

## EFFECTS OF LATENT TOXOPLASMOSIS ON AUTOIMMUNE THYROID DISEASES IN PREGNANT WOMEN - DUHOK CITY-KURDISTAN REGION- IRAQ

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### ABSTRACT

**Background:** Toxoplasmosis is a common zoonotic disease has a global distribution and can infect many hosts causes several clinical symptoms in humans and attack many body organs leading to hormonal and behavioral alterations in infected hosts. Latent or dormant form toxoplasmosis is the common form that can affect pregnancy course. Autoimmune thyroid disease (AITD) belongs to the known risk factors for adverse pregnancy outcomes. The aim of this study was the detection of anti-Toxoplasma antibodies and its effects on autoimmune thyroid disease among pregnant women.

**Methods:** A total of 220 pregnant women were included in the current study from August 1st 2021 to February 1st, 2022. Toxoplasma status in pregnant women was detected for seropositivity of anti-Toxoplasma IgG antibodies using ELISA technique. Free triiodothyronine (FT3), free thyroxine (FT4), Thyroid stimulating hormone (TSH) and thyroperoxidase antibodies (TPO) were assessed by ELISA. The blood parameters of examined pregnant women were measured by a Coulter counter machine.

**Results:** Overall, 95(43.2%) of the examined pregnant women had seropositivity for anti-Toxoplasma IgG antibodies and 28(12.7%) were screened positive for AITD. Out of 95 seropositive IgG antibody cases, 18 (18.9%) had AITD. Out of 220 pregnant, 90 (40.9%) had normal thyroid, while thyroid disorders are classified to subclinical hypothyroidism, clinical hypothyroidism, subclinical hyperthyroidism, and clinical hyperthyroidism, 64(29.1%), 41 (18.6%), 22(10.0%), and 3 (1.4%) respectively. From 95 seropositive anti-Toxoplasma IgG antibodies, high rates were detected in subclinical hypothyroidism 35(36.8%), followed by 31(32.6%) with normal thyroid, and 19 (20.0%) had clinical hypothyroidism, while only, 9 (9.5%), and 1(1.1%) had subclinical hyperthyroidism, and clinical hyperthyroidism respectively. High infection rate was recorded among women in third trimester of pregnancy 60 (44.4%) compared to lower infection rate in first trimester women 20(40.0%). The IgG antibodies seropositive women had more often highly elevated TPO antibodies than negative ones, and the latent toxoplasmosis was associated disturbance of thyroid hormones. The antibodies seropositive women had high total leucocyte count, high lymphocytes, high count of granulocytes, low total red blood cells count and low hemoglobin level.

**Conclusions:** The current study indicated that latent toxoplasmosis is associated with alteration in thyroid functions and autoimmunity during pregnancy. Pregnant women should be tested for FT3, FT4, TSH hormones, and TPO antibodies with measurement of hematological parameters in order to reduce the risk in both mother and fetus and provides early therapies.

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**Keywords:** Autoimmune thyroid diseases, ELISA, Pregnancy, Thyroid hormones, *Toxoplasma gondii*.

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**T** *oxoplasma gondii* is a common apicomplexan intracellular protozoan parasite that causes toxoplasmosis<sup>1</sup>. Many studies have indicated that about one-third of the world population has been infected with toxoplasmosis<sup>2</sup>. The members of the Felidae family, which represents domestic cats and their relatives, are the only definitive hosts, while many mammals and birds act as intermediate hosts, the disease occurs by raw or uncooked meat consumption infected with tissue cysts, or by swallowing of sporulated oocysts excreted with at feces in the water, soil, and contaminated food<sup>3</sup>. The disease is asymptomatic, while it's considered as a life-threatening condition with many serious complications especially in immune-compromised patients<sup>4</sup>. Congenital toxoplasmosis occurs during pregnancy through parasite transmission via the placenta of the infected mother to her fetus which leads to miscarriage, brain damage, retinitis, and stillbirth<sup>5</sup>. *T. gondii* has the ability to attack many human body organs, thyroid gland as an example, which produces the most important hormones T4 and T3<sup>6</sup>. Thyroid hormones are important for healthy fetus growth thus, thyroid diseases are considered as common endocrine disorders among pregnant populations, they cause spontaneous abortion, fetal damage, and preterm delivery<sup>7</sup>. The pregnancy period is affected by many environmental and endogenous factors, thyroid hormones of pregnant women have fundamental roles in the neurological development of fetus exactly in the first trimester, and because of the fetus thyroid hormones are not

produced until 16-20 weeks<sup>8</sup>. Thyroid diseases are the most risk factors for adverse pregnancy outcomes. Nowadays, researchers have found that *T. gondii* is associated with autoimmune thyroid diseases (AITD) and other autoimmune diseases. AITD is linked to various serious forms ranging from infertility, premature delivery, cesarean section, spontaneous abortion, and fetal death<sup>9</sup>. It has been described that people with genetic backgrounds may develop AITD after an infectious disease<sup>10</sup>. Molecular mimicry is recognized by resemblance and similarities in pathogen molecular components and the human thyroid gland auto antigens which lead to autoimmune disease<sup>11</sup>.

## **PATIENTS AND METHODS**

The current study was included the collection of 220 blood samples from pregnant women who attended to gynecology and obstetrics hospital in Duhok city. A special questionnaire for each woman with complete demographic characteristics (age, residency, education, occupation, pregnancy period, drinking unsterilized water, eating habits, and cat contact) was completed by the individual before the blood samples were drawn.

This study was approved by the ethical committee of the University of Duhok Scientific and Ethical approvals for the study were granted by the Scientific Committee of the College of Medicine/Duhok University. The ethical approval was obtained from the Research Ethical Committee of the Directorate General of Health, Duhok, Iraq. no. 13072021-7.

Five ml of venous blood was drawn from each woman by using a disposable sterile syringe and placed into two tubes, 1 ml of the blood was placed in a first tube with anticoagulants for measurements of blood parameters such as (total white blood cells (WBCs), lymphocytes (LYM), granulocytes (GRA), total red blood cells(RBCs), hemoglobin(Hb) and platelets(PLT). While 4 ml of the blood was placed in a second tube without anticoagulant, which was left for 20 minutes at room temperature for coagulation and then centrifuged for 10 minutes at 3000 rpm to obtain sera, the serum was placed in clean, sterile Eppendorf tubes, then kept in freezing at -200C till use<sup>12</sup>.

