both the rats in the control and treatment group were kept for 30 days before dissected and remove the brain to observe the changes that take place in the microstructure of the Hippocampus of the brain.

The use and dispose of these animal samples were followed according to the procedure of the Ethical Commission at the end of the research. The samples that have been used will be disposed of appropriately in a plastic bag and will be buried. Independent variable of this research is castration done to the male rats to lower the testosterone level. Whereas, the dependent variable is the number of neurons, neuroglia and vasculature in the Hippocampus of the brain. The controlled variables in this research are the age, gender, food and environment of the samples.

Analysis of research data to calculate the mean and standard deviation of the number of neurons, neuroglia and vasculature of Hippocampus. Normality analysis data of the number of neurons, neuroglia and vasculature by the test of Shapiro Wilk, with the sense of significance $a = 0.05$, obtained result $p > 0.05$ which means data in all groups were normally distributed so that it continued with parametric tests. Analysis of variance homogeneity data of the number of neurons, neuroglia and vasculature by using Levene test, variance result obtained homogeneity as the value of $p > 0.05$. Data were analyzed using SPSS version 17 for Windows to determine the significant difference between two groups in the number of neurons, neuroglia and vasculature.

**RESULTS**

Normality test was carried out by using the Shapiro Wilk test because the sample of subjects was small in less than 50 subjects. The results of the analysis show that the data were normally distributed ($p > 0.05$), and the data variance was also homogeneous ($p > 0.05$) (Table 1).

Data analysis with an Independent T-Test have been done to determine the mean difference in the number of neurons, neuroglia and vasculature between groups. The Independent T-Test results were described in Table 2, and Figure 1 show a data descriptive regarding the number of neuron, neuroglia and vasculature between groups.

The test showed that the mean number of neurons between groups was significantly different ($p = 0.007$). Also, the mean number of neuroglia between groups was significantly different ($p=0.014$). Besides, the test showed that the mean number of vasculature between groups was significantly different ($p = 0.004$) (Table 2). The image of differences in the hippocampus cell in the control and treatment groups can be seen in Figure 2.

### Table 1 Normality test by Shapiro Wilk

<table>
<thead>
<tr>
<th>Variable</th>
<th>Group</th>
<th>Statistic</th>
<th>Df</th>
<th>Sig.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of Neurons</td>
<td>Control</td>
<td>0.943</td>
<td>5</td>
<td>0.688</td>
</tr>
<tr>
<td></td>
<td>Treatment</td>
<td>0.940</td>
<td>5</td>
<td>0.665</td>
</tr>
<tr>
<td>Number of Neuroglia</td>
<td>Control</td>
<td>0.935</td>
<td>5</td>
<td>0.631</td>
</tr>
<tr>
<td></td>
<td>Treatment</td>
<td>0.874</td>
<td>5</td>
<td>0.282</td>
</tr>
<tr>
<td>Number of Vasculatures</td>
<td>Control</td>
<td>0.887</td>
<td>5</td>
<td>0.342</td>
</tr>
<tr>
<td></td>
<td>Treatment</td>
<td>0.882</td>
<td>5</td>
<td>0.318</td>
</tr>
</tbody>
</table>

### Table 2 Independent t-test to the number of neuron, neuroglia and vasculature between groups

<table>
<thead>
<tr>
<th>Variables</th>
<th>Group</th>
<th>Mean Difference</th>
<th>SE Difference</th>
<th>$p$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of Neurons</td>
<td>P0</td>
<td>33.664</td>
<td>9.326</td>
<td>0.007</td>
</tr>
<tr>
<td></td>
<td>P1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Number of Neuroglia</td>
<td>P0</td>
<td>22.466</td>
<td>7.123</td>
<td>0.014</td>
</tr>
<tr>
<td></td>
<td>P1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Number of Vasculatures</td>
<td>P0</td>
<td>22.600</td>
<td>5.768</td>
<td>0.004</td>
</tr>
<tr>
<td></td>
<td>P1</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

P0=Control; P1=Treatment; SE: Standard Error; P-value was considered significant if less than 0.05
DISCUSSION

Circulating androgens are converted to metabolites that have a wide range of biological activities, in different regions of the brain.\textsuperscript{10} This extensive metabolism provides mechanisms for both control, in terms of regulating the cellular responses that can be obtained via modulation of the activity of the relevant rate-limiting enzymes, as well as for diversification of the cellular effects of the circulating hormones.\textsuperscript{11}

Testosterone is made into estradiol in few areas of the brain, including the Hippocampus.\textsuperscript{12} It is also being used, in both neurons and glia, to convert to the potent androgen, dihydrotestosterone (DHT).\textsuperscript{12,13} Called as aromatase, the enzyme that converts testosterone to estrogen was produced in the brain’s Hippocampus and cerebral cortex in a variety of species that includes humans.\textsuperscript{13}

Aromatase in Hippocampus serve as a medium in making long-term memories and spatial memory, and the cerebral cortex.\textsuperscript{14} This happens as the synaptic connection takes place in the Hippocampus with the presence of aromatase.\textsuperscript{14} The interconnection between the neurons aids in the process of interpreting information which is essential to memory, attention, awareness and thought. Removing out aromatase also reduces expression of CREB, a major transcription factor known to play a primary role in learning and memory, as well as a neuron-nourishing brain-derived neurotrophic factor.\textsuperscript{15}

The result of androgens affecting the Hippocampus includes a difference in neurogenesis in the subgranular zone of the dentate gyrus.\textsuperscript{16} The changes cause by androgens on the hippocampal structure are the difference in glia, in particular astrocytes.\textsuperscript{16} These effects are important not only because of the role of glia in neurosteroid biosynthesis but also because glia mediate some of the neuronal effects of steroid hormone action.\textsuperscript{17}

Result of this research shows that the rats which were castrated had its testosterone level to be decreased. This eventually reduces the microstructure of the Hippocampus, which are the neuron, neuroglia and vasculature. The result of this research shows the number of neurons, neuroglia and vasculature in the treatment group are lower compared to the control group. A previous study has been noticed that low testosterone level will affect the adult neurogenesis through several pathways.\textsuperscript{18}

CONCLUSION

Based on the present research that has been conducted, we conclude that the number of neurons, neuroglia, and vasculature in the Hippocampus of castrated rats is lower compared to the number of neuron in the Hippocampus of non-castrated rats.

CONFLICT OF INTEREST

There is no competing interest regarding the manuscript.
ETHICS CONSIDERATION

This research has been approved by the Ethical Commission of Faculty of Medicine, Universitas Udayana, Bali, Indonesia.

FUNDING

None.

AUTHOR CONTRIBUTION

All of the authors are equally contributed to the study from the conceptual framework, data gathering, data analysis, until reporting the results of study through publication.

REFERENCES


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