# Biochemical Study of Gonad Hormones in Sera of Iraqi Patients with Thyroid Disorder

Fatin F. Al-Kazazz\*

Mohammed M.AL-Bahadlii\*

Received 16, September, 2012 Accepted 17, December, 2012

#### **Abstract:**

The objective of this study was to evaluate the alteration in levels of gonado trophins hormones i.e., Leutizing (LH), Follicular (FSH) in sera of patients with thyroid disorders and molecular binding study of (LH, FSH) with their antibodies

The study was conducted at the specialized center for endocrinology and diabetes from January / 2009 to March / 2010. Two hundreds and twenty three Iraqi subjects, 109 patients with thyroid disorders at age range between (40-50) years and 114 healthy individuals as control group were included in this study.

The majority of patients were female with hyperthyroidism and (49.54 %) were at age range between(40 - 50) years.

The levels of hormones(LH,FSH.tri iodothyronine(T3).thyroxine(T4), thyroid stimulating hormoneTSH) were measured by using Enzyme Linked Flourscent Assay(ELFA, while residual studies used Immuno Raido Metric Assay(IRMA) method only. It was found that production of TSH was increased with about (15-20) fold in hyperthyroidism over that in control.Sixty percent of patients with hyperthyroidism were underweight BMI $\leq$ 20.0 (kg/m²) while (44.5%) of patients with hypothyroidism were overweight BMI(25.0-29.9)kg/m².

The mean level of LH and FSH had increased significantly (p<0.01) in patients with hyperthyroidism and in females more than in males compared to control group. In contrast , for hypothyroidism the level of LH& FSH were decreased significantly (p<0.01)

It can be concluded that patients with hypo and hyper thyroidism would altered their serum LH&FSH levels in female more than in male due to the effects of T3 and T4 on pituitary secretions

Key words: Gonadal Hormones, LH, FSH, Thyroid Disorder.

### **Introduction:**

Thyroid gland is the largest organ specialized for endocrine function in human body(1). It produces hormones, L-Thyroxin (T4), Tetraiodothyronine and (Triiodo-thyronine, T3) [1,2]. Thyroid disorders can be Hyperthyroidism classified to which (thyrotoxicocsis), is excessive of secretion thyroid hormone while Hypothyroidism (Myxedema) is a decrease in the production of thyroid hormones. There several other are types

hypothyroidism goiter, such Hashimotos disease [3,4].Glycoprotein hormones. including luteinizing hormone (LH), Follicle – stimulating hormone (FSH), Thyroid – stimulating hormone (TSH). They secreted from pituitary gland. LH and FSH both are subjected to feed back regulation (short feed back loops inhibition on the anterior loop of the pituitary)by ovarian and hormones testosterone & progesterone [5-8] The FSH ,LH, and **TSH** 

<sup>\*</sup> Dep. of Chemistry College of Science Al-Mustansiriya University Baghdad-

receptors are members of the super family of G- protein coupled receptors in the cell membrane. They interact with target proteins, such as adenylate cyclase c-AMP [2]. Increased levels of LH and FSH are seen in primary ovarianfailure, while the decrease in their levels are seen in secondary ovarian failure which is an indication of a pituitary or hypothalamus disorder [9]. The current study was conducted to:

1.Evaluate the levels of (LH and FSH) in sera of patients with thyroid disorder and find out whether this disease would altere or has an effect on (LH and FSH) levels then studying the correlation between the age,BMI; sex and the disease according to the Gonadotrophins hormone levels.

2.Study the molecular binding between each 125I-anti LH or125I-anti FSH antibody with its hormones (LH or FSH) in sera of thyroid disorder females patients with in low & control hormonal levels.

#### **Subjects & Methods**

Patients and Control Study Design

The present study was conducted at the specialized center for endocrinology and diabetes with period of collection from January / 2009 to March / 2010. Patients with thyroid disorders were 109, age range (12-50)years (69females & 40 males)., while the healthy subjects were 114 (77females & 37 males)as control group, age range(13-50)years.

#### **Materials and Methods:**

1- Total serum protein was determined by using Lowry *et al.*, method. <sup>[10]</sup>

- 2- ELFA & IRMA methods were used to estimate the hormones levels of (TSH, T3, T4,FSH, and LH) in sera of patients & control groups
- 3-Quantitative determination was made using ELFA (enzyme linked fluorescent assay) technique on the Vidas Instrument by using kit for LH, FSH, TSH, T3, T4 hormone. Biomerieux-(France)
- 4- The molecular binding of gonadotrophins (LH & FSH ) with its labeled antibodies(Abs) were made and studied.

