Effects of hyperthyroidism on expression of vascular endothelial growth factor (VEGF) and apoptosis in fetal adrenal glands

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Abstract

This study investigated the expression of vascular endothelial growth factor (VEGF), vascular density, and apoptosis in fetal rat adrenal glands with hyperthyroidism in late gestation. Twelve mature female Wistar albino rats with the same biological and physiological features were used for this study. Rats were divided into two groups: control and hyperthyroidism. Hyperthyroidism was induced by daily subcutaneous injections of L-thyroxine (250 µg/kg) before pregnancy for 21 days and during pregnancy. Rats in the control and hyperthyroidism groups were caged according to the number of male rats. Zero day of pregnancy (Day 0) was indicated when the animals were observed to have microscopic sperm in vaginal smears. Pregnant rats were sacrificed on the 20th day of pregnancy; blood from each animal was collected to determine the concentrations of maternal adrenocorticotropic hormone and thyroxine. Rat fetuses were then quickly removed from the uterus, and the adrenal glands of the fetuses were dissected. VEGF expression, vascular endothelial growth factor (VEGF), and CD31 in fetal rat adrenal glands were measured using immunohistochemistry. The results showed that hyperthyroidism significantly stimulates metabolism and causes sudden weight loss, a rapid or irregular heartbeat, sweating, and irritability. Graves’ disease is prevalent in women of gestational age and affects approximately 0.2% of pregnant women. However, neonatal hyperthyroidism has been occurring in roughly 1% of neonatal born to mothers with Graves’ disease.

In the early phases of organ development, some fetal organs are dissimilar in structural or functional organization seen in adult organs. The fetal adrenal systems of humans and other mammalians are structurally different from those of adult structures. The human fetal adrenal gland is composed of the definitive zone, the transitional zone, and the fetal zone, from the outer to the inner part. Conversely, the rat fetal adrenal gland is almost established around the time of birth.

Ross et al. demonstrated that the rat adrenal cortex on gestational day 18 seems to show histological zonation, from the outer to the inner part, namely, zona glomerulosa, zona fasciculata, and zona reticularis. However, Mitani et al. showed the presence of a fourth zone between zona glomerulosa and zona fasciculata when performing immunohistochemical staining of an adult rat adrenal gland. This zone is called the undifferentiated zone and contains stem cells for the adrenal cortex.

In humans, the fetal adrenal vasculature is established by the eighth week of gestation when the adrenal gland is supplied by arteries from the descending aorta, and the capillaries within the organ form a continuum with a common circulation. One of the most important angiogenic factors that may be involved in the regulation of adrenal vascularization is the vascular endothelial growth factor (VEGF), the actions of which are limited to the vascular endothelial cells. VEGF is a potent regulator of blood vessel formation, and it enhances vascular permeability.

CD31 [platelet endothelial cell adhesion molecule (PECAM-1)] is found on the surface of platelets, Kupffer cells, T/NK cells, and megakaryocytes, and it makes up a large portion of endothelial cell intercellular junctions. Under normal circumstances, CD31 is observed at high levels in the vascular endothelium.

Angiogenesis and/or vasulogencis is essential for a variety of embryonic processes, including tissue development, growth, and differentiation. Important factors act on fetal adrenal growth through intra-adrenal growth factors, and these factors include fibroblast growth factor, insulin-like growth factor II, and epidermal growth factor. Previous studies have suggested that extracellular components (fibrinogen, collagen IV) are also important in coordinating proliferation, migration, and differentiation.

In the present study, we investigated the expression of VEGF and CD31 in fetal rat adrenal glands with hyperthyroidism in late gestation (Day 20). We also examined the changes in apoptosis in the corticomedullary region of fetal adrenal glands.

Materials and Methods

Animals

Twelve mature female Wistar albino rats with the same biological and physiological features were used in this study. These rats were bred in the Research Department of Experimental Animals in Trakya University, and their weights varied between 200-230 g. The study was approved by the Institutional Animal Ethical Committee of Trakya University, Edirne, Turkey (permission number: TUHDYEK-2015/22). In this experiment, all subjects were fed with rat pellet feed (Purina) containing 21% pure protein and were given drinking water in an optimum laboratory atmosphere (22±1°C, 12-h light/dark cycle).