#### **EXCLUSION CRITERIA**

Women with other infectious diseases, immunosuppressed and chronic diseases were excluded.

#### **INCLUSION CRITERIA**

All pregnant women with different pregnancy periods.

#### **STUDY DESIGN**

Anti *T. gondii* IgG antibodies was measured using Toxoplasma ELISA kit (Bioactiva diagnostica, Germany), the determination of thyroid hormone levels was done by ELISA to measures FT3, FT4, and TSH hormones, all hormones kits (AccuBind-Monobind, USA) and TPO antibodies detection by ELISA kit (Aeskulisa, Germany) and the blood parameters were detected by coulter counter machine.

Detection of anti-Toxoplasma IgG antibodies by ELISA

Firstly, the serum samples were diluted (1:101) by distributing 10 µl of serum into

1 ml of sample diluents, then 100 µl of diluted sera, and the calibrators were pipetted to plate wells, then incubated for 30 minutes at 370C, and the wells were washed four times with washing solution, blotted and dried by inverting the plate on absorbent material. About, 100µl of enzyme-labeled secondary antibody was added to the wells and incubated for 30 minutes at 370C, the wells were washed four times, and then 100µl of TMB chromogen solution was added and incubated for 15 minutes at room temperature. Finally, 100µl of stop solution was added to the wells, the optical density (O.D) of the samples was measured at 450 nm using a plate reader.

#### **Detection of FT3 by ELISA**

Fifty µl of the calibrators, controls and sera pipetted to the wells and 100 µl of FT3 enzyme reagent was added to the wells, incubated for one hour at room temperature, the wells were washed three times by washing solution, then 100 µl of the working substrate was added and incubated at room temperature for 15 minutes, then 50 µl stop solution was added to the wells and the absorbance read at 450nm.

#### **Detection of FT4 by ELISA**

A total 50µl of the calibrators, controls and serum samples were pipetted into the microplate wells, then 100µl of FT4 enzyme reagent was added and incubated for one hour at room temperature. The wells were washed three times with washing solution, and 100µl of the working substrate was added and incubated at room temperature for 15 minutes. About 50µl of stop solution was pipetted to the wells, and the results were read at 450nm.

**Detection of TSH by ELISA**

A total 50µl of the calibrators, controls and sera samples were pipetted into the wells, 100µl of TSH enzyme reagent was added to the wells, and incubated for one hour at room temperature, the wells were washed three times with washing solution, and 100µl of the working substrate was added to the wells, incubated at room temperature for 15 minutes, then add 50µl of stop solution to the wells and the absorbance read at 450nm.

**Detection of TPO by ELISA**

A total of 100µl of each of the calibrators, negative control, positive control and serum were added to the wells and incubated for 30 minutes at 20-320C, the wells were washed three times with washing buffer, and after that, 100µl conjugate was added to the wells and incubated for 30 minutes at 20-320C, then washed three times with washing buffer, and 100µl TMB substrate was added, incubated for 30 minutes at 20-320C, after that, 100µl stop solution pipetted to the wells, incubated for 5 minutes and the absorbance read at 450 nm.

**STATISTICAL ANALYSES**

All the data were statistically analyzed using the SPSS version 22, and P-value ≤ 0.05 was considered as significant.

**RESULTS**

In the current study, overall 220 pregnant women were enrolled with mean ages of 29 years (15-50). A total of 95(43.2%) were seropositive for anti-Toxoplasma IgG antibodies, and 125 (56.8%) had IgG seronegativity. The autoimmune thyroid disease (AITD) had been indicated due to the presence of TPO antibodies. It had noticed that 28(12.7%) were screened positive for autoimmune thyroid disease, while 192(87.3%) were without AITD. Table 1, shows that high rates of IgG seropositivity were found among old aged women than young ones 69.2% and 35.7%, respectively statistical analysis was significant with P-value ≤0.05. Rural women were highly seropositive than urban residents 23(46.0%), and 72(42.3%) respectively, P-value ≥0.05. Low rates of seropositivity were recorded in high educational levels 18(38.2%), P-value ≥0.05. Unfiltered water drinker women with more seropositive 85(46.1%) than filtered water 27.7% with a statically correlation P-value ≤0.05. Cat contact women were more highly seropositive than non-cat contacts 15(62.5) and 80(40.8%), respectively.

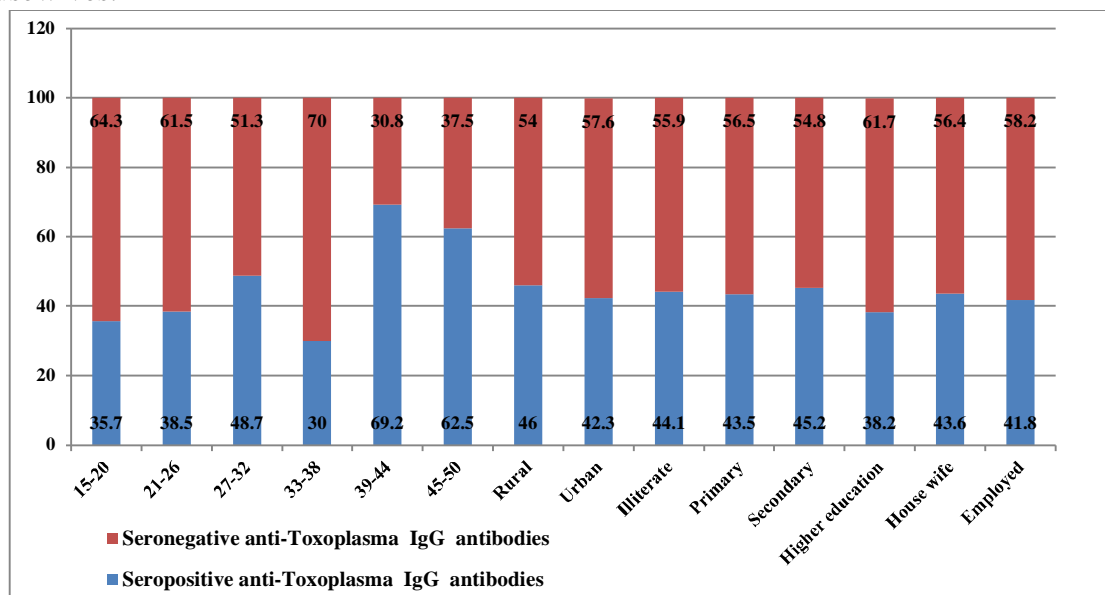
**Table1: Seropositivity of anti-Toxoplasma IgG antibodies on some demographic characteristics in pregnant women no=220**