#### **Results & Discussion**

Depending on the results in Table (1) patients with thyroid disorders can be classified into two groups: group A : patients with hyperthyroidism in which their mean T3 levels were significantly higher than control group (p<0.01)  $(8.727\pm16.61),(1.988\pm0.96)$ nmol\ml respectively, also the mean value of T4 showed a significant increase(p <0.01) compared to control  $(105.5 \pm 36.75)$  ,  $(87.58 \pm 18.33)$ nmole\ml respectively, while TSH level shows a significant decrease (p <0.01) (0.062  $\pm$  0.0422) compared to control group  $(1.8517 \pm 1.0097)$ nmole\ml.

The results were in agreement with previous studies. They found that hypo thyroidism can be characterized by the increase in TSH production with about (15-20) fold over that in control and T4 levels were low, while elevated T3 & T4 concentrations are indicated of hyperthyroidism. It is well known that secretion of TSH is regulated by negative feed back from serum thyroid hormones concentrations [11-14].

Table (1): Levels of TSH, T3 and T4 in sera of patients and control groups.

Group	No.	TSH Mean±Std. (μIU/ ml)	Std. Error	T3 Mean±Std. (nmol/ ml)	Std. Error	T4 Mean±Std. (nmol/ ml)	Std.	P. Value
Control	114	1. 8517±1. 0097	0. 094	1. 988 ± 0. 96	0. 09	87. 58±18. 33	1.71	
Hyper thyroidism	35	$0.062 \pm 0.0422$	0. 007	8.727 ± 16. 61	2. 8	105. $5 \pm 36.75$	6. 21	P<0.01
Hypo thyroidism	74	21. 616 ± 19. 93	2. 3	$1.416 \pm 0.78$	0. 09	58. 93 ± 29. 19	3. 39	P<0.01

The distribution of subjects according to age is illustrated in Table (2) which shows that the majority of patients (49.54 %) were at age range between (40-50) years.[15]

Table (2) Distribution of patients, controls according to age range.

Age range (Year)	Patient No.	Patient %	Control No.	Control %
10-19	10	9.18	13	11.4
20-29	11	10.09	21	18.4
30-39	34	31.19	22	19.3
40-50	54	49.54	58	50.9
Total	109	100	114	100

The effect of gender on patients of thyroid disorder is shown in Table (3). This Table reveals that most of the patients were females (57.1%) for group A hyperthyroidism and (66.2%) for group B hypothyroidism. Baskin, Surks Freal and found that hypothyroidism is more hyperthyroidism and women show a much higher risk of hypothyroid than more with greater than 5:1 ratio and eight folds in hyperthyroidism [16-18].

According to the data found at the statically department of bio and health at the Ministry of Health in Iraq (statically guide 2009), the number of patients with hypothyroidism are more than hyperthyroidism and female patients are more than male, and in the north, and middle of Iraq more than in the south, this may be explained due to the environment or kinds of diet in Iraq [19].

Female Female Male Male Total Total Groups % No. No. % No. % Group A: 20 57.1% 15 42.9% 35 33.11 Hyperthyroidism Group B: 49 66.2 % 25 33.8% 74 67.89 Hypothyroidism 69 40 Total 109 100

**Table (3) Distribution the patients according sex** 

Body mass index (BMI) is another factor that had been studied in this research. As shown in Table (4), most patients with hyperthyroidism (60%) were underweight BMI≤20.0 (kg/m²) while (44.5%) of patients with

hypothyroidism were overweight BMI 25-29.9(kg/m²). These percentages agree with Larsen &others; symptoms of hyperthyroidisum include weight loss while for hypothyroidism include weight gain[20-22].

Table (4) Distribution of patients according to body mass index (kg/m<sup>2</sup>).

$(kg \backslash m^2)$	Classes	Hypert	thyroidism	Hypothyroidism		
		No.	%	No.	%	
<20.0	under weight	21	60%	10	13.5 %	
20→24.9	health weight	6	17.1%	3	4%	
25→29.9	over weight	3	8.6%	33	44.5%	
≥30.0	obesity	5	14.3%	28	38 %	
Total		35	100%	74	100%	

In measurement the level of LH,FSH hormones in the sera of the patients under study it cleared from Table (5), the mean level of LH &FSH had increased significantly (p<0.01) in patients with hyperthyroidism and in females more than males compared to control group .

In contrast , in hypothyroidism patients, levels of LH and FSH were decreased significantly (p<0.01) and in females more than males. The data

agree with Ashok khar et al who found that T3,T4 were significantly inhibit the release of gonadotrophin in the presence of gonadotrophin releasing hormone GnRH (using rat pituitary cell calture)[23].JF Bruni, et. al studied the effect of hypo &hyper thyroidism on serum LH&FSH in both intact and castrated male & female rats; they found that hypothyroidism results in decreased release of LH and FSH in rats [24].

