

THE DENTATE GYRUS NEURONAL DENSITY IN OVARIECTOMIZED AND NORMAL RATS

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Summary

Ovarian hormones loss and memory loss have been shown to be related in those suffering AD. Estrogen receptors have been found in several places throughout the brain and regulate a wide variety of non-reproductive functions in the central nervous system by interacting with several molecular and cellular processes. The aim of this study is the evaluation of dentate gyrus neuronal density in ovariectomized rats in compare with normal rats. Eighteen female Wistar Rats weight (300-350 g) were completely divided into two experimental groups and one control group. Animals were anesthetized under interaperitoneal injection of a mixture (1:2) of 10% ketamin and 2% xylazine then the gonads were removed. After one month and two, Animals were decapitated and their brain dissected, fixed in 10% formalin, sectioned in 7 μ m thickness and stained by H.E. By applying stereological techniques and systematic random sampling scheme the neuronal density of dentate gyrus were estimated. Statistical analyses showed significant decrease ($p>0.05$) in the dentate gyrus neuronal density in experimental groups. Then reduceing the steroidal hormones hasnot effect in neuronal degeneration in dentate gyrus.

Key words: dentate gyrus, stereology, ovaectomy

Running title: dentate gyrus neuronal density in ovariectomized rats

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Introduction

It appears that estrogen stimulates brain cell growth in the hypothalamus and hippocampus. Therefore, estrogen apparently plays a significant role in promoting brain activity in regions that are responsible for cognition. Accumulating evidence indicates that ovarian hormones regulate a wide variety of non-reproductive functions in the central nervous system by interacting with several molecular and cellular processes (1). Estrogen loss and memory loss have been shown to be related in those suffering AD. Estrogen receptors have been found in several places throughout the brain including the hypothalamus, preoptic area, anterior pituitary, and most importantly, the hippocampus (2).

The hippocampus is an important structure for memory processing. It is a particularly vulnerable and sensitive region of the brain that is also very important for declarative and spatial learning and memory (3). Hippocampus neurons are vulnerable to seizures, strokes, and head trauma, as well as responding to stressful experiences. At the same time they show remarkable plasticity, involving long-term synaptic potentiation and depression, dendrite remodeling (synaptic turnover, and neurogenesis in the case of the dentate gyrus (4).The hippocampus has been implicated in certain short-term memory. Indeed hippocampus lesions often produce short-term memory deficits (5).The hippocampus is preferentially susceptible to a wide variety of toxic insults and disease processes, including hypoxia-ischemia and hypoglycemia (6).Metabolic diseases such as diabetes and obesity have been associated with increased vulnerability to stress (7) and cognitive dysfunction (8).

In animal models, ovarian hormones have been shown to influence memory via actions on neurons, particularly in the hippocampus. Gonadal hormones affect the nervous system in ways that extend beyond their essential actions of regulating gonadotropin and PRL secretion and modulating sexual behavior(9). Confronting the efficacy of a regenerative therapeutic is the degenerative environment that is characterized by neuronal loss, physical plague and glial scar barriers and inflammation(10,11), But perhaps more fundamentals from a regenerative prospective, are changes in the biochemical milieu of steroid and peptide growth factors, cytokines and neurotransmitter systems(11,12,13). Data from multiple researches indicate that gonadal steroid hormones and their metabolites can promote neural health whereas their decline or absence is associated with reduction in neural health and increased risk of neurodegenerative diseases including Alzheimer's (14,15). In ovariectomized mice with or without estrogen replacement, regeneration of the sciatic nerve after crush injury was studied. Functional recovery, quantified with sciatic functional index was significantly accelerated in estrogen – treated mice throughout the regeneration (10). Sexosteroid hormones are potential neuroprotective candidates following CNS injury (16) and can play an important role in promoting and enhancing repair after traumatic brain injury and stroke. Although many of its specific actions on neuroplasticity remain to be discovered. There is growing evidence that these hormones may be a safe and effective treatment for traumatic brain injury and other neural disorders in humans (17). In this research, the neuroprotective actions of testosterone on populations of injured rat spinal motoneurons will be discussed. The aim of present experimental design was to evaluation of dentate gyrus neuronal density in ovariectomized rats in compare with normal rats.

