

Available online freely at www.isisn.org

Bioscience Research Print ISSN: 1811-9506 Online ISSN: 2218-3973 OPEN ACCESS

RESEARCH ARTICLE

Journal by Innovative Scientific Information & Services Network



Effect of chronic toxoplasmosis in the function of pancreas, liver, and kidney in pregnant women in Makkah, Saudi Arabia

Khalil Mohamed

Department of Epidemiology, Faculty of Public Health & Health Informatics, Umm Al-Qura University, Saudi Arabia

*Correspondence: kmismail@uqu.edu.sa Received:06-05-2023, Revised: 29-05-2023, Accepted: 12-06-2023 e-Published: 17-06-2023

Toxoplasmosis is a zoonotic parasitic disease worldwide which causes abortion to pregnant women. The aim of the current study to explore the effectiveness of chronic toxoplasmosis to the functions of pancreas, liver, and kidney to pregnant women. Case control study was performed in pregnant women in Makkah using ELISA IgG to differentiate chronic toxoplasmosis. Different biochemical tests were used to detect different substances indicated the functions of pancreas, liver, and kidney. ANOVA test was used to analysis the data in SPSS. Approximately, 326 pregnant women were participated in the present study. The results showed significance in the accumulated blood sugar in the grandmultigravida group with chronic toxoplasmosis *p*- value < 0.05. The current study also found significant changes in AST which was increased in grandmultigravida with chronic toxoplasmosis significantly *p*- value < 0.05. Also, serum urea increased at the first trimester stage in pregnant women *p*- value < 0.005. Serum creatinine was significant in the pregnant women at first trimester *p*- value < 0.001.chronic infection with T. gondii in pregnant women is associated with hypo-glycemia particularly in multigravida. Moreover, association was detected between liver necrosis particularly in grand multigravida and chronic toxoplasmosis. In addition, exposure to chronic toxoplasmosis may cause acute or chronic damage to the kidney

Keywords: Chronic toxoplasmosis; Pregnant women; Hypoglycemia; Liver necrosis; Kidney damage.

INTRODUCTION

Toxoplasmosis is zoonotic parasitic disease caused by protozoan known as *Toxoplasma gondii* (*T. gondii*). The definitive host of the parasite is cat and all feline groups (Sonar and Brahmbhatt, 2010). The parasite can infect wide range of hosts included man and has global distribution (Sedlak and Bartova, 2012). People may acquire infection horizontally by ingestion raw meat contains bradyzoites or water contains oocysts or vertically from mother to her foetus or via blood transfusion by tachyzoites (Robert-Gangneux and Dardé, 2012). Toxoplasmosis was defined according to serological prevalence data as the most common disease in human throughout the world and the disease was common in worm climates area (Sonar and Brahmbhatt, 2010).

Infection with *T. gondii* distributed worldwide and considered as one of famous prevalent parasite to human (Tenter, 2000).

The infection with *T. gondii* can lead to several immunological changes in the body of the host which are led to product different immunoglobulins such as IgM, IgA, and IgG (Dubey, 1994).

In acute stage of the disease, the parasite can be seen in the blood or others liquids of the body such as

cerebrospinal fluid, tears, semen, urine and saliva. The acute infection usually leads to abortion in human and animals (Guruz and Ozcel, 2007; Mahmood, 2016).

The disease classified to acute, sub-acute, and chronic according to immunoglobulins detected in the blood. Detection of IgM indicated the acute phase while detected of IgG leads to chronic phase of the disease (Lynn et al. 2023; Liu et al. 2015).

Few studies were done previously to elucidate the relationship between *T. gondii* infection and liver or kidney diseases. In Turkey some study associated liver cirrhosis with *T. gondii* infection (Kodym et al. 2023). Some other study found the association between toxoplasmosis and kidney function (Atmaca et al. 2012). Moreover, toxoplasmosis causes acute and severe damage in the infected liver (Atmaca et al. 2013; El-Sayed et al. 2016). The relationship between raising level of urea and creatinine in pregnant women infected with *T. gondii* were explored (Alvarado-Esquivel et al. 2011).

Previous study in pregnant women in Makkah found that the prevalence of chronic toxoplasmosis was 5.6% (Dalimi and Abdoli, 2012).

The main aim of the current study was to detect the effectiveness of the chronic infection of *T. gondii* to the liver, pancreas, and kidney functions.

