

A Study of Prolactin, Thyroid Stimulating Hormones, Malondialdehyde and Ceruloplasmin Levels in Infertile Women, in Thi-Qar Governorate/Iraq

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ABSTRACT

OBJECTIVES: The study designed to investigate the level of prolactin (PRL), thyroid-stimulating hormones, Malondialdehyde (MDA) and ceruloplasmin (Cp) levelsof the groups of infertile and fertile females in Nasiriyah province, southern Iraq.

METHODOLOGY: Blood samples were collected from two groups of females (females with infertility and fertile women) with a total number of 99 infertile women and 70 fertile females at Bent Al-Huda maternity and children teaching hospital in Thi-Qar governorate. Autoanalyzercobas e411 instrument was used to estimate of prolactin, Thyroid-stimulating hormones and Ceruloplasmin levels in sera of patients and control group using Elecsys and cobas immunoassay analyzers. The MDA level of the serum was measured by the modified method of Fong et al. (1973). The concentration of plasma MDA was computed according to the equation of coefficient of MDA equal to $1.56 \times 10^3 \text{ mol}^{-1} \cdot \text{cm}^{-1}$.

RESULTS: Prolactin and TSH levels were significantly increased ($p < 0.0001$) in infertile females as compared to fertile women. A similar pattern was observed to concentrations of MDA and Ceruloplasmin were significantly increased ($p < 0.0001$) in patients when compared to control group.

CONCLUSION:The results of this study indicate that oxidative stress does occur in infertile females. Patients with infertility showed are higher significant rise in serum Prolactin hormone and thyroid-stimulating hormone levels in infertility patient group. All the patients with a high Prolactin level should have TSH estimation.

INTRODUCTION:

Infertility is the inability of a couple to achieve pregnancy over an average period of one year (in a woman under 35 years of age) or 6 months (in a woman above 35 years of age) despite adequacy¹. Infertility may also be

referred to as the inability to carry a pregnancy to the delivery of a live baby. Infertility can be due to the woman, the man, or both; primary or secondary. Ovulation problems are the most common causes of infertility and can arise as a result of a defect in hypothalamus, pituitary or ovary¹.

Hypothalamic disorder is most commonly due to weight loss, eating disorder, excessive exercise and hypothalamic lesions. Pituitary disorders include hyperprolactinemia, thyroid disease, Cushing disease and Sheehan syndrome². Ovarian dysfunctions are most commonly due to polycystic ovarian syndrome (POS), less common aetiology is premature ovarian failure³.

Hyperprolactinemia is a common problem in reproductive dysfunction affecting about one third of infertile women. It has been suggested that hypogonadism seen in hyperprolactinemic women is due to circulating levels of prolactin interfering with the action of the gonadotrophins at the ovarian level and impaired gonadal steroid secretion. Gonadotrophins, in turn, alters positive feedback effects on the hypothalamic and pituitary levels. This leads to lack of gonadotrophin cyclicity and infertility. Prolactin can inhibit the follicular estradiol production and this results in infertility⁴.

TSH stimulates the thyroid gland to secrete the hormones T4 and T3. The production of TSH is controlled by a thyrotropin releasing hormone (TRH), which is manufactured in the hypothalamus and transported to the anterior pituitary gland via the superior hypophyseal artery, where it increases TSH production and release. A high level of TSH stimulates prolactin secretion and causes ovulatory dysfunction and leading to infertility⁵. When cells use oxygen to generate energy, free radicals are created as a consequence of ATP (adenosine triphosphate) production by the mitochondria. These by-products are generally reactive oxygen species (ROS) as well as reactive nitrogen species (RNS) that result from the cellular redox process⁶.

These species play a dual role as both toxic and beneficial compounds. At low or moderate levels, ROS and RNS exert beneficial effects on cellular responses and immune function. At high concentrations, they generate oxidative stress, a deleterious process that can damage all cell structures⁷. Many human studies reported the presence of ROS in the female reproductive tract and their role in its physiological functions such as oocyte maturation, ovarian steroidogenesis and corpus maturation⁸. Also ROS may be produced during embryo metabolism and from its surroundings⁹.

Nitric oxide (NO) is synthesized during the enzymatic conversion of L-arginine to L-citrulline by nitric oxide synthase (NOS). It is a highly reactive free radical, damages proteins, carbohydrates, nucleotides and lipids results in cell and tissue damage. The two common examples of reactive nitrogen species are nitric oxide (NO) and nitrogen dioxide. NO is produced by the enzyme NO synthase¹⁰. Inducible NO synthase (iNOS) is present in monocytes and macrophages and produces a large amount of NO. During the follicular development in the ovaries, NO synthase is expressed on the surface of the oocyte⁹. Antioxidants are molecules that can prevent the oxidation of other molecule. Oxidation is a chemical reaction that transfers an electron from a molecule to an oxidizing agent. There are two types of antioxidants: enzymatic and non-enzymatic. These two types of antioxidant system are working together to ameliorate any harmful effects of oxidant in the cell, both of the enzymatic and nonenzymatic antioxidants detoxify ROS in the intracellular and extracellular environments¹¹. Many human studies reported the presence of ROS in the female reproductive tract

and their role in its physiological functions such as oocyte maturation, ovarian steroidogenesis and corpus maturation⁸.

Ceruloplasmin is a ferroxidase enzyme that in humans is encoded by the CP gene¹². Ceruloplasmin binds copper; appears to be more important as a copper storage pool than as a transport protein; integrates iron and copper homeostasis¹³. In addition, Cp is an effective antioxidant because of its ability to oxidize highly toxic ferrous iron to the relatively nontoxic ferric form and thus help prevent oxidative damage to proteins, lipids and DNA¹⁴. Malondialdehyde is an end product of lipid peroxidation (LPO) which is a process where reactive oxygen species degrade polyunsaturated lipids. This compound is reactive aldehyde and is one of the many reactive electrophile species that cause toxic stress in cells and for advanced glycation end products¹⁵. The production of this aldehyde is used as a biomarker to measure the level of oxidative stress in an organism¹⁶.

MATERIALS AND METHODS:

This study was conducted as a case-control study at Bent Al-Huda maternity and children Teaching Hospital in Thi-Qar governorate. The study was carried out in the unit of the laboratory Department of Clinical Biochemistry. Informed consent was obtained verbally from all participants. A total of ninety nine women with infertility of the ages 18 – 45 years and duration of infertility more than one year who were diagnosed by gynaecology and obstetrics. To compare the results, seventy healthy age matched females with a history of at least one childbirth were also enrolled.

Five ml of blood samples from patients and control groups was collected by vein puncture using sterile disposable

syringe in plain plastic tubes. The serum was separated immediately in order to allow to clot at room temperature. The blood was centrifuged at 3000 rotor per minute (rpm) for 10 minutes and stored in plain tubes at – 200 C until used or immediately analysed.

Determination of Serum Malondialdehyde (MDA):

In this method, lipid peroxidation reacts with thiobarbituric acid (TBA) in coexisting trichloroacetic acid (TCA) to give a pink chromophore. The MDA level of the serum sample was estimated by the modified method of Fong et al., (1973)¹⁷. The level of MDA was computed according to the equation of coefficient of MDA equal to $1.56 \times 10^3 \text{ mol}^{-1} \cdot \text{cm}^{-1}$.

Estimation of serum prolactin and TSH:

Prolactin and TSH determination kits were obtained from Roche diagnostics and determined in Elecsys 2010, autoanalyzer cobas e411 instrument.

Estimation of serum Antioxidants (serum Ceruloplasmin)

Ceruloplasmin estimation kit was obtained from Roche diagnostics and determined by clinical chemistry analyzer cobas c311 in serum samples using cobas e immunoassay analyzer instrument.

Statistical Analysis

Statistical analysis was done using the software SPSS version 15.0; the results were expressed as mean \pm standard deviations (SD). One way ANOVA-test was used to compare parameters in different studied groups. P-values ($P < 0.0001$ and $P < 0.05$) were considered statistically significant.

RESULTS:

In this study, we estimated the levels of Prolactin, TSH, MDA and ceruloplasmin among infertile and fertile females. The levels of serum prolactin hormone in the infertile women's group showed a highly significant increase ($P < 0.0001$) in comparison with their control group (Table 1). The mean serum TSH levels in infertile patients were 2.70 ± 1.42 μ IU/ml against the controls who had a mean serum TSH level of 1.89 ± 1.11 μ IU /ml and both groups are statistically significantly different to each other (p value = < 0.0001) (Table 2). The serum ceruloplasmin and MDA levels were found to be significantly increased ($P < 0.0001$ and $P < 0.0001$, respectively) in patients with infertility compared to control group (Table 3 and Table 4).

Table 1: Serum Prolactin level in infertile females compared with control group.

| Parameter | Group | No. | Mean \pm SD |
|---|-----------|-----|----------------------------------|
| Prolactin Level (μIU) | Control | 70 | 10.35 \pm 6.08 |
| | Infertile | 99 | 20.29 \pm 17.09 ^{a**} |

(a) :significant between patients with infertility and control group.

** $P < 0.0001$

Table 2: Serum TSH level in infertile women compared with control group.

| Parameter | Group | No. | Mean \pm SD |
|---------------------------------------|-----------|-----|--------------------------------|
| TSH Level (μIU) | Control | 70 | 1.89 \pm 1.11 |
| | Infertile | 99 | 2.70 \pm 1.42 ^{a**} |

- Legend as in table (1)

Table 3: Level of Serum Ceruloplasmin in Infertile females and control group.

| Parameter | Group | No. | Mean \pm SD |
|----------------------------------|-----------|-----|--------------------------------|
| Ceruloplasmin Level (g/L) | Control | 70 | 0.29 \pm 0.08 |
| | Infertile | 99 | 0.40 \pm 0.09 ^{a**} |

- Legend as in table (1)

Table 4: Level of Serum Malondialdehyde in Infertile females and control group.

| Parameter | Group | No. | Mean \pm SD |
|--------------------------|-----------|-----|----------------------------------|
| MDA Level (mg/ml) | Control | 70 | 54.79 \pm 7.46 |
| | Infertile | 99 | 75.08 \pm 15.18 ^{a**} |

- Legend as in table (1)

DISCUSSION:

The failure to identify a clear cause of the infertility after a full screening of females is defined as infertility of unknown cause¹⁸. The increased function of the thyroid is likely to cause disorders in menstrual cycle and an increase or decrease in women's sexual activity. The decrease function of the gland causes a decrease in sexual activity. Also, other problems may be due to the cause of thyroid dysfunction. Disorders of the pituitary gland, on the other hand, may cause decreased function of the thyroid¹⁹.

The mean serum TSH levels in infertile patients were 2.70 ± 1.42 μ IU/ml against the controls who had a mean serum TSH level of 1.89 ± 1.11 μ IU /ml and both groups are statistically significantly different to each other. These results are in agreement with results of^{20, 21}.

Thyroid hormones have profound effects on reproduction and pregnancy. TRH is under negative feedback

control of TSH through a short negative feedback loop; that is any increase in TSH will decrease the release of TRH which in turn will inhibit the secretion of prolactin and will also normalize the TSH levels. The relationships between TSH, prolactin and female infertility are multiple and complex. Thyrotropin-releasing hormone is a potent stimulus of prolactin and the association between hypothyroidism and hyperprolactinemia is well appreciated. Hyperprolactinemia, because of a variety of causes, can reduce pulsatile Gonadotropin releasing hormone secretion and interfere with ovulation²².