Variables		Total	Seropositive anti-Toxoplasma IgG antibodies (%)	Seronegative anti-Toxoplasma IgG antibodies (%)	P-value
Age	15-20	14	5 (35.7)	9(64.3)	≤0.05*
	21-26	65	25(38.5)	40(61.5)	
	27-32	80	39(48.7)	41(51.3)	
	33-38	40	12(30.0)	28(70.0)	
	39-44	13	9(69.2)	4(30.8)	
	45-50	8	5(62.5)	3(37.5)	
Residency	Rural	50	23(46.0)	27(54.0)	≥0.05

Variables		Total	Seropositive anti-Toxoplasma IgG antibodies (%)	Seronegative anti-Toxoplasma IgG antibodies (%)	P-value
Education	Urban	170	72(42.3)	98(57.6)	≥0.05
	Illiterate	34	15(44.1)	19(55.9)	
	Primary	46	20(43.5)	26(56.5)	
	Secondary	93	42(45.2%)	51(54.8)	
Occupation	Higher education	47	18(38.2%)	29(61.7)	≥0.05
	Housewife	165	72(43.6%)	93(56.4)	
	Employed	55	23(41.8%)	32(58.2)	
Pregnancy period	1 <sup>st</sup> trimester	50	20(40.0%)	30(60)	≥0.05
	2 <sup>nd</sup> trimester	35	15(42.9%)	20(57.1)	
	3 <sup>rd</sup> trimester	135	60(44.4%)	75(55.6)	
Drinking water	Non filtered	184	85(46.1%)	99(53.8)	≤0.05*
	Filtered water	36	10(27.7%)	26(72.2)	
Eating habits	Restaurant foods	57	20(35.1%)	37(64.9)	≥0.05
	Home foods	163	75(46.0%)	88(54.0)	
Cat contact	Yes	24	15(62.5%)	9(37.5)	≤0.05*
	No	196	80(40.8%)	116(59.2)	

\* Statically significance (p-value ≤0.05)

In figure 1, the high seropositive rates were detected in old aged women, rural, illiterates and housewives.



**Figure 1: Demographic factors associated with seropositivity and seronegativity of anti-Toxoplasma IgG antibodies in pregnant women.**

figure 2, shows high seropositive rates had been recorded in pregnant women in the third trimester 60(44.4%), in groups of drinking unfiltered water 85(46.1%), and cat contact ones 15(62.5%).

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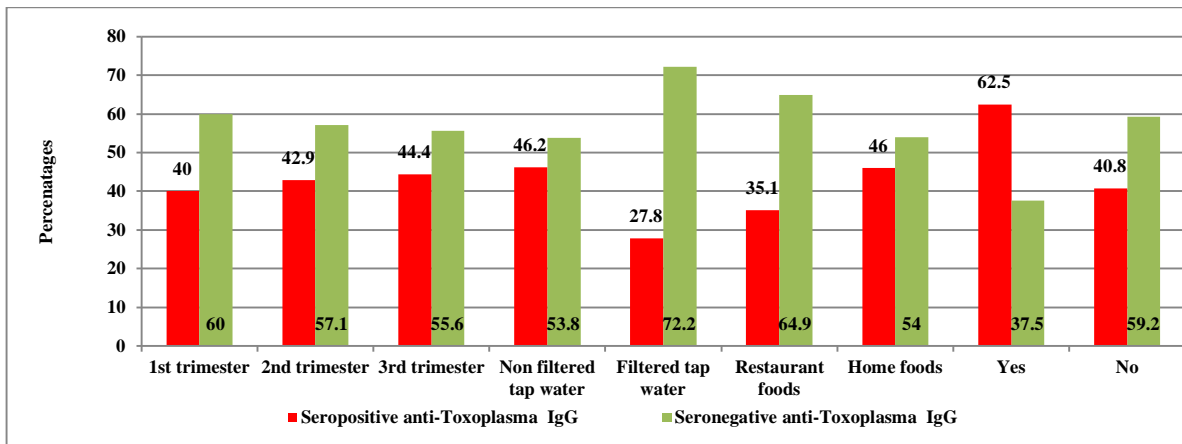


Figure 2: The period of pregnancy first trimester *Toxoplasma* IgG positive women 20 (40.0%), second trimester 15(42.9%) and third trimester 60 (44.4%), type of drinking water, non-filtered water 85(46.1%), filtered water 10(27.7%), food habits as restaurant foods 20(35.1%) and home foods75(46.0%), contact with cats 15 (62.5%) and non-cat contacts 80(40.8%).

Figure 3, shows the distribution of normal and abnormal cases of thyroiSd among women, 0.9% of cases had normal thyroid, while 29.1% and 18.6% had subclinical and clinical hyperthyroidism respectively.

On the other hand, low rates of hyperthyroidism were recorded, 10.0% and 1.4% represented subclinical and clinical hyperthyroidism respectively.

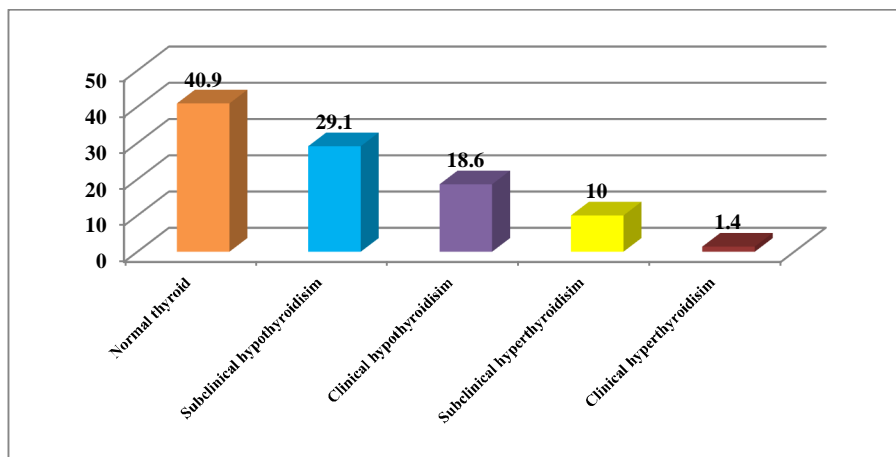


Figure 3: Thyroid status among pregnant women.

Table 2 shows the normal and abnormal thyroid cases among IgG seropositive and seronegative ones. Out of 95 IgG seropositive cases 31(32.6 %) had normal thyroid, while 64 (67.4%) were with thyroid diseases. Out of 125 IgG seronegative women about 59 (47.2%) with normal thyroid and 66 (52.8%) had thyroid disorders. Thyroid disorders which classified into subclinical hypothyroidism,

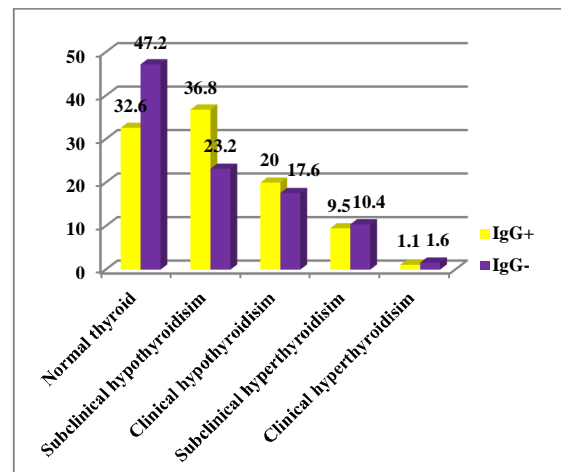
and clinical hypothyroidism, subclinical and clinical hyperthyroidism as 64 (29.1%), 41 (18.6%), 22(10.0%) and 3 (1.4%) respectively. A total of 95 IgG seropositive high rates of subclinical hypothyroidism 35(36.8%) had detected, while low rates were recorded in clinical hyperthyroidism 1(1.1%).



**Table 2: Analysis of normal thyroid and thyroid disorders among women.**

Cases	Total (%)	IgG + (%)	IgG- (%)
Normal thyroid	90(40.9)	31(32.6)	59(47.2)
Subclinical hypothyroidism	64(29.1)	35(36.8)	29(23.2)
Clinical hypothyroidism	41(18.6)	19(20.0)	22(17.6)
Subclinical hyperthyroidism	22(10.0)	9(9.5)	13(10.4)
Clinical hyperthyroidism	3(1.4)	1(1.1)	2(1.6)

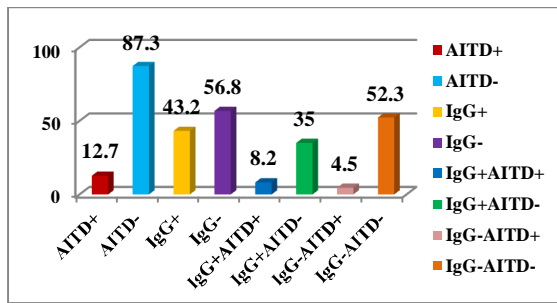
Regarding to FT3, FT4 and TSH levels, it had been shown that decreased levels of FT3 (<1.8 pg/ml) were found in 12 (12.6%) seropositive women compared to 16(12.8%) seronegative women, while elevated levels of FT3 (>4.2 pg/ml) were obtained in 5(5.3%) and 4(3.2%) in seropositive and seronegative ones respectively. The low levels of FT4 (<0.76 ng/dl) were detected in 18 (18.9%) and 18(14.4%) in IgG seropositive and IgG seronegative women respectively, while high FT4 levels (>2.24  $\mu$ IU/ml) indicated in 2(2.1%) in seropositive and 2 (1.6%) in seronegative women. On the other hand, decreased levels of TSH (<0.39  $\mu$ IU/ml) were detected in 11(11.6%) IgG seropositive and 20(16.0%) in IgG seronegative ones. Elevated TSH levels (> 6.1  $\mu$ I U /ml) were found in 51(53.7%) in seropositive and 46 (36.8%) in seronegative cases. The analysis showed that an association of IgG seropositivity and TSH levels and thyroid hormones P-value ( $\leq 0.05$ ) .It's obvious from figure 4 that high rate of IgG seropositive women had subclinical hypothyroidism 36.8%and lowest rate 1.1% noticed in clinical hypothyroidism, while most of IgG seronegative women with normal thyroid 47.9%, and 1.6% related to clinical hyperthyroidism cases.



**Figure 4: Distribution of normal thyroid and thyroid disorders among IgG seropositive and seronegative pregnant women.**

Figure 5 analyses that from overall 220 women, 28 (12.7%) had autoimmune thyroid diseases while 192 (87.3%) without autoimmune thyroid diseases. The highest rates 115 (52.3%) belonged to IgG seronegative women and without autoimmune thyroid disease and the lowest rates 10(4.5%) related to IgG-seronegative cases with autoimmune thyroid disease. Among 95 IgG seropositive cases, 18 (18.9%) had seropositivity for TPO antibodies, while from 125 IgG seronegative women, 10 (8.0%) screened positive for TPO antibodies. The analysis showed that those women with seropositivity to Toxoplasma were more frequently highly positive for TPO antibodies than seronegative ones (P-value  $\leq 0.05$ ) with a statistically significance.

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**Figure 5: Seropositivity of Toxoplasma IgG and AITD status among women.**

According to table 3, IgG seropositive women had increased counts of white blood cells, lymphocytes, granulocytes, and a decrease in red blood cells and hemoglobin compared to IgG seronegative ones.

**Table 3: Mean values of WBCs, LYM, GRA, RBCs, HGB, and PLT in seropositive IgG and seronegative ones.**

Parameters	Seropositive IgG (mean± SD)	Seronegative IgG (mean± SD)
WBCs (10 <sup>9</sup> /l)	10.80±3.04	7.92±1.75
LYM (10 <sup>9</sup> /l)	2.67±1.43	1.99±0.80
GRA (10 <sup>9</sup> /l)	6.81±2.74	3.77±1.86
RBCs(10 <sup>12</sup> /l)	3.61±0.46	4.25±0.34
HGB(g/dl)	11.19±1.18	12.36±0.89
PLT (10 <sup>9</sup> /l)	213.62±68.52	228.30±66.01

WBCs: white blood cells, LYM: lymphocytes, GRA: granulocytes, RBCs: red blood cells, HGB: hemoglobin, PLT: platelets.