Experimental protocol

Female rats were randomly divided into two groups: control and hyperthyroidism. Rats from the hyperthyroidism group were injected subcutaneously (sc) with 250 µg/kg/day of thyroxin hormone (in normal saline; L-thyroxine, Sigma, St. Louis, MO, USA) for 21 days. The control group received daily sc injections of a saline vehicle. After thyroxin administration for 21 days, vaginal smears were taken from
each animal, and animals in the estrus phase were placed in a cage attached to another cage with a male rat. All females mated with males (ratio 1:1). The morning when spermatozoa were present, the vaginal smear was designated as day zero of pregnancy. During pregnancy, L-thyroxine was injected once every 48 h, and the drug was stopped on the 20th day of pregnancy. Pregnant rats were anesthetized intraperitoneally with xylazine-ketamine and sacrificed on the 20th day of pregnancy. Blood from each animal was collected, and serum was separated to determine the concentrations of maternal ACTH and T4. Rat fetuses were then quickly removed from the uterus, and the adrenal glands of fetuses were dissected. In this study, 24 fetuses were examined for each group, and the fetuses were collected from six different mothers. The rat fetal adrenal glands were fixed with buffered formaldehyde solution for 24 h. The fetal adrenal gland samples were dehydrated and embedded in paraffin according to standard histological procedures. Serial cross-sections of 5 µm were prepared from each rat fetal adrenal gland. Fetal placental weights were also measured, and the number of fetuses was counted and recorded.

Chemical immunoassay of ACTH and T4 levels

Serum concentration levels of maternal adrenocorticotropic (ACTH) and thyroxine (T4) in both groups were measured using 100 µL samples of serum with commercially available chemical immunoassay kits (Siemens, Immulite 2000 analyzer and Advia Centaur XP, respectively).

Immunohistochemical examination

The avidin–biotin complex method was used for immunohistochemical staining as previously reported. Sections of 5 µm thick were examined by light microscopy on the preparations obtained from each rat. The paraffin sections were incubated with specific monoclonal anti-VEGF antibodies (1:100; ab6154, Abcam, Cambridge, MA, USA). In the negative controls, the primary antibodies were replaced with phosphate-buffered saline (PBS). Fetal adrenal tissue sections were examined under light microscopy (400X). Cells with red cytoplasmic staining were considered positive, and the number of VEGF positive cells were counted in random high-power sections using a light microscope (Olympus BX51, Japan) and incorporating a software analysis system (Argentin Kameram, ver. 2.11.5.1, Istanbul, Turkey). In fetal adrenal glands, the number of VEGF positive cells were counted in 10 random (each rat) sections was counted. All scores were converted to the number of VEGF positive cells per unit area (mm²).

Figure 1. Effects of hyperthyroidism on maternal ACTH (a) and T4 (b) levels. *P<0.05 compared with the control group.

Figure 2. Expression of fetal rat adrenal VEGF in the zona glomerulosa and zona fasciculata. VEGF expression was more prominent in the zona fasciculata than in other zones in both the control and hyperthyroidism groups. a) Control group. b) Control group. c) Hyperthyroidism group. d) Hyperthyroidism group. e) Analysis of VEGF positive cells in the fetal rat adrenal gland. VEGF positive cell number in the hyperthyroidism group was significantly increased compared with the control group. GZ, zona glomerulosa; FZ, zona fasciculata; arrows, VEGF positive cells. *P<0.05 compared with the control group. Scale bars: a,c) 100 µm; b,d) 50 µm.
Microvessel density

Fetal adrenal sections were stained with the anti-CD31 (Abcam, 1:100; BS-0195R- Bios, Woburn, MA, USA) antibodies and then analyzed. In the negative controls, the primary antibodies were replaced with PBS. The number of microvessels in the adrenal cortex was counted for five randomly selected fields in each section at 400x magnification. The value was defined in area mm² for the microvessel density.