. Materials and methods

All experiment was conducted in faculty of science, Islamic Azad University of Mashhad, Iran (2011-2011).

Animal subjects:

Eighteen female, Wistar rats weighting between 300-350 g served as subjects for these experiments. All animals were housed individually and maintained on a 12/12 light/dark cycle, with lights on at 6.00h. Ambient temperature in the animal facility was kept at $22\pm 2^{\circ}\text{C}$. Food and water was given ad libitum.

Groups:

1. Controls (N=8)

2.,3. Ovariectomized groups (N=8)

(In group2, the time for brain sampling was 1 month after ovariectomy but in group 3 was 2 month.)

Surgery:

Animals were anesthetized under interaperitoneal injection of an initial dose of 60mg kg⁻¹ketamine and 6 mg kg⁻¹ xylazine (ip). Then gonads were removed. In group 2 after one month and in group 3 after 2 month, Animals were anesthetized with sodium pentobarbital(64mg/kg) and decapitated. The whole brain was removed and fixed in 10% paraformaldehyde. NaCl was added to the fixative to make the tissue float in order to overcome deformities during the fixation period(18). Paraffin embedded tissue blocks were sectioned at 7μ thickness coronally and stained. A uniform random sampling scheme was employed so that about 10 sections from each block were sampled. With each section thus selected its immediately preceding neighbor was also collected. Sections were stained with toluidine blue staining method with special buffer of acetic acid, sodium acetate and distilled water (PH=4.65). Neuronal density in dentate gyrus was determined, using stereological counting technique; physical dissector (19).

The dissector principle was used to determine the numbers of neurons in each section. Form each section and it's adjacent neighbor two photos were taken, one from each section with a final magnification of 100. A two-dimensional unbiased counting frame was overlaid in a uniform, random manner on to regions of any two photos taken of both sections. Those cell nuclei selected by the frame on the reference plane but not appearing on the adjacent look-up frame section were deemed to have their tops in the volume described by the product of the area of the counting frame and the distance between sections. These nuclei were counted (Q) to provide the numerical density of cells (NV) in the ventral horns of 100-spinal cord according to the equation:

$$NV = \frac{\sum a}{\sum frame \times V_{di\ sec\ tor}}$$

Where $\sum a$ is the sum of counted neurons, h is the depth of the dissector equal to the section thickness (7 micron) and a (frame) is the scaled area of the dissector frame (20).

Statistical analyze

The ratio of numerical density of neurons in samples of brain was then used as an index of neuronal death. All quantitative data were analyzed using ANOVA and t-test. Student's t test was used for comparison when only 2 groups were analyzed. Statistical significance was chosen as p<0.05.All results are reported as mean ± SEM.

Results

The effects of gonad hormone on the numbers of intact neurons in the dentate gyrus region of hippocampus after ovariectomy in rats are shown in (Fig.1,2,3).

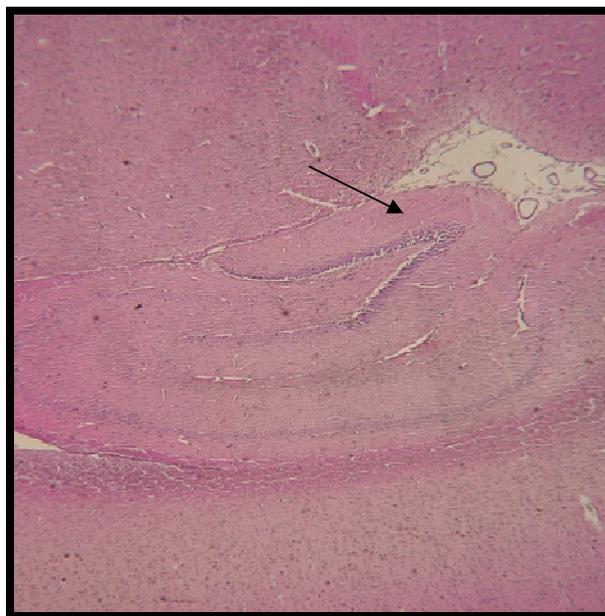


Fig.1. Photomicrograph illustrates neurons of the dentate gyrus region of hippocampus at magnification of (20×). spiks show the dentate gyrus.