### DISCUSSION

In the current study 220 pregnant women were included, 95 (43.2%) had seropositivity to anti-Toxoplasma IgG antibodies. This result showed agreement with the results of other studies Mizani et al.,<sup>13</sup> and Mohamed et al.,<sup>14</sup> in the seroprevalence of toxoplasmosis which reported 43.0 % and 44.1% respectively, while disagreeing with a study done by Al-Saeed and his group<sup>15</sup> which reported 10.0% and other studies by Imam<sup>16</sup>, Murad et al.,<sup>17</sup>, and Eissa et al.,<sup>18</sup> which showed 21.3%, 21.1% and 61.1% respectively. It has been indicated that high infection rates were found in old aged women than younger ones 69.0% with a statistically significance p-value  $\leq 0.05$ , which proves the association of infection with age, this

may be due to that old aged women with prolonged exposure to the risk factors of *T. gondii* transmission during their lives than younger ones, lack of public awareness of infection and prevention methods. This is concordant with studies by Imam<sup>16</sup> and Mizani<sup>13</sup> and disagrees with Al-Qurashi et al.<sup>19</sup>. Rural women were highly infected than urban ones which showed similar agreements with studies Tammam et al.,<sup>20</sup> and Senthamaria et al.,<sup>21</sup> this is an indication that rural women keep cats, dogs, birds and domestic animals that feed and hunt freely with low care and awareness about the disease. Also, women with cat contact showed increased infection with similar reports were observed in other studies Kolbekova<sup>22</sup>, Ngu<sup>23</sup>. High infection rates were found in illiterates, housewives, women who drinking nonfiltered water with agreements with another studies Tammam et al.,<sup>20</sup> and Senthamaria et al.,<sup>21</sup>. It is indicated that toxoplasmosis causes abnormalities in thyroid hormones and TSH levels which



come in accordance with several studies Al-Khamesi<sup>24</sup> and Hashim with his group<sup>25</sup>, while disagrees with studies of Adday<sup>26</sup> and Eissa et al.,<sup>18</sup>. The prevalence of normal thyroid and thyroid disorders were 40.9% and 59.1% respectively. The abnormalities in the levels of FT3, FT4 and TSH hormones may increase the risks of pregnancy complications such as miscarriage, preterm delivery, placental abruption and low birth weights. The results of the current study found that seropositive IgG cases had highly elevated TPO antibodies that agrees with other studies Kankova et al.<sup>7</sup> and Valizadeh et al.,<sup>27</sup>. According to the effects of thyroid disorder on both mother and fetus, TPO antibodies screening should be included during pregnancy with performing the thyroid function tests and checkup by a gynecologist must be considered. Regard to the white blood cells, lymphocytes and granulocytes were increased in IgG positive women compared to negative ones, which agrees with studies Flegr with Stiiz<sup>28</sup> and Leka<sup>29</sup>, this is due to the activation of immune system and immune cells during infection and disagree with a study Hassen et al.,<sup>30</sup>. A decrease in red blood cells and hemoglobin had observed in seropositive women which come in accordance with other studies Flegr<sup>28</sup>, Mohamed<sup>31</sup>, and Salih et al.,<sup>32</sup> this reduction in hemoglobin in seropositive for anti-Toxoplasma IgG antibodies women may be due to the multiplication of the parasite inside the host body cells and degradation of red blood cells which may cause anemia, that affects directly on the fetus and lead to preterm birth (when delivery occurs before 37 complete weeks of pregnancy) or have a low birth weight baby and postpartum depression.

## CONCLUSION

The seropositivity of anti-Toxoplasma IgG antibodies, thyroid hormones and TPO antibodies among pregnant women by using ELISA technique are useful. Increasing the public awareness programs, and prevention and control strategies are needed. It's very important to test for Toxoplasma antibodies, thyroids hormones and TPO antibodies to reduce the risks on both mother and fetus in order to provide early therapies and safe the health of the fetus. It's necessary to measure blood parameters during pregnancy period. It seems that Toxoplasma gondii may cause thyroiditis because of antigenic similarity of the parasite with thyroid peroxidase and lead to cross reactivity in immune system resulting in AITD. This could provide new clues to the complex pathogen of autoimmune thyroid disease. Thus, many studies should be done with focusing on discovering molecular similarities between thyroid peroxidase and Toxoplasma antigens.

## CONFLICT OF INTEREST

The author declared that there is no conflict of interest.

## REFERENCES

1. Foroutan M., Fakhri Y., Riahi S., Ebrahimpour S., Namroodi S., Taghipour A., Spotin A., Gamble H.R., Rostami A. The global seroprevalence of Toxoplasma gondii in pigs: A systematic review and meta-analysis. *Veterinary Parasitology*. 2019;1; 269: 42-52.
2. Mahmood O.I. Effect of Toxoplasmosis on hematological, biochemical and immunological parameters in pregnant women in Tikrit city, Iraq. *Tikrit Journal of Pure Science*. 2018; 21(3): 24-27.

