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**ТHE EFFECT OF** **INTERMITTENT COLD EXPOSURE**

**AND BROWN ADIPOSE TISSUE ON THE STATE**

**OF ADIPOCYTOKINES INRATS**

**WITH DEHYDROEPIANDROSTENEDIONE –INDUCEDPOLYCYSTIC OVARY SYNDROME**

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**Introduction.** Polycystic ovary syndrome (PCOS) is one of the most common diseases in women of reproductive age and includes cystic ovarian degeneration, hyperandrogenism and chronic anovulation [1]. In addition, it is associated with a high risk of developing insulin resistance, obesity, dyslipidemia and cardiovascular disease. Because no etiological factor is able to fully cover all clinical signs, the pathogenesis of PCOS is largely unknown. PCOS has been shown to be associated with the secretion of adinocytokines (leptin and adiponectin) [2]. Leptin activates specific leptin receptors. An important regulator of leptin secretion is hyperinsulinemia (GI). Adipocytes produce leptin in response to increased insulin levels. Adiponectin is secreted by white adipose tissue and affects the release of gonadotropins and can be regulated by insulin and serve as a marker of insulin resistance [3]. One of the possible factors in the pathogenesis of PCOS, at present, is called dysfunction of adipose tissue (even in the absence of obesity), which is primarily associat-ed with insulin resistance. There are two main types of adipose tissue in hu-mans and mammals: white adipose tissue, which stores energy in the body, and brown adipose tissue (BAT), whose main function is thermogenesis and maintaining body temperature under cooling [4].The aim of the study.To determine the changes in adiponectin and lep-tin in rats during experimental PCOS before and after intermittent cold exposure and the role of brown adipose tissue in restoring the structure and function of the ovaries on the background of insulin resistance.

**Material and methods**. The study was performed on 32 white female rats of the population WAG / GSto, aged 5-6 months, body weight, before the study, 240 ± 14,5 grams. To model PCOS, rats were divided into 2 groups: main (25 rats) and control (7 rats). The control group consisted of healthy animals, and was selected to compare the results with the main group. The experiments were performed in accordance with the «General Principles of Animal Experiments», approved by the I National Congress of Bioethics (September 20, 2001, Kyiv, Ukraine) and agreed with the provi-sions of the Convention on the Protection of Vertebrate Animals for experi-mental and other scientific purposes.Modeling of PCOS in rats was performed using exogenous dehydroepi-androstenedione (DHEA). In this case, the mechanism of insulin resistance is receptor. Due to the increase in the area of adipocytes there is a decrease in the number of insulin receptors per unit area. The level of sex hormones, insulin, antimullerian hormone (AMG) in rats was determined by ECLIA immunoas-say using automatic analyzers and reagents from Roche Diagnostics (Germany) and enzyme-linked immunosorbent assay of Immunotech kits (Czech Republic). The concentration of adipocytokines in the serum was de-termined by enzyme-linked immunosorbent assay using the sandwich method (ELISA) with a kit manufactured by Diagnostics Biochem Canada Inc.Intermittent cold exposure (ICE) was carried out by keeping the animals for 4 hours in a chamber in which thelight regime and a temperature of + 4C. The remaining 20 h of the animal were under normal conditions of ambient temperature and light regime. Rats were cooled daily for 20 days.Statistical processing of the obtained data was performed using the soft-ware package Statistica10.0 (StatSoftInc., USA).

**Results.** Hormonal examination of rats with experimental PCOS showed a significant increase in peripheral blood luteinizing hormone (LH), ovarian androgens, a tendency to decrease the level of estrodiol (ES) and insufficient content of folliculo-stimulating hormone(FSH) and progesterone (р<0,01). It was determined that adiponectin does not affect the concentration of FSH, but reduces the secretion of LH.In our study, we found that the level of adiponectin in the serum of rats of the control group increases, in accordance with the increase in the concen-tration of antimullerian hormone (AMG) and LH.The concentration of leptin in the blood increased proportionally with the degree of obesity and insulinemia. In particular, in rats of the main group with experimental PCOS, the insulin content was almost 2,5 times higher than in rats of the control group. Decreased levels of adiponectin and increased leptin concentrations in rats of the main group compared with the control group, confirm the dysfunction of adipose tissue of rats with PCOS.Morphometric analysis allowed a quantitative assessment of changes in the cytomorphological profile of adipocytes of the BAT under intermittent cold exposure. A significant decrease in the number of adipocytes in the BAT of animals compared with the control. Visually, BAT acquired a denser packaging and a pronounced darkbrown color. There was a significant de-crease in the area of adipocytes in rats of the experimental group with PCOS, in which in adipocytes of the same type there was asignificant decrease in the diameter of lipid vacuoles after the animals were subjected to ICE. Summarizing the results, it can be noted that under the influence of cold in BAT animals are adaptive reactions aimed at mobilizing lipid reserves to generate heat. This is expressed in a decrease in the number of adipocytes, as well as the area of cells and the diameter of lipid droplets in them.Our results coincide with those of other authors who observed similar morphological changes in BAT in rats under cold exposure. With the intro-duction of DHEA, changes are observed, which can also be described as the activation of BAT in the preservation of the morpho-functional state of the ovaries in experimental PCOS. This was expressed in the fact that the trans-plantation of BAT in rats on the background of the introduction of DHEA decreased the formation of cysts in the ovaries, while in the case of BAT transplantation, ovarian cysts are formed in approximately 85% of animals.This may indicate a protective effect ofactivated BAT on the regulation of follicle maturation. It is known that rats injected with exogenous DHEA are characterized by a significant decrease in energy metabolism. The mech-anism of influence of BAT on DHEA-induced cyst formation in the ovaries isassociated with the secretion of adipocytokines (leptin, adiponectin) and other systemic regulators of metabolism.The level of adiponectin decreased in rats with experimental PCOS on the background of induced cyst formation in the ovaries, possibly due to other systemic regulators of metabolism. ICEleads to increased levels of adipocytokines, which may indirectly affect the secretion of reproductive hormones and prevent the development of cystic changes in the ovaries.

**Conclusions.** Experimental polycysticovaries syndrometo degenerative cystic changes in the ovaries of rats, which contributes to the secretion of gonadotropic hormones and indicators of adiponectin and leptin associated with insulin resistance.Intermittent cold exposures and brown adipose tissue transplantation help to restore the structure and function of ovarian tissue, normalizetion the synthesis of adipocytokines and prevent the development of polycystic ovary syndrome

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