Prolactin hormone shows a highly significant increase in the infertile women's group in comparison with their control group. Prolactin is one of several hormones that is produced by the pituitary gland. The most important role of prolactin is to stimulate milk production in women after the delivery of a baby²³. This study is agreement with the findings of Mohammed A. Z (2003) increased prolactin in patients with infertility in comparison with control group may be due to low oestrogen and progesterone levels in the infertile women results in a decrease in serum LH and FSH in infertile women²⁴. According to EmoKpae et al, there maybe failure of the hypothalamus or pituitary gland, which result in increased serum prolactin levels and leads to infertility²³. This study agrees with a study done by Nissreen F. A. et al. (2014) and Iqbal Naji Tawfiq (2013) suspected that the infertile women with ovulation disorder had increased prolactin due to an inadequate corpus luteal progesterone secretion^{25, 26}. Elevated levels of prolactin cause infertility because prolactin inhibits the GnRH secretion. When the GnRH secretion is low, the FSH and LH

secretions are also low and so they do not stimulate the gamete production and the gonadal steroid synthesis^{27, 28, 29}. The high level of serum ceruloplasmin in infertile women may be due to its role in the ferroxidase activity which is of greatest importance as it converts reduced (ferrous) linked with transferrin to oxidized (ferric) iron linked with ferritin. Fe+2 acts as a pro-oxidant agent because of its readiness to change from one valency state to another. In its free form, iron is one of the most effective antioxidant catalysts³⁰. There are several mechanisms which have been suggested for ceruloplasmin antioxidant activity, including the protecting the organism as a whole from within the possible ill effects caused by the release of free radical oxidation products. This suggests that the organism might respond by raising the antioxidant efficiency of plasma by elevating ceruloplasmin levels³¹. The results of this study are similar to the study of Al-Helaly and Al-Kado on infertile females in which they showed that higher levels of ceruloplasmin were in infertile women rather than in the control group³². In this study, we revealed significant highly increase in the serum MDA level in patients with infertility in comparison with control females. These results are in agreement with other study published elsewhere in the world^{33, 34, 32}. The increase in serum MDA level in patients with infertility could be due to changes in the balance system of oxidants and antioxidants. The elevated MDA levels in serum of infertile women, rather than in the control group, signifies that the oxidative damage in infertile females results from the increase of unsaturated lipid peroxidation of various cells including ova cells. The elevated MDA levels, are indicators of a state of oxidative

stress in infertile females. MDA is an indicator of lipid peroxidation and increased levels of the peroxidation products which have been associated with a variety of acute and chronic pathophysiological processes in human beings³⁵.

CONCLUSION:

The results of this study indicate that oxidative stress does occur in infertile females. Patients with infertility showed are higher significant rise in serum Prolactin hormone and thyroid-stimulating hormone levels in infertility patient group. All the patients with a high Prolactin level should have TSH estimation.

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دراسة مستويات هرمون الحليب ، هرمون محفز الغدة الدرقية، المالوندهايد والسيرولوبلازمين لدى النساء العقيمات في محافظة ذي قار / العراق

رواء عبدالمطلب محمد حسين رائد معلك حنون ساهر عبدالرضا علي

الخلاصة

دراسة أجريت لقياس مستوى هرمون الحليب ، هرمون محفز الدرقية ، مالونداي الديهايد ومستوى السيريلوبلازمين لمجموعتين من النساء العقيمات والغيرعقيمات بواقع 99 امرأة عقيمة و70 امرأة غيرعقيمة كمجموعة سيطرة للمقارنة في مستشفى بنت الهدى للولادة والأطفال التعليمي في الناصرية وأظهرت النتائج زيادة معنوية في مستويات هرمون الحليب وهرمون محفز الدرقية لدى العقيمات مقارنة بالغيرعقيمات كذلك كانت هناك زيادة معنوية بتركيز مستويات المالونداي الديهايد والسيرولوبلازمين لدى النساء العقيمات عند مقارنتها مع مجموعة السيطرة أيضا . كما ان نتائج الدراسة تشير إلى حدوث جهد تأكسدي لدى المريصات العقيمات وزيادة في مستوى الهرموني لدى النساء العقيمات