MATERIALS AND METHODS

Study design and population

A case control study was performed to study the effectiveness of chronic toxoplasmosis to liver and kidney functions in pregnant women. Pregnant women in different stages of pregnancy and different ages were considered in the current study. The study was done in two groups of pregnant women; the first group with chronic toxoplasmosis (IgG positive) and the second group without chronic toxoplasmosis (IgG negative) both groups were (IgM negative). Also, the effectiveness was studied according to the age-group; pregnant women were divided into two groups; group one with chronic toxoplasmosis and age between 16-26 years with control from the same age group and the group two ages between 27-46 years with control. The study also considered the stages of pregnancy, each stage with control.

Data collection

Data was collected from each subject after signed the consent form and accepted to include in the study. The questionnaire was distributed to individual; the questionnaire included socio-demographic information including age, education, with clinical information such as history of abortion, and pregnancy stage.

Samples collection

Blood sample of 5 ml was collected from each participant and divided into two EDTA tubes. The first tube was separated and plasma was stored in -20 °C to be used in ELISA to differentiate between pregnant women in getting chronic toxoplasmosis. The second tube was used to measure some enzymes and other biochemical tests.

ELISA IgG and IgM

Using ELISA IgG and IgM to differentiate between chronic infection, acute infection, and control. Human TOXO kit- Germany was used according to manufactory constructions. All plasma samples from pregnant women were analysed using the above-mentioned kits. The results divided the participants into three groups: group one which were positive to IgM and excluded from the study, group two positive to IgG and group three which were negative to IgG and IgM. Both group two and three were considered in the current study.

Biochemical analysis

Biochemical analysis was carried out to determine the functions of pancreas, liver, and kidney. Different kits were used to determine different enzymes according to the certain organ. The techniques were followed the manufacturing procedures for each kit.

Data analysis

Analysis data was performed using SPSS program version 26. The form of variable put in mean \pm SD. Data were analysed using independent t test besides ANOVA test. Significance was approved for *p*-values less than 0.05.

RESULTS

Chronic toxoplasmosis was found more in grandmultigravida at third trimester with no history of abortion and their age group between 28 and 45 of pregnant women when compared with other groups as seen in table (1).

	Тох	o lgG					
Variable	Positive Negative (Infected (Control		Total				
	Group)	Group)					
	Age Grou	р					
	Years						
17-27	21(17.9%)	96(82.1%)	117				
28-45	48(23.0%)	161(67.0%)	209				
	Gravity						
Primgravida	5(12.5%)	35(87.5%)	40				
Multigravida	14(23.0%)	47(77.0%)	61				
Grand	40(22.70/)	100/76 20/)	160				
multigravida	40(23.7%)	129(70.3%)	109				
Pregnancy							
	Stage						
First trimester	17(24.3%)	53(75.7%)	70				
Second	20(22 50/)	65/76 50/)	05				
trimester	20(23.5%)	05(70.5%)	00				
Third trimester	32(19.0%)	136(81.0%)	168				
History of							
	Abortion	l					
Yes	23(18.3%)	103(81.7%)	126				
No	46 (23.0%)	154(77.0%)	200				

The effectiveness of chronic toxoplasmosis to pancreas activity detected by using the level of blood glucose, lactate dehydrogenase (LDH), and an amylase test. The results showed significance changes in the accumulated blood sugar in the multigravida group with chronic toxoplasmosis p- value< 0.031table (2) while no significance changes in lactate dehydrogenase or in an amylase.

The effectiveness of chronic toxoplasmosis to liver functions detected by measuring the level of serum alanine amino transferees (ALT), serum aspartate aminotransferase (AST), and serum alkaline Phosphatase (ALP).