Groups		No.	LH mean±Std	Std. Error	FSH mean±Std	Std. Error	P. Value
Нуро	Female	49	2.907±3.365	0.48	5.392±8.056	1.15	P<0.01
thyroidism	Male	25	0.227±1.911	0.38	2.046±2.512	0.50	
Hyper	Female	20	8.11±9.615	2.15	18.016±29.158	6.5	P<0.01
thyroidism	Male	15	15.234±16.02	4.13	33.597±28.03	8.67	r<0.01
Control	Female	77	5.125±7.046	0.80	10.149±18.74	2.1	
Control	Male	58	0.6619±0.574	0.09	12.546±2.643	0.41	

Table (5): LH& FSH levels according to sex of patients &control groups

In conclusion for the current study ,the following points can be drawn:

- 1- The majority of Iraqi patients with thyroid disorder were at ages (40-50) years and in hypothyroidism more than those with hyperthyroidism, and in females more than in males.
- 2- The levels of LH and FSH were elevated for patients with hyper thyroidism while the low LH&FSH levels were found in patients with hypothyroidisum.
- 3-Most of the patients in hypothyroidism were with over weight BMI 25-29.9 Kg\m², while patients in hyperthyroidism most of them was considered to be under weight BMI<20.0

From all above, we can say that Iraqi females with hypothyroidisum would decrease releasing LH &FSH. In other word they will be infertility in the future.

#### **References:**

1. Elisei et al., 2006. Treatment with Drugs Able to Reduce Iodine Efflux Significantly Increases the Intracellular Retention Time in Thyroid Cancer Cells Stably

- Transfected with Sodium Iodide Symporter Complementary Deoxyribonucleic Acid. The Journal of Clinical Endocrinology & Metabolism June 1, 91 (6):2389-2395
- 2.Thomas M. Devlin, 2006. "Text book of Biochemistry with Clinical Correlations"6<sup>th</sup>ed. Jhon willy &sons,inc.,publication chapter 23; pp: 904-916.
- 3. H. Jack Baskin, 2002. Evaluation and treatment of hyperthyroidism and hypothyroidism. American Association of Clinical Endocrinologists, Thyroid Guidelines, Endocrine Practice 8 (6).
- 4. Brooks W. Vetinary Partner .2008. "Hypothyroidism" The Pet Health Library:457-469.
- 5. Joe M. Chehade, M.D.2009.Unusual Presentation of a Malignant Follicular Carcinoma of the Thyroid in Struma Ovarii. Academic physican; 3(3):5-7
- Brent GA, Larsen PR, Davies TF. 2008. Hypothyroidism and thyroiditis. In: Kronenberg: HM, Shlomo M, Polonsky KR, Larsen PR, eds. Williams Textbook of

- Endocrinology". 11th ed. Philadelphia, Pa: Saunders Elsevier;:chap 12.
- 7. Sehu S, Reddy K, Fleseriu M.2007 Management of pituitary, thyroid and adrenal disorders. In: Falcone T, Hurd WW, eds. Clinical reproductive medicine and surgery. New York: Elsevier;:311-34.
- 8. William , J.; Bremner ; Huhtanieni , I. ; and Amory , J.K. 2001. Pituitary gonadotropins and their disorders . In : Becker ,K. L. et al . (eds ) . Principles and Practice of Endocrinology and Metabolism Lippincott Williams and Wilkins. 3<sup>rd</sup> ed. Philadelphia ,Chapter 16: 170-176 .
- 9. Mycec, M. J.;Harry ,R. A.; and Champe ,P. C. 2009. Lippincott's Illustrated Reviews Pharmacology Wolters Kluwer company, 1<sup>st</sup> ed., London Chapter 23: 276-283.
- 10.Lowry OH. Rosebergh NJ., Farr AL. and Randall RJ.1951." Protein measurement with the Folin phenol reagent". J. Biol. Chem.", 193(1): 265-75.
- 11. Braverman ,L. E. 1996. Evaluation of thyroid status in patients with thyrotoxi- cosis .Clin . Chem . 42 (1): 174-178,
- 12. Lavado Autric, R. Auso, Garcia Velasco, J. V., Arufe Mdel c, Eschobar del Rey, F, Barbel, P. J. 2003. Early maternal hypothyroxinemia alters histogenesis and cerebral cortex cytoarchitecture of the progeny J. Clinical Investigations. 111(7):1073-1082
- 13. Grossman, M. , Weintroub, Szkudlinski, M. W. B.D., and 1997. Novel in sights into molecular mechanism of human thyrotrophic action structural. physiological, and therapeutic implications for the glycoprotein hormone family ,Endocrine Rev; (4) 18:476-501.