TUNEL assay

The terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL) method, which detects the fragmentation of DNA in the nucleus during apoptotic cell death in situ, was employed using an apoptosis detection kit (Apop Tag plus Peroxidase in Situ Apoptosis Detection Kit S7101, Millipore, Billerica, MA, USA), as previously reported. The distribution of TUNEL-positive cells was conducted in the same manner as the anti-VEGF.

Statistical analysis

The statistical analyses were determined by one-way analysis of variance (ANOVA) using the SPSS version 19.0 (Chicago, IL, USA) software. Groups were compared using ANOVA, one-way analysis of variance (ANOVA) using Tukey's test for post hoc analysis or by a non-parametric Kruskal-Wallis test whenever appropriate. Significance was accepted at P<0.05.

Results

Serum ACTH and T₄ levels

Both ACTH and T₄ levels tended to increase in the hyperthyroidism maternal serum samples compared with the control maternal serum samples (Figure 1 a,b; P<0.05). Fetal, placental weights, and number of fetuses were not significantly different between the hyperthyroidism and control rat groups (data not shown).

VEGF expression

The adrenal cortex expression of VEGF was examined by immunohistochemical staining methods. The VEGF positive cell numbers are summarized in Figure 2. The zona fasciculata contained a greater number of VEGF positive cells than the zona glomerulosa in all control and hyperthyroidism maternal offspring fetuses. In addition, the number of VEGF positive cells was higher in the outer zona fasciculata than in the inner zona fasciculata areas. The number of VEGF positive cells in the hyperthyroidism fetal group was significantly increased compared with the control fetal adrenal group (Figure 2e).

Microvessel density

The vessel density values in the hyperthyroidism group were significantly increased compared with the control fetal adrenal group. (Figure 3; P<0.05). Microvessel density was more intensive in the zona glomerulosa than in the zona fasciculata (Figure 3).

TUNEL assay evaluations

The TUNEL positive cell count is summarized for both groups in Figure 4. A statistically significant increase in the number of TUNEL positive cells was detected in the hyperthyroidism group (P<0.05) compared with the control group. Furthermore, the TUNEL positive cell density was more intense in the zona fasciculata than in the zona glomerulosa of fetal adrenal glands (Figure 4).

Discussion

During embryonal and fetal development, angiogenesis is concomitant with proliferation, and microvessels provide oxygen, nutrients, and growth factors to tissue cells. The significance of angiogenesis in mammalian adrenal gland and activity shows the gland as one of the most highly vascularized organs in the mammalian fetus (e.g., human and rhesus). The steroid products of fetal adrenal glands are considered essential for fetal tissue maturation.

Adrenal and thyroid secretions play an important role in fetal development, protection of homeostasis throughout fetal term, and producing the steroid precursor dehydroepiandrosterone, which is converted to estrogen by the placenta. In the present study, maternal serum ACTH and free T₄ levels...
were significantly increased in the hyperthyroidism group compared with the control group. ACTH secreted by the fetal pituitary plays an important role in the prominent development of the human fetal zone. Additionally, ACTH is the primary regulator of adult adrenal cortex trophicity. Human and other mammalian species may have developmental failures related to adrenal development-induced pathological conditions. Fetal hyperthyroidism may cause fetal tachycardia, increase in thyroid gland size, and intrauterine growth retardation. In this study, hyperthyroidism (induced with L-thyroxine 250 µg/kg) showed a modulatory effect on VEGF and CD31 (PECAM-1) expressions in the fetal adrenal glands. However, hyperthyroidism did not change the body weight of offspring, placental weights, and the number of fetuses (data not shown). Johnson et al. showed that adrenal weights significantly increased at 7 and 60 days in hyperthyroid rats compared with euthyroid rats.

In previous studies, VEGF expression has been identified in human fetal adrenal glands and rhesus monkey adrenal glands. The expression of VEGF in the midgestation human fetal adrenal gland has been demonstrated in the definitive and fetal zones.27 We demonstrated for the first time that VEGF-positive cells were denser in the periphery of the rat fetal zona fasciculata (glomerulosa-fasciculata border) in late gestation (days 20). By contrast, the number of VEGF positive cells was less in the zona glomerulosa and the central region of the fasciculata. According to a previous study, VEGF-A was expressed strongly in the glomerulosa and fasciculata layer cells in adult bovine adrenal glands.