	10	JICZ. TH	circuivenes.				paricicas func		pregnant wome	1		
		P	ositive IgG (Infec	ted Group)			Negative IgG (Control Group)					
Variable	Glucose	P-	LDH	P-	A	P-	Glucose	P-	LDH	P-	A	P-
	mmol/L	value	IU/L	value	Ату	value	mmol/L	value	IU/L	value	Ату	value
				-	Age Gro	oup Years					•	
17-27	4.3705±0.69	0.766	202.2±29.5	0.090	61±17.8	0.074	4.4317±1.1	0.766	182.5±31.7	0.090	48.6±17.5	0.074
28-45	5.0852±1.87	0.766	194.14±33	0.089	45±13.7	0.274	4.9698±1.78	0.766	178.04±44	0.069	37.6±18.8	0.274
					Gr	avity						
Primgravida	4.5280±0.45		173.0±42.6		50.5±36.1		4.4960±0.64		173.0±42.6		50.5±36.1	
Multigravida	4.1957±0.73	0.021	208.5±20.1	0.691	41.0±0.0	0.000	4.7687±1.1	0.021	183.1±32.9	0.691	47.6±15.4	0.000
Grandmult igravida	5.2088±1.9	0.031	196.1±35.4	0.001	52.5±18.7	0.693	5.0717±1.8	0.031	178.5±45.1	0.001	39.1±19.6	0.693
					Pregna	ncy Stage						
First trimester	4.9129±1.3		161.0±24.8		67.5±10		4.7338±0.8		169.5±27.1		25.6±4.5	
Second trimester	5.5110±2.4	0.785	189.5±27.6	0.089	49.6±20	0.274	4.8149±1.9	0.785	182.1±32.3	0.089	37.3±3.7	0.274
Third trimester	4.4244±0.9		207.3±29.5		50.2±17	50.2±17	4.8237±1.5		183.1±46.5		51.1±16.8	
History of Abortion												
Yes	4.9400±1.3	0 790	180.6±36.2	0.090	39.00±12.7	0.274	5.0872±1.9	0.790	207.63±24.26	0.090	37.11±18.8	0.074
No	4.5896±1.8	0.760	173.7±41.5	0.089	60.00±14.7	0.274	4.6107±1.2	0.780	186.17±38.5	0.069	48.88±17.3	0.274

Table2: The effectiveness of chronic toxoplasmosis on pancreas functions in pregnant women

Table3: The effectiveness of chronic toxoplasmosis on liver functions in pregnant women

	Positive IgG (Infected Group)					Negative IgG (Control Group)						
Variable	ALT IU/L	P-value	AST IU/L	P-value	ALP IU/L	P- value	ALT IU/L	P- value	AST IU/L	P- value	ALP IU/L	P- value
Age Group Years												
17-27	19.99±50.2	0.175	25.5313±32.4	0.067	108.0±49.7	0.550	10.19±6.8	0.175	20.0247±5.1	0.067	123.2±66.1	0.550
28-45	10.43±6.1	0.175	20.7703±8.0	0.067 0.067	104.5±55.4	0.550	9.96±9.4	0.175	19.1266±5.5	0.007	105.9±61.3	0.550
Gravity												
Primgravida	7.1667±3.7		18.1333±4.5		-		11.1367±9.8		19.3533±6.2		144.9±100.8	
Multigravida	8.9077±7.7	0 122	21.2154±10.5	0.013	110.25±54.1	0.670	10.7189±12.5	0 1 2 2	19.1622±6.3	0.013	133.65±65.1	0.670
Grand multigravida	16.8448±34.5	0.125	24.9815±24.9	24.9815±24.9	104.38±55.1	0.079	9.4363±6.5	0.123	18.8057±4.5	0.015	99.42±54.2	0.079
Pregnancy Stage				-								
First trimester	12.2909±6.2		19.3000±6.4		64.4±17.9		12.6364±6.5		18.7978±4.2		69.1±22.9	
Second trimester	22.0643±49.7	0.180	27.6667±37.5	0.070	77.7±34.1	0.493	11.3556±11.2	0.180	19.2833±5.7	0.070	72.1±22.8	0.493
Third trimester	8.0000±5.6		21.2179±8.2		145.5±64.7		8.3679±7.2		19.8991±5.6		144.7±50.1	
History of Abortion												
Yes	11.5647±4.7	0.172	21.2125±6.2	0.069	107.1±65.8	0.511	9.9540±6.4	0.172	18.9978±4.6	0.069	100.7±60.1	0.511
No	13.7057±31.9		22.6378±5.4		118.9±62.1		10.1025±9.7		19.8628±5.4		108.6±49.4	