1-The control group revealed healthy neuronal cells amounted by $(19642 \times 10^{-9} \pm 544)$ intact neurons. The neuronal cells in ovariectomized (1) were $(18803 \times 10^{-9} \pm 665)$ and ovariectomized (2) group $(19634 \times 10^{-9} \pm 479)$.

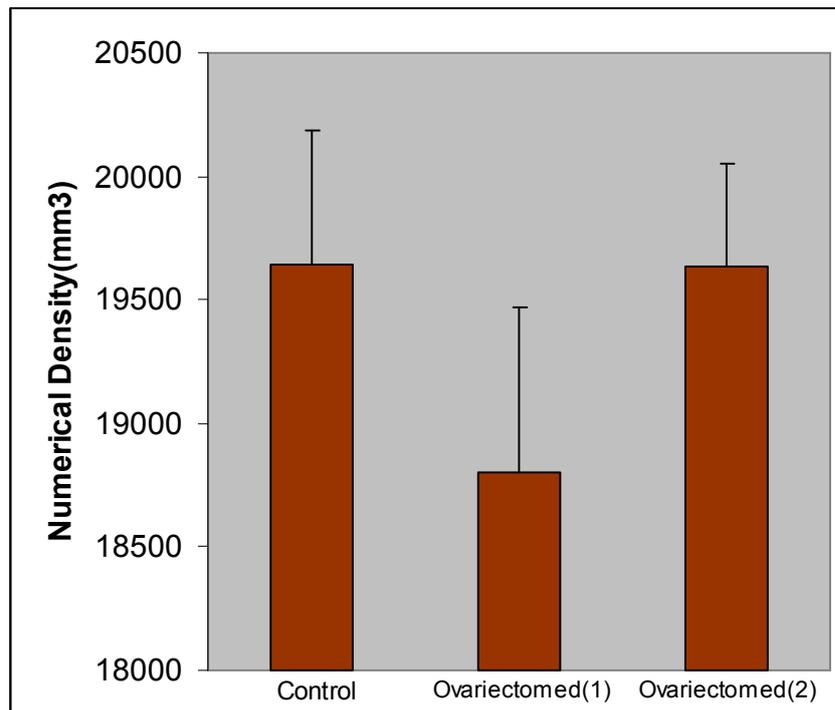


Fig.2: Effects of gonad hormone on the intact neurons numerical density of hippocampus dentate gyrus region in rats. Results are expressed as Mean± SD of 8 rats and data were analyzed by one-way ANOVA followed by Tukey-kramer multiple comparisons test.

2- As has shown in fig.2, ovariectomy resulted in massive neuronal damage manifested remarkable decrease in the number of normal appearing neurons after one month.

3- After two month this reduction was not clear more. There was not a remarkable decrease in this group in compare with ovariectomized (1) ($P>0.05$).

4- When compare ovariectomized (1) with (2), there was not any significant changes in numerical density (Fig.3).