- 14. Stockigt. J. 2003. Assessment of thyroid function: towards an integrated laboratory-clinical approach, Clin. Biochem. Rev: 24(4): 109-122
- 15. Iham, A.A. 2005. Study of the Biochemical parameters and someTumor markers in the sera of patients with thyroid gland disorders and thyroid carcinoma M.Sc. thesis college of science, Al-Mustansiryah University.
- 16. H. Jack Baskin, M.D. Mace,
  2006. American Association of Clinical Endocrinologists
   Endocrine practice 8(6):46-65
- 17.Freal, H. R. 2008. Leptin and some biochemical parameters in goiter patients. M.Sc. thesis ,college of science ,Al Mustansirya University.
- 18. Surks, M.I, Ocampo E. 2006. Sub clinical thyroid disease. Am J. Med; 100.217-223.
- 19. Moran, L. J.; and Norman, R. J. 2002. The obese patient with infertility: a practical approach to diagnosis and treatment. Nut.Clin. Care, 5(6): 290-297
- 20.Yong Soon Park<sup>1</sup> and Jun-Su Kim. 2012. Obesity Phenotype and Coronary Heart Disease Risk as Estimated by the Framingham Risk Score J Korean Med SciMarch; 27(3): 243–249.
- 21. Landi et al., .2012.: Methods measurement of body mass index(BMI) and waist circumference. S Afr J Clin Nutr;25(2):67-72.
- 22.Larsen, P.R., Davies ,T.F., Hay ID.1998.The thyroid gland In: wilson JD, Foster DW, kronenberg HM, and larsen PR, Text book of endocrinology. W.B saunders Co.,9<sup>th</sup> ed. Philadalphia:389-515.
- 23. Ashok, Khar,T. Taverny-Bennardo and Marian Jutisz .1980.

Effect of thyroid hormones on gonadotrophin release and biosynthesis using rat pituitary cell cultures. J. endocrinology 85, DOI10.1677: 229-235.

24. J.F. Bruni, S.Marshall , J.A.Dibbet and J. Meites. 1975. Effects of

hyper-and hypothyroidism on serum LH and FSH levels in intact and gonadetomized male and female rats. Endocrinology, endocrine society, 97: 558-563.

# دراسة كيمياحياتية لهرمونات المناسل في امصال المرضى العراقيين بخلل الدرقية فاتن فاضل القزاز\*

\*الجامعة المستنصرية /كلية العلوم /قسم الكيمياء

## الخلاصة

تهدف الدراسة الحالية الى تقييم التغيرات فى مستويات هرمونات المناسل المغديةللدرقية (اللوتوزينى وهرمون الجريب)فى امصال المرضى بخلل الدرقية ودراسة الارتباط الجزيئي مابين () ومضاداتها .

الجزء العملى: اجريت الدراسة في المركز التخصصي للغددلصم والسكرى بجمع مئتان وثلاثة وعشرين عينة من امصال العراقيين ابتداءا كانون الثاني/2009 الى ادار/2010. شملت العينات 109 لمرضى بخلل الدرقية بمدى العمر ما بين (40-50)سنة و114 لاشخاص اصحاء كمجموعة سيطرة بنفس المدى من العمر اغلب المرضى كانوا نساءمصابات بارتفاع الدرقية وان (45.49%)منهن كانوا بمدى من العمر مابين (40-50) سنة. تم قياس مستويات هرمونات الثايروكسين (T4) وثايرونين ثلاثى اليود(T3) والمحفز للدرقية (TSH) و (ELFA) باستخدام تقنية الفحص التفلورى للانزيم المرتبط(ELFA) واستخدم تتقنبة الفحص الراديوى المناعى المترى لبقية الدراسات(IRMA)

النتائج: اظهرت نتائج الدراسة ارتفاع انتاج TSH بمقدار 20-30 مرة عند المصابين بارتفاع الدرقية اكثر من TSH = TSH الاصحاء ان 60%من المرضى بارتفاع الدرقية لديهم انخفاض فى الوزن TSH = TSH = TSH الاصحاء ان 60%من المرضى بانخفض الدرقية كانواعندهم زيادة فى الوزن . TSH = TS

الاستنتاج يمكن القول ان مرضى ارتفاع وانخفاض الدرقية سبعانون من تغير فى هرمونات المناسل LH&FSH لاسيما النساء اكثر من الرجال بسبب تاثيرات هرمونات الدرقية T3وT4 على افرازات الغدة النخامية.