Variable		Po: (Infec	sitive IgG cted Group)		Negative IgG (Control Group)				
	Urea (mg/dl)	P-value	Creatinine(mg/dl)	P-value	Urea (mg/dl)	P-value	Creatinine(mg/dl)	P-value	
Age Group Years									
17-27	2.2±0.8	0.788	47.9±5.7	0.778	2.7±1.6	0.788	47.9±6.0	0.778	
28-45	2.8±2.0		48.0±4.8		2.6±1.5		48.1±6.4	1	
Gravity		•	•			•			
Primgravida	2.2±0.6	0.034	48.6±8.2	0.531	2.2±0.6	0.034	47.1±5.2	0.531	
Multigravida	3.2±2.6		47.3±3.2		2.9±1.7		48.9±6.2	1	
Grandmultigravida	2.5±1.6		48.1±5.2		2.7±1.4		48.4±6.7	1	
Pregnancy Stage		•	•			•			
First trimester	2.9±2.1	0.009	50.3±4.5	0.001	3.1±1.6	0.009	50.4±6.4	0.001	
Second trimester	2.6±1.7		47.1±6.3		2.7±1.6		47.9±6.3	1	
Third trimester	2.4±1.5		47.3±4.2		2.4±1.2		47.2±5.9]	
History of Abortion									
Yes	2.2±1.0	0.258	50.6±4.6	0.433	2.6±1.3	0.258	47.9±5.9	0.433	
No	2.8±2.0		46.7±4.7]	2.7±1.6		48.2±6.5	7	

	Table4: The effectiveness of	chronic toxoplasmosis	on kidney functions in	pregnant women
--	------------------------------	-----------------------	------------------------	----------------

The results which obtained from the current study found significant changes in AST which was increased in grandmultigravida with chronic toxoplasmosis p- value < 0.013as showed in table (3) but no significance changes were detected in serum aspartate aminotransferase (AST), and serum alkaline Phosphatase (ALP).

The effectiveness of chronic toxoplasmosis to kidney functions clarified by estimation the level of urea and creatinine in the serum. According to the results obtained, the chronic toxoplasmosis increased the urea in multigravida infected with chronic toxoplasmosis significantly *p*- value < 0.034. Also, serum urea decreased at the first trimester stage in pregnant women *p*- value < 0.009 as appeared in table (4). Serum creatinine was significant in the pregnant women at first trimester *p*- value < 0.001 as mentioned in table (4)

DISCUSSION

Very little is known about the effectiveness of chronic toxoplasmosis in pregnant women. The current study adjusts to go behind epidemiological studies and study the effectiveness of latent *T. gondii* in a natural host without manipulation, particularly recent studies confirmed that latent toxoplasmosis unsafe for human (Flegr et al. 2003). The majority of the studies connected between latent toxoplasmosis and the behaviour changes in human (Novotna et al. 2008; Lindova et al. 2006; Mihu et al. 2020). This study aimed to clarify the effectiveness of chronic toxoplasmosis on the function of some organs via studying the changes in the level of some important secession.

As the study targeted the impact of chronic toxoplasmosis therefore, it was ordinary to find that the majority of chronic infection in grandmultigravida or in aged women. Similar study found that chronic toxoplasmosis tended to increase with age in Romina (Majidiani et al. 2016).

According to the finding in the current study, the association between chronic toxoplasmosis and hypoglycemia observed in the multigravida. Several studies done recently concluded that there is no correlation between infection with toxoplasmosis and type-1 or type-2 diabetes mellitus (Khalili et al.2018; Catchpole et al. 2023). Other study found correlation between toxoplasmosis infection and type-1 diabetes mellitus (Molan et al. 2020). Moreover, some studies found correlation between infection of toxoplasmosis and type-2 diabetes mellitus (Roller et al. 1987). The target group of our study was different, as the study here targeted pregnant women beside the result showed hypoglycemia in pregnant women at multigravida stage. This finding needs more studies as hypoglycemia during pregnancy may lead to complicates to women and her foetus included difficult labours and additional monitoring.

Studying hepatic enzymes are good marker for hepatocellular injury (Mahmood and Dawood, 2012). In

the present study, the results shown significant in AST compared with control and the significant found exactly in grandmultigravida which is elevated. The current results matching with different results publishing (Limdi and Hyde, 2003; Babekir et al. 2022a). This result may link between chronic toxoplasmosis and liver necrosis particularly in grandmultigravida. As in some previous study found that *T. gondii* exposure was associated with an elevated relative risk of chronic liver disease andnonalcoholic fatty liver disease (Ocak et al. 2005) therefore, clinical study should be performed to confirm the effectiveness of chronic toxoplasmosis in liver cells.

