Prenatal Water Deprivation Induces Apoptosis in Sexual Dimorphic Nuclei of the Brain of Male New Born Sprague-Dawley Rats

Deprivación Prenatal de Agua Induce Apoptosis en Núcleos Dimórficos Sexuales del Cerebro de Ratas Sprague-Dawley Macho Recién Nacidas

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SUMMARY: Considering the size of some nuclei and area, sex hormones control the sexual development of the brain. The sexual development of the brain can also be influenced by environmental stress. This study aimed to clear the effect of prenatal water deprivation on the development of sexual dimorphic nucleus (SDN) of the brain. In this research, pregnant rats were divided into two groups (control and treated). For the treated animals, water was removed from the ewes for 48 h at the end of third trimester of gestation (19-21 days). TUNEL staining was used for detection of apoptosis in paraffin embedded diencephalon selected sections. The ratio of apoptotic cells to non-apoptotic ones was calculated as apoptotic index. Differences of apoptotic index and serum testosterone were examined for statistical significance using Paired T-test (p<0.05). The apoptotic index was 0.0160±0.01174% for control and 0.1870±0.02541% for treated groups. The concentration of serum testosterone was 22.4±1.3 for control and 13.37±3.3 for treated groups. Prenatal water deprivation induces apoptosis in developing SDN nucleus of male rats that is derived by reducing the concentration of serum testosterone. The study shows the importance of low concentration acting testosterone for development of SDN nucleus that can be affected by environmental stress.

KEY WORDS: Apoptosis; Rat; SDN nucleus; Stress, TUNEL

INTRODUCTION

The brain develops differently in males and females under the influence of sex steroid hormones. The sex regions of the brain mainly exist in the hypothalamus where the most prominent of them is called medial pre-optic area (MPOA). Sexual dimorphic nucleus of POA (SDN-POA) circumscribed region within the medial part of the MPOA, of male rats exhibits about five fold greater in nuclear volume than females (Morris et al., 2004; Brann, 1995; Chung et al., 2000).

In humans and also in rats, the SDN-POA becomes sexually dimorphic as a result of prenatal hormone exposure and is not affected by steroidal environment in adulthood. Indeed, during intrauterine development, the SDN-POA is affected by testosterone according to the involvement of androgen receptor (ARs) neurons among this nucleus. Generally, testosterone binds with ARs in the cytoplasm, forms complexes and travels into cell nucleus, where it then binds with DNA to stimulate gene expression of apoptosis regulators (Mooradian et al., 1987; Arai et al., 1996; Lund et al., 2000).

It has been reported that neonatal treatment with testosterone to females was exerted on size of the SDN-POA and resulted in almost the same size between females and untreated males. Conversely, when newborn female rats are injected with tamoxifen (an antiestrogen) the volume of their SDN-POA is decreased (Vancutsem & Roessler, 1997).
Prenatal stress or castration of male rats on the first day of life reduces the volume of this nucleus permanently (Kerchner & Ward, 1992; Anderson et al., 1985). Moreover, stress induced by crowding during the final third of pregnancy, reduced sexual behavior and fertility in mice (Crump & Chevins, 1989). It also observed that prenatal stress and alcohol did not reduce the level of testosterone hormone in newborn mice but pregnancy, reduced sexual behavior and fertility in mice but stress induced by crowding during the final third of (Kerchner & Ward, 1992; Anderson et al., 1985).

The present study was concerned with prenatal water deprivation on the development of SDN-POA nucleus of male newborn rats.

MATERIAL AND METHOD

Twenty inbred female Sprague-Dawley rats aged 6–8 weeks (250 ± 5 g) were obtained from a closed bred colony at Islamic Azad University. The animals were maintained in an air-conditioned animal house at a temperature of 23-25 °C, relative humidity at ~50 % and photo-cycle of 12:12 h under ventilation at 38°C. The animals were provided with standard diet pellets and water ad libitum. Rats were maintained in these facilities for one week before the experiment. All experiments were carried out according to the Guidelines of the Animal Care and Use Committee of our University.

Experimental procedure. Following mating, beginning of pregnancy was determined by daily evaluation of sperm presence in vaginal smear by light microscopy analysis (ZEISS, magnification x100). Pregnant rats were divided randomly into two control and treated groups (n= 10). For the control animals, they were provided with both food and water. For the treated animals, water was removed from the ewes for 48 h at the end of third trimester of gestation (19-21 days). Following delivery, the newborns were perfused via the carotid artery by a 16-gauge needle with 0.1 M phosphate buffered saline (PBS) followed by 4% paraformaldehyde (PFA) in 0.1 M phosphate buffer under anesthesia.

The perfusion period was about 6–8 min, and the newborns were decapitated during perfusion. The brain was removed immediately following perfusion. Post fixation was performed in the same PFA solution for 12 h, after which the brain was placed in 20% sucrose in 0.01 M phosphate overnight.