3. Eskandarian A., Jahani S., Hejazi H., Yousefi H., Raissi V. Investigation of *Toxoplasma gondii* infection in Cutaneous Leishmaniasis patients of the Isfahan province. *Int J Infect.* 2017; 4 (2): 40260.
4. Wang Z.D., Liu H.H., Ma Z.X., Ma H.Y., Li Z.Y., Yang Z.B., Zhu X.Q., Xu B., Wei F., Liu Q. *Toxoplasma gondii* infection in immunocompromised patients: a systematic review and meta-analysis. *Frontiers in microbiology.* 2017; 9; 8: 389.
5. Singh S. Congenital toxoplasmosis: Clinical features, outcomes, treatment, and prevention. *Tropical parasitology.* 2016; 6(2): 113.
6. Zhang W, Wang Y, Wei Z, Chang G, Luo Q, Abudusailamu N, Nurula M, Tao S, Li H, Chen Y. Endocrine Hypertension. In *Secondary Hypertension* Springer, Singapore. 2020; 249-347.
7. Kankova S., Prochazkova L., Flegr J., Calda P., Springer D., Potlukova E. Effects of latent toxoplasmosis on autoimmune thyroid diseases in pregnancy. *PloS one.* 2014; 28; 9(10): e11087.
8. Machumi I., Mirambo M., Ruganuz D., Rambau P., Massinde AN, Kihunrwa A., Mshana S., Morona D. Factors associated with *Toxoplasma gondii* IgG and IgM antibodies, and placental histopathological changes among women with spontaneous abortion in Mwanza City, Tanzania. *The East African Health Research Journal.* 2017; 1(2): 86.
9. Stagnaro-Green A, Abalovich M, Alexander E, Azizi F, Mestman J, Negro R, Nixon A, Pearce EN, Soldin OP, Sullivan S, Wiersinga W. Guidelines of the American Thyroid Association for the diagnosis and management of thyroid disease during pregnancy and postpartum. *Thyroid.* 2011; 21(10): 1081-125.
10. Morohoshi K., Takahashi Y., Mori K. Viral infection and innate pattern recognition receptors in induction of Hashimoto's thyroiditis. *Discovery medicine.* 2011; 12(67): 505-11.
11. Benvenega S., Santarpia L., Trimarchi F., Guarneri F. Human thyroid autoantigens and proteins of *Yersinia* and *Borrelia* share amino acid sequence homology that includes binding motifs to HLA-DR molecules and T-cell receptor. *Thyroid.* 2006; 16(3): 225-36.
12. Murad M. A. Eassa S.H., Ibrahim S.A. Serosurvey of anti-toxocara antibodies and risk factors in relation to vitamin d levels among pregnant women of Duhok city–Kurdistan Region. *Duhok Medical Journal.* 2021; 15(2): 52-61.
13. Mizani A., Alipour A., Sharif M., Sarvi S., Amouei A., Shokri A., Rahimi M.T., Hosseini S.A., Daryani A. Toxoplasmosis seroprevalence in Iranian women and risk factors of the disease: a systematic review and meta-analysis. *Tropical medicine and health.* 2017; 45(1): 1-3.
14. Mohamed, Z., Hajissa, K. Seroprevalence of *Toxoplasma gondii* infection among patients in Hospital Universiti Sains Malaysia. *Trop. Biomed.* 2016; 33, 78–83.
15. Al-Saeed A.T., Eassa S.H., Murad M.A. Detection of toxoplasmosis among women with abortion using molecular and serological tests in Duhok city. *Medical.* 2016; 10(2): 56-68.
16. Imam A. *Toxoplasma gondii* infection during pregnancy. *Journal of Taibah University Medical Sciences.* 2016; 11(3) : 255-259.
17. Murad M.A., Eassa S.H., AL-Saeed A.T. Serodiagnosis of torch infections among aborted women in Duhok city/ Kurdistan Region/Iraq. *Journal of Duhok University.* 2019; 22(2): 185-94.
18. Eissa, I., Alfaki Tayseer, Al-Jaili M., Khery M., Adam R., Holie M., Elbasheir M. "The relationship between people with toxoplasmosis

- and changes in thyroid hormone levels between 2017 and 2018, Khartoum, Sudan", *Merit Research Journal of Medicine and Medical Sciences*. 2019; 7(9 ): 308-315.
19. Al-Qurashi A.R., Ghandour A.M., Obied O.E., Al-Mulhim A., Makki S.M. Seroepidemiological study of *Toxoplasma gondii* infection in the human population in the Eastern Region. *Saudi medical journal*. 2001; 22(1): 13-8.
  20. Tammam A.E., Haridy M.A., Abdellah A.H., Ahmed S.R., Fayed HM, Alsammani M.A. Seroepidemiology of *Toxoplasma gondii* infection in women with first trimester spontaneous miscarriage in Qena governorate, Egypt. *Journal of Clinical and Diagnostic Research: JCDR*. 2013; 7 (12): 2870.
  21. Senthamarai S., Sivasankari S., Apurba S., Sandhya B., Kumudavathi M., Anitha C., Amshavathani S. Seroprevalence of toxoplasmosis in pregnant women with bad obstetric history in a tertiary care hospital, Kanchipuram-a pilot study", *Disease Journal*, 2013; 3(9): 29-31.
  22. Kolbekova P., Kourbatova E., Novotna M., Kodym P., Flegr J. New and old risk-factors for *Toxoplasma gondii* infection: prospective cross-sectional study among military personnel in the Czech Republic. *Clinical Microbiology and Infection*. 2007; 1;13(10): 1012-1017.
  23. Ngui R., Lim Y.A, Amir N.F., Nissapatorn V., Mahmud R. Seroprevalence and sources of toxoplasmosis among Orang Asli (indigenous) communities in Peninsular Malaysia. *The American journal of tropical medicine and hygiene*. 2011; 85(4): 660.
  24. Al-Khamesi M.B. Effect of Toxoplasmosis on lipid profile and thyroid hormones in aborted women. *Al-Nahrain Journal of Science*. 2016; 19(4): 122-6.
  25. Hashim M.F., Dawood S.A., Ahmad Y.I. Evaluation of some Biochemical parameters in serum patients with thyroid gland dysfunction. *Basrah J. Sci*. 2018; 36(1): 42-51.
  26. Adday L. Study of hematological changes and hypertension in patients with hyperthyroidism *Journal of Al-Kufa University for Bio*. 2014; 16(1): 156-161.
  27. Valizadeh G., Khamseh M.E., Kashaniyan M, Rafiei-Sefiddashti R, Hadighi R. Role of *Toxoplasma gondii* in Thyroiditis of Pregnant Women. *Russian Journal of Infection and Immunity*. 2022.
  28. Flegr J. and Stiiz I. Potential immunomodulatory effects of latent toxoplasmosis in humans. *BMC Infectious Diseases*. 2011; 11(1): 1-7.
  29. Leka A., Shrook M. Seroprevalence of Toxoplasmosis in Rural Populations Among Pregnant Women in Babylon Province. *Journal of University of Babylon*. 2012; 20(3). 949-952
  30. Hassen A., Ali M., Ekhnafar A. Effect of *Toxoplasma gondii* Infection on Haematological and liver function parameters among abortive women in El-Beida City. *Saudi Journal of Biomedical Research*. 2019; 4(8): 295-303.
  31. Mohamed K. Hematological Changes during Chronic *Toxoplasma gondii* Infection in Pregnant Women in Makkah, Saudi Arabia. *American Journal of infectious diseases*. 2020; 16(2): 36-39.
  32. Salih J. M., Salih Mero W. M., and Eassa. S. H. Seroprevalence and some demographic factors associated with *Toxoplasma gondii* infection among female population in Duhok province, Iraq. *International Journal of Research in Medical Sciences*. 2020; 8(3): 921-926.