In the kidney the results being contrasting regarding urea level, the level of urea increased significantly in multigravida while decreased significantly in the first trimester. The level of creatinine was decreased particularly in the first trimester. Exposure to *T. gondii* may cause acute or chronic damage to the kidney, triggering injury, which can affect the exposure over their life course. Prior studies have found a link between undergoing dialysis and an increased rate of *T. gondii* infection (Saadat et al. 2020; Babekir et al. 2022b).

CONCLUSION

In conclusion, chronic infection with *T. gondii* in pregnant women is associated with hypoglycemia particularly inmultigravida. Moreover, association was detected between liver necrosis particularly in grandmultigravida and chronic toxoplasmosis. In addition, exposure to chronic toxoplasmosis may cause acute or chronic damage to the kidney

CONFLICT OF INTEREST

The authors declared that present study was performed in absence of any conflict of interest.

ACKNOWLEDGEMENT

The author would like to thank Mr. Mohand Gafar and Mr. Abdelrahman Almhmadi for their kind help and good collaboration.

Copyrights: © 2023@ author (s).

This is an open access article distributed under the terms of the **Creative Commons Attribution License (CC BY 4.0)**, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author(s) and source are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

REFERENCES

Alvarado-Esquivel C, Torres-Berumen JL, Estrada-Martínez S, Liesenfeld O, Mercado-Suarez MF. *Toxoplasma gondii* infection and liver disease: a

Effect of toxoplasmosis in pregnant women

Khalil Mohamed.

case-control study in a northern Mexican population. Parasit Vectors. 2011; 13;4:75. doi: 10.1186/1756-3305-4-75.

- Atmaca HT, Gazyagci AN, Canpolat S, Kul O. Hepatic stellate cells increase in *Toxoplasma gondii* infection in mice. Parasite & Vectors. 2013; 6:1-66.
- Atmaca HT, Ocal N, Babur C, Kul O. Reactivated and clinical *Toxoplasma gondii* infection in young lambs: Clinical, serological and pathological evidences. Small Rum Res. 2012;105: 335- 340.
- Babekir A, Mostafa S, Minor RC, Williams LL, Harrison SH, Obeng-Gyasi E. The Association of Toxoplasma gondii IgG and Liver Injury in US Adults. Int J Environ Res Public Health. 2022 Jun 19;19(12):7515. doi: 10.3390/ijerph19127515.
- Babekir, A.; Mostafa, S.; Obeng-Gyasi, E. The Association of *Toxoplasma gondii* IgG Antibody and Chronic Kidney Disease Biomarkers. Microorganisms 2022, 10, 115. https://doi.org/10.3390/microorganisms10010115
- Catchpole A, Zabriskie BN, Bassett P, Embley B, White D, Gale SD, Hedges D. Association between *Toxoplasma gondii* Infection and Type-1 Diabetes Mellitus: A Systematic Review and Meta-Analysis. International Journal of Environmental Research and Public Health. 2023; 20(5):4436. ttps://doi.org/10.3390/ijerph20054436
- Dalimi A, Abdoli A. Latent Toxoplasmosis and Human. Iran J Parasitol. 2012; 7(1): 1–17.
- Dubey JP. Toxoplasmosis, J. Am. Vet. Med. Assoc. 1994; 205: 1593-1598.
- El-Sayed NM, Ramadan ME, Ramadan ME. *Toxoplasma* gondii Infection and Chronic Liver Diseases: Evidence of an Association. Trop Med Infect Dis. 2016 1;1(1):7. doi: 10.3390/tropicalmed1010007.
- Flegr J, Preiss M, Klose J, Havllcek J, Vitakova M, Kodym P. Decreased level of psychobiological factor novelty seeking and lower intelligence in men latently infected with the protozoan parasite *Toxoplasma gondii* Dopamine, a missing link between schizophrenia and toxoplasmosis? Biol Psychol. 2003;63:253–268.
- Guruz AY, Ozcel MA. Toxoplasmosis. In Özcel A (ed) TıbbiParazitHastalıkları, Meta basımMatbaacılıkHizmetleri, İzmir. 2007; 141-189.
- Khalili M, Mahami-Oskouei M, Shahbazi A, Safaiyan A, Mohammadzadeh-Gheshlaghi N, Mahami-Oskouei L. The Correlation between Serum Levels of Anti-*Toxoplasma gondii* Antibodies and the Risk of Diabetes. Iran J Parasitol. 2018 ;13(4):637-642.
- Kodym P, Kurzova´ Z, Berenova´ D, Maly´ M () Detection of persistent low IgG avidity–an interpretative problem in the diagnosis of acute toxoplasmosis. PLoS ONE 2023; 18(4): e0284499. https://doi.org/10.1371/journal.pone.0284499
- Limdi J, Hyde G. Evaluation of abnormal liver function tests. Post Med 2003; 79: 307 312.