The paraffin blocks were prepared following routine processing protocol of paraffin preparation sampling method. Seven micrometer coronal sections were cut through the newborn brains. Diencephalic sections selected for TUNEL staining method.

TUNEL Assay. Brain cell apoptosis was assessed by terminal deoxynucleotidyl transferase (TdT)-mediated deoxy-UTP nick end labeling (TUNEL) assay using ApopTag alkaline phosphatase in situ Apoptosis Detection kit (Roche; 11684809910), according to the manufacturer’s instructions. Following TUNEL staining, sections were counterstained with Harris Hematoxylin and mounted under glass cover slips. The sections were examined and scored under a light microscope (Nikon; YS100) equipped with a digital camera (Motic) (100X). Apoptotic index was the ratio of TUNEL-positive cells to negative ones and for this purpose sections from 10 animals per group in at least 5 random fields per brain were analyzed. As negative controls, a section from each animal was processed, but terminal transferase was omitted from the TdT labeling buffer and for positive controls DNAse was added on the sections (Hsu et al., 2001).

Serum testosterone level. The serum samples were extracted with diethyl ether and allowed to freeze in a dry ice-ethanol mixture. The ether was decanted into another tube and dried under ventilation at 38°C. The dried residue was dissolved in 0.01 M PBS (pH 7.4) containing 0.1% gelatin. Aliquots of the PBS-gelatin—dissolved steroids were used for RIA as previously described without further chromatographic separation of testosterone (Yu et al., 1988; Hsu et al.). The sensitivity of the testosterone assay was 1.3pg/assay tube and the intra-assay and inter-assay coefficients of variation were 2.95 and 10.3%, respectively.

Statistical Analysis. All data were analyzed using Sigma Stat (SPSS Inc, Chicago, Illinois). Apoptotic indices and the concentration of blood testosterone are presented as means ± SEM. Differences between groups were examined for statistical significance using Paired T-test (p< 0.05).

RESULTS

A clear difference between control and treated male newborn rats in the number of the cells of SDN-POA was confirmed. Indeed, the numbers of stained cells of SDN-POA by H&E were higher in control than the treated group (Figs. 1A, B).

The apoptotic cells exhibited bright labeling of fragmented nuclear DNA by TUNEL staining (Figs. 2A and B). The apoptotic index was .0160 ± .01174% for treated and .01174% for control and .1870 ± .02541% for treated groups (Fig. 2A). The concentration of serum testosterone was 22.4 ± 1.3 for control and 13.37 ± 3.3 for treated groups (Fig. 2B).
The mechanisms by which morphological sex differences arise in the central nervous system are not completely understood. Experimental and human data suggest that stresses, of either maternal or fetal origin, act in the developing embryo (Vrekoussis et al., 2010).

On female reproduction, the stress system suppresses the hypothalamic-pituitary-ovarian axis at the hypothalamic, pituitary, ovarian, and uterine levels. These include preterm birth of the offspring, low birth weight, and the development of adult diseases ranging from the metabolic syndrome to several neurodevelopmental disorders (Vrekoussis et al.). In line with these studies, our data suggest that water deprivation has an effect on the development of SDN-POA nucleus of newborn male rats.

The formation of sexually dimorphic nuclei involves the control of cell number in such nuclei by neurogenesis, neuron migration, apoptosis and differentiation (Tobet, 2002; Forger, 2006). Furthermore, previous studies showed that prenatal stress reduces the size of sexually dimorphic nucleus (Anderson et al., 1985; Rhees et al., 1999; Kerchner & Ward).

The study in parallel with this work confirmed that water deprivation affect on the development of sexually dimorphic nucleus of males by inducing apoptosis.

Moreover, our data complete these studies by showing that the reduction of cell number is caused by apoptosis but not blocking neurogenesis, neuron migration or inducing differentiation.

It seems that the role of testosterone for development of sexually dimorphic nuclei is very important as the study showed the role of testosterone declining by prenatal stress. This data confirms Anderson et al. (1986) but is in contrast with Ward et al.

The difference in the volume of SDN-POA nucleus has been attributed by the researchers to the prevention of neurons from apoptosis developed by the influences of circulatory androgen after being converted to estrogen (Davis et al., 1996). In line with this idea, testosterone was exposed prenatally in rats that prevent apoptosis of SDN-POA nucleus of females but not in the males (Yang et al., 2004). This data confirmed that testosterone in low concentration controls the development of sexually dimorphic nuclei of males and high concentration for the females. Here we suggest that when the rats were exposed to prenatal water
deprivation, the concentration of testosterone reduced from the minimum needs for masculinization of the nucleus.

Summing up, prenatal water deprivation reduces the concentration of plasma testosterone which leads to reduction of the size of sexually dimorphic nucleus. The reduction in the size of the nucleus is derived by induction of apoptosis.

ACKNOWLEDGEMENTS. We acknowledge the staff of Fertility and Infertility research center of Kermanshah.

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Received: 21-12-2010
Accepted: 09-03-2011