## پوخته

کارتیکرنا نه خووشیا پشیکا یا دوومدریژ ل سه رنه خووشین به رگیا خوویی یا په ریزادی ل ده ف نافرته تین دووگیان ل  
 باژیژی دهوک □ مه ریما کوردستان - عراق

**پشنشین و نارمانج:** نه خووشیا پشیکا نه خووشیهکا بهر به لاقه ل سه رانسهری جیهانی و مشه خووره کی ئیک خانه توکسوپلازمه گوندی یی بهر پرسیاره بو فی نه خووشی. ئەف مشه خووره دشیته تووشی گه لهک زینده وهران بیت و دبیته ئەگه ری نیشانی نه خووشی ل دهف مروفی و هیرش دکه ته گه لهک نه ندامین له شی و ده ستکاری ل تیگچوونا هورموونی و رهوشتی دکه ت ل ناف زینده وهرین تووشبووی. نه خووشیا پشیکا یا دووم دریژ بهر به لاقترین شیوهیه و دشیته کارتیکرنا ل سهر ماوی دووگیان بوونی بکه ت. نارمانج ژفی فه کوولینی، ده ستیشانکرنا به لاقبوونا نه خووشیا پشیکا و په یوه ندیا وی ل گه ل نه خووشیا بهر گریا خووی یا په ریزادی ل ده می دووگیان بوونی دا.

**ریکین کاری:** ل فی فه کوولینی دا، 220 نافرته تین دووگیان ب ده ستخووفه گرته نه ژ 1 ته باخا 2021 هه تا 1 شواتا 2022. توکسوپلازمه هاتبوو ده ستیشانکرنا بریکا دژه ته نی اج ج بریکا ب کارئینانا الایزا. ده رباری په ریزادی هورمونین ف ت 3، ف ت 4، ت س ج هاتیه پیفان ب الایزا و دژه ته نین ت پ و دیسان ب ریکا الایزا هاته پیفان. پیگهاتنین خوویی هاتنه پیفان ب ریکا نامیری س ب س.

**نه نجام:** ژ 220 نافرته تین دووگیان 95 (43.2%) ژ نافرته تین دووگیان سیروپوزه تیف بوون بو دژه ته ن اج ج یی توکسوپلازما 28 (12.7%) نه خووشیا بهر گریا خووی یا په ریزادی هه بوون ژ 95 نافرته تین سیروپوزه تیف 18 (18.9%) نه خووشیا بهر گریا خووی یا په ریزادی هه بوو. ژ 220 نافرته تین دووگیان 90 (40.9%) خودان نورمال په ریزاد بوون، تیگچوونین په ریزادی دهینه پولینکرنا بوو کیم رژینا په ریزادی یا نه کلینیکی، کیم رژینا په ریزادی یا کلینیکی، زیده رژینا په ریزادی یا نه کلینیکی و زیده رژینا په ریزادی یا کلینیکی (29.1%): 64، (18.6%): 41، (10.0%): 22 (1.4%) 3 ل دیف ئیک. ژ 95 نافرته تین سیروپوزتیف بو توکسوپلازما ی IgG ریژا زیده هاتیه دیارکرنا ل کیم رژینا په ریزادی یا نه کلینیکی 35 (36.8%)، 31 (32.6%) نورمال په ریزادی، 19 (20.0%) کیم رژینا په ریزادی یا کلینیکی و بتنی 9 (9.5%) زیده رژینا په ریزادی یا نه کلینیکی و 1 (1.1%) زیده رژینا په ریزادی یا کلینیکی هه بوون. نافرته تین ته مه ن مه زن ریژین زیده تین تووشبوونی هه بوون ژیین عه مر بچیک 5 (62.5%)، 5 (35.7%) ل دیف ئیک. نافرته تین گوندان زیده ترین ریژین تووشبوونی هه بوون 23 (46.0%) و 72 (42.3%) نافرته تین باژیژان. نافرته تین نه خاندنه فان 15 (44.1%) زیده ترین ریژین تووشبوونی و نافرته تین خودان بلین خاندنا بلند ریژین کیم 18 (38.2%). کابانین بهر مالا زیده ترین ریژین تووشبوونی هه بوون ژ نافرته تین شولکر 72 (43.6%) و 23 (41.8%) ل دیف ئیک. نافرته تین ل دوماهیك سی هه یقیته دووگیان بوونی زیده ترین ریژین تووشبوونی هه بوو 60 (44.4%) و کیم ریژین تووشبوونی ل ماوی دووگیان بوونا نیکی 20 (40.0%). نافرته تین نافا نه فله ته رگری فه دخوون زیده ترین ریژین تووشبوونی هه بوو ژ نافرته تین نافا فله ته رگری فه دخوون 85 (46.1%) و 10 (27.7%) ل دیف ئیک دا. نافرته تین تیگه ل ل گه ل پشیکا زیده ترین ریژین تووشبوونی ژ نافرته تین بی تیگه ل پشیک هه بوون 15 (62.5%) و 80 (40.8%) ل دیف ئیک. دووگیان سیروپوزه تیف بوون بو دژه ته ن اج ج یی توکسوپلازمه بلندترین ناستین ت پ هه بوو ژ نافرته تین دووگیان سیرونیکه تیف و خووشیا پشیکا یا دووم دریژ هاتبوو گریدان ب تیگچوونا هورمونین په ریزادی. نافرته تین تووشبووی گه لهک خرووکی خینی یین سبی و زیده بوونا لیمفوسایت و کیم خرووکی سوورین خوویی و کیم هیموگلوبین هه بوو.

**ده ست کهفتین فه کوولینی:** نه خووشیا پشیکا ناریشه که ل دهف نافرته تین دووگیان. گه لهک یا فهره زیده کرنا ریکارین ناگه هداریا و ناموژکاری ل دهف خه لکی ل دوور فه گوو هاستن و فاکته رین مه ترسیدارین تووشبوونی ب ریکا ناگه هداریا بو خووپاراستن و کونترولکرنا و کیمکرنا فی نه خووشی. هاته دیارکرنا کونه خووشیا پشیکا یا هاتیه گریدان ل گه ل تیگچوونین فهرمانین په ریزادی و نه خووشیا بهر گریا خووی یا په ریزادی ل ده می دووگیان بوونی دا، یا پیویسته ل سهر نافرته تین دووگیان پشکینیا بکه ن بوو هورمونین ف ت 3، ف ت 4، ت س ج و دژه ته نین په ریزادی ت پ و ل گه ل پیفانا پیگهاتنین خوویی ژبو کیمکرنا مه ترسی ل ده ف دایک و زارووی و دانا بهل بوو ده رمانان.

## الخلاصة

تأثير داء القطط الكامن على امراض الغدة الدرقية المناعية الذاتية في النساء الحوامل في مدينة دهوك - إقليم كوردستان - العراق

**الخلفية والاهداف:** داء القطط مرض شائع ذو انتشار عالمي ويسببه طفيلي احادي الخلية توكسوبلازما غوندي يمكن أن يصيب العديد من المضيفين بسبب العديد من الأعراض السريرية لدى البشر ويهاجم العديد من أعضاء الجسم مما يؤدي إلى تغيرات هرمونية وسلوكية في العوائل المصابة. داء المقوسات الكامن هو الشكل الشائع الذي يمكن أن يؤثر على مسار الحمل. يرتبط مرض الغدة الدرقية المناعي الذاتي (AITD) إلى عوامل الخطر المحددة جيداً لنتائج الحمل السلبية. الهدف من هذه الدراسة هو الكشف عن داء القطط وآثاره على أمراض الغدة الدرقية المناعية الذاتية بين النساء الحوامل.

**طرق العمل:** تضمنت الدراسة الحالية حوالي 220 امرأة حامل في فترة 1-1 اب 2021 الى 1-شباط 2022. وتم الكشف عن حالة توكسوبلازما من خلال الكشف عن الأجسام المضادة IgG للتوكسوبلازما عن طريق مقايضة الممنز المناعي المرتبط بالإنزيم (ELISA) وهورمونات FT3,FT4,TSH, الأجسام المضادة (TPO) بتقنية الايلايزا. تم قياس مكونات الدم بواسطة جهاز CBC.

**النتائج:** بشكل عام، 95 (43.2%) من النساء الحوامل لديهن إيجابية مصلية للأجسام المضادة IgG المضادة للتوكسوبلازما و28 (12.7%) تم فحصهن إيجابياً ل AITD. من 95 امرأة IgG ذات إيجابية المصل ، 18 (18.9%) كان لديهن AITD. من 220 امرأة حامل، 90 (40.9%) لديهن غدة درقية طبيعية، و اضطرابات الغدة الدرقية تنقسم الى قصور الغدة الدرقية تحت الإكلينيكي وقصور الغدة الدرقية فوق الإكلينيكي وفرط نشاط الغدة الدرقية تحت الإكلينيكي وفرط نشاط الغدة الدرقية فوق الإكلينيكي (29.1%)، 64 (29.1%)، 41 (18.6%)، 22 (10.0%)، 3 (1.4%) تعاقبياً. من 95 امرأة IgG ذات إيجابية، نسب عالية كانت قصور الغدة الدرقية تحت الإكلينيكي 35 (36.8%)، تاليا غدة درقية طبيعية 31 (32.6%) و 19 (20.0%) لديهن قصور الغدة الدرقية فوق الإكلينيكي، فقط 9 (9.5%) و 1 (1.1%) لديهن وفرط نشاط الغدة الدرقية تحت الإكلينيكي وفرط نشاط الغدة الدرقية فوق الإكلينيكي تدريجياً. النساء كبار العمر لديهن نسب اصابة عالية اكثر من النساء الشابات 5 (62.5%) و 5 (35.7%) تدريجياً. النساء القرويات لديهن نسب اصابة عالية 23 (46.0%) مقارنة بنساء المدن 72 (42.3%). النساء الاميات لديهن نسبة اصابة 15 (44.1%) والنساء ذات التحصيل الدراسي العالي لديهن نسبة اصابة 18 (38.2%). ربات البيوت كانوا اكثر اصابة من النساء المتعينات 72 (43.6%) و 23 (41.8%). سجلت نسب اصابة عالية في الاشهر الثلاثة الاخيرة من فترة الحمل 60 (44.4%) مقارنة بنسبة اصابة واطئة في ثلاثة الاشهر الاولى للحمل 20 (40.0%). النساء الشاربات المياة الغير المفطرة ذات نسب اصابة عالية اكثر من النساء الشاربات المياة المفطرة 85 (46.1%) و 10 (27.7%) تدريجياً. النساء ذات الالتماس مع القطط ذات نسب اصابة عالية اكثر من اللواتي لا يلتصون مع القطط 15 (62.5%) و 80 (40.8%) تدريجياً غالباً كان لدى النساء المصابات أجسام مضادة لTPO مرتفعة للغاية مقارنة بالأجسام غير المصابة، وداء المقوسات الكامن مرتبطاً باضطراب هرمونات الغدة الدرقية وكان لدى النساء المصابات زيادة في اعداد الخلايا دم بيضاء، زيادة في اعداد الليمفوسايت وزيادة في اعداد الخلايا الحبيبية البيضاء وقلة في اعداد خلايا الدم الحمراء وانخفاض في هيموغلوبين الدم.

**الاستنتاجات:** لايزال داء القطط يمثل مشكلة بين النساء الحوامل، ومن الضروري للغاية رفع مستوى الوعي بين الناس حول عوامل خطر العدوى والانتقال من خلال العديد من الاستراتيجيات والطرق للوقاية من المرض ومكافحته. يشار إلى أن داء المقوسات يرتبط بالتغيير في وظائف الغدة الدرقية والمناعة الذاتية أثناء الحمل. يجب اختبار النساء لحوامل بحثاً عن هرمونات FT3 و FT4 و TSH والأجسام المضادة لـ TPO مع قياس معايير الدم من أجل تقليل المخاطر في كل من الأم والجنين وتزويد الأدوية.