- Lindova J, Novotna M, Havlicek J, Jozifkova E, Skallova A, Kolbekova P, Hodny Z, Kodym P, Flegr J. Gender differences in behavioural changes induced by latent toxoplasmosis. Int J Parasitol. 2006;36:1485–1492.
- Liu, Q., Wang, ZD., Huang, SY. et al. Diagnosis of toxoplasmosis and typing of *Toxoplasma gondii*. Parasites Vectors.2015; 8, 292,. https://doi.org/10.1186/s13071-015-0902-6.
- Lynn MK, Aquino MSR, Self SCW, Kanyangarara M, Campbell BA, Nolan MS. TORCH Congenital Syndrome Infections in Central America's Northern Triangle. Microorganisms. 2023; 11(2):257. https://doi.org/10.3390/microorganisms11020257
- Mahmood O.Effect of Toxoplasmosis on hematological, biochemical and immunological parameters in pregnant women in Tikrit city, Iraq. Tikrit Journal of Pure Science.2016; 21(3);24-27.
- MahmoodNAJ, DawoodMN. Liver function tests in toxoplasmosis. Ann. Coll. Med. Mosul 2012; 38 (2): 68-72.
- Majidiani H, Dalvand S, Daryani A, Galvan-Ramirez L, Foroutan- M. Is chronic toxoplasmosis a risk factor for diabetes mellitus? A systematic review and metaanalysis of case–control studies. The Brazilian Journal of Infectious Diseases. 2016: 20(6): 605-609.
- Mihu AG, Balta C, Marti DT, Paduraru AA, Lupu MA, Olariu TR. Seroprevalence of *Toxoplasma gondii* infection among women of childbearing age in an endemic region of Romania, 2016-2018. Parasite. 2020;27:59. doi: 10.1051/parasite/2020057.
- Molan, A., Nosaka, K., Hunter, M. The association between *Toxoplasma gondii* and type 2 *diabetes mellitus*: a systematic review and meta-analysis of human case-control studies. Bull Natl Res Cent 2020; 44, 7. https://doi.org/10.1186/s42269-019-0256-x.
- Novotna M, Havlicek J, Smith P, Kolbekova P, Skallova A, Klose J, Gasva Z, Pisacka M, Sechovska M, Flegr J. *Toxoplasma* and reaction time: role of toxoplasmosis in the origin, preservation and geographical distribution of Rh blood group polymorphism. Parasitology. 2008;135:1253–1261.
- Ocak, S.; Duran, N.; Eskiocak, A.F.; Aytac, H. Anti-*Toxoplasma gondii* antibodies in hemodialysis patients receiving long-term hemodialysis therapy in Turkey. Saudi Med. J. 2005, 26, 1378–1382.
- Robert-Gangneux F, Dardé ML. Epidemiology of and diagnostic strategies for toxoplasmosis. Clin Microbiol Rev. 2012;25(2):264-96. doi: 10.1128/CMR.05013-11.
- Roller A, Bartlett A, Bidwell D. Enzyme Immunoassay with special reference ELISA technique. J Clin Path 1987; 31: 507 520.
- Saadat, F.; Mahmoudi, M.R.; Rajabi, E.; Roshan, Z.A.; Shad, B.M.; Karanis, P. Seroepidemiology and associated risk factors of *Toxoplasma gondii* in hemodialysis patients. Acta Parasitol. 2020, 65, 906–

- Sedlak K, Bartova E. The prevalence of *Toxoplasma gondii* IgM and IgG antibodies in dogs and cats from the Czech Republic. VeterinárníMedicína. 2012; 51(12): 555-558.
- Sonar SS, Brahmbhatt MN. Toxoplasmosis: An Important Protozoan Zoonosis. Veterinary World. 2010; 3(9):436-439.
- Tenter A, Heckeroth A, Weiss L. *Toxoplasma gondii*: from animals to humans. Int. J. parasitol.2000; 30: 1217-1258.