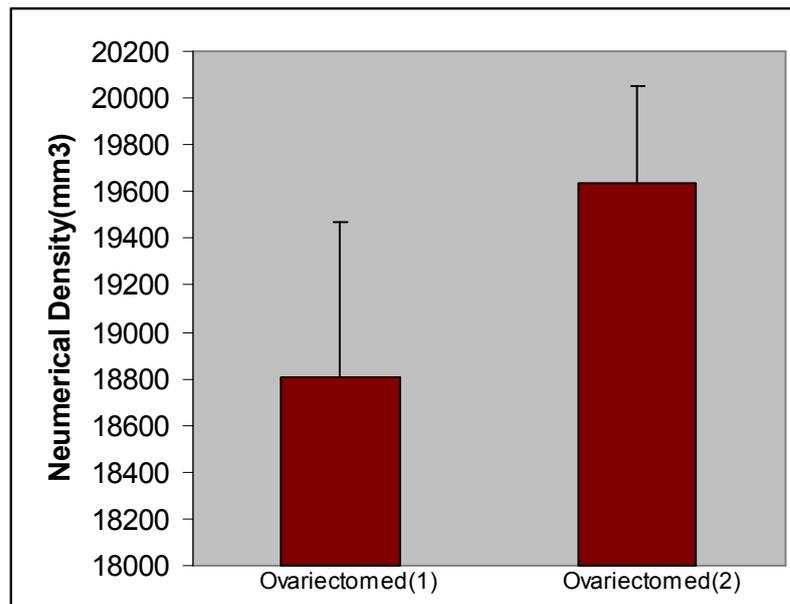


Fig.3. Comparison between dentate gyrus neuronal density in experimental 1 and 2. Results are expressed as Mean \pm SD of 8 rats and data were analyzed by one-way ANOVA followed by Tukey-kramer multiple comparisons test.

Discussion

Our findings demonstrate that endogenous estrogen plays an important role in the maintenance and survival of the cell. The data show that there is a remarkable change in the number of neurons in different groups. Endogenous estrogen in control animals resulted in a remarkable increase in the number of intact neurons, respectively as compared to ovariectomized groups (Fig. 2). Results show that the rate of degeneration was not increased in ovariectomized groups during the experiment. It means that in the control group endogenous estrogen acts as a survival factor and suppresses the degeneration phenomena. It is also recommended that whenever in the animal gonad hormone decreased, programmed cell death occurred. In ovariectomized groups, even omitting the hormone was longer, the numerical density was not reduced more (Fig.3).

The hippocampus is especially important because it has been shown to be involved in certain aspects of learning and memory (21) the degeneration of which are primary effects of Alzheimer's. The reasons for this degeneration are many-fold. According to one study, estrogen is associated with an increase in the release of certain brain chemicals which transmit information across the space between two nerve cells, in the brain (2). Therefore, when estrogen presence decreases or disappears, neural chemicals which previously were available in high quantities are no longer largely present, causing a decrease in brain activity. A study of estrogen-treated rats tends to support this theory in that the treated rats had higher memory performances than did estrogen-deprived rats (2).

Gonadal androgens (testosterone) can act as a regulator of the expression of receptors for trophic factors, proteins critical for the maintenance of normal structure and function. The cellular effects of gonadal hormone can be grouped into genomic and non-genomic categories. Genomic effects are related to transcription and translation of genes, but non-genomic effects occur very rapidly and involve ion movements and/or initiation of signal transduction cascades (22). Testosterone might act directly through androgen pathways or indirectly via conversion to estrogen (8). There is a consensus that these

hormones increase secretion of neuromediators, stimulate formation of new synapses and activation of certain genes, responsible for production of anti-apoptotic proteins and growth factors (23). Furthermore, accumulated evidence suggests that TP manifests its effect on neuronal regeneration in the emendate post operative or pre regenerative phase by altering the cellular stress response (24).

A report argues for a previously unrecognized antioxidant cycle for estrogen derived compounds (25). Other researchers reported that exogenous administration of testosterone immediately after nerve injury impacted positively on functional recovery through actions mediated by the androgen receptors (AR) (26). They have suggest that mechanism, by which steroidal enhancement of the regenerative properties of the injured motoneurons occurs, may involve pre-existing AR, modulation of the cellular stress response and heat shock proteins. It is interesting that steroid hormones regulate ribosomal gene expression and nuclear ultra structure in target tissues (8).

Furthermore, induction of hippocampal LTP is facilitated in ovariectomized rats treated with estrogen as compared to untreated ovariectomized rats (1). Generally, the results of present study indicated that endogenous estrogen may change the media for regenerating neurons or survival the cell. It is concluded that estrogen with the protective role is clinically beneficial in the cases of neuronal death. If estrogen provides neuroprotection as seen in rats, estrogen treatment would act to save a number of patients from CNS damage.

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