The human fetal adrenal gland is highly vascularized at midgestation, with a distinct patterned vascular network in the inner fetal and outer definitive zones. VEGF-positive cell density was greater in the fetal zone than in the definitive zone. Shifren et al. suggested that after the administration of ACTH, VEGF expression increased in human fetal adrenal glands. The nourishing effect of ACTH could be mediated through the vasculature. This argument is supported by the fact that ACTH stimulates the expression and secretion of VEGF in primary cultures of adrenal cortical cells.

In the literature, no study has examined the microvessel density of rat fetal adrenal gland in experimentally induced hyperthyroidism. We investigated microvessel density in fetal adrenal glands in both the zona glomerulosa and the zona fasciculata using immunohistochemistry. In this study, vascular density in the hyperthyroidism group was significantly increased in fetal adrenal tissue compared with normal healthy fetus. Terada suggested that the human fetal adrenal cortex was negative for CD31 at week 7 of gestation.24 In the current study, microvessel density immunohistochemistry revealed that the vessels were denser in the zona glomerulosa than in the zona fasciculata. We observed that hyperthyroidism increased vascular density of the fetal adrenal glands in rats. This condition may be associated with the increased maternal blood levels of ACTH, which increases vasularization. Muench et al. suggested that human fetal adrenal cells were 6% CD31 positive in all cell populations, positive cells and vessels were observed in the capsule, and cells were dispersed in the definitive zone.2 A previous study found that capillary activities were not detected on day 16 of gestation in the rat fetal adrenal gland. However, the same study showed that the capillaries were detected immunohistochemically in the rat fetal adrenal gland on gestational days 19 and 20. Apoptosis has a critical role in the development of adrenal glands, and the disruption of developmental apoptosis may produce conditions such as hyperthyroidism and steroidogenic hormone secretion disorders. In the present study, fetal adrenal glands of the hyperthyroidism group showed higher numbers of apoptotic cells than the control group glands. In both groups, the TUNEL positive cells were particularly found in the inner region of the zona fasciculata. Spencer et al. suggested that human fetal adrenal apoptotic cells were detected at low levels in the definitive zone. In the present study, TUNEL positive cells, regardless of the group, were rarely observed in the zona glomerulosa of fetal rat adrenal glands. The number of apoptotic cells was low in the control fetal adrenal glands at day 20 of gestation, but it increased markedly in the fetal adrenal tissue of the hyperthyroidism rat group. These results are similar to those of previous studies. For example, Jirasek detected cellular apoptosis in human fetal adrenal glands, mainly in the central portions of the fetal zone. Mitani et al. suggested that apoptosis was scattered during the late gestational period in the rat fetal adrenal gland.

Previous studies have suggested that fetal pituitary ACTH is required for normal adrenal growth. However, fetal zone regression and the postnatal involution of the adrenal gland do not coincide with an apparent decrease in ACTH exposure. With regard to fetal zone regression and cell apoptosis, the current thinking is that other factors, such as the peptides activin A and TGFβ, can stimulate apoptosis in several cell types. We demonstrated changes in the distribution of VEGF expression, microvessel density, and apoptosis in the rat fetal adrenal gland after L-thyroxine-induced experimental hyperthyroidism. An abundance of VEGF protein expression was found on the outer side of the zona fasciculata in the rat fetal adrenal gland in the offspring of hyperthyroid mothers. Microvessels were dense in the capsule and in the zona glomerulosa in the hyperthyroidism group compared with the control group. These issues may be clarified by a detailed investigation of the development of adrenal gland cells during fetal growth.

Figure 4. Representative photographs of the TUNEL assay. a) Control group. b) Hyperthyroidism group. c) The number of TUNEL positive cells per mm² was counted in the zona glomerulosa and zona fasciculata of each group. Arrows, TUNEL positive cells. *P<0.05 compared with the control group. Scale bars: 50 µm.
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