# Changes in the personality profile of young women with latent toxoplasmosis

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Abstract. Latent toxoplasmosis is the most widespread parasite infection in developed and developing countries. The prevalence of *Toxoplasma gondii* infection varies mostly between 20 to 80% in different territories. This form of toxoplasmosis is generally considered to be asymptomatic. Recently published results, however, suggest that the personality profiles of infected subjects differ from those of uninfected controls. These results, however, were obtained on non-standard populations (biologists or former acute toxoplasmosis patients). Here we studied the personality profiles of 191 young women tested for anti-*Toxoplasma* immunity during gravidity. The results showed that the differences between *Toxoplasma*-negative and *Toxoplasma*-positive subjects exits also in this sample of healthy women. The subjects with latent toxoplasmosis had higher intelligence, lower guilt proneness, and possibly also higher ergic tension. The difference in several other factors (desurgency/surgency, alaxia/protension, naiveté/shrewdness, and self-sentiment integration) concerned changes in the variances, rather than the mean values of the factors.

One of the most widespread parasites in the Czech Republic is the protozoan Toxoplasma gondii Nicolle et Manceau, 1908. The longtime prevalence in normal population of this parasite is about 30%, while the prevalence in higher age categories and within some risk groups increases above 50% (Sengbusch and Sengbusch 1978, Němec et al. 1988, Yamaoka and Konishi 1993). Four distinct forms of toxoplasmosis exist. The most serious form is the congenital toxoplasmosis which often leads to serious malformations including hydrocephaly, intra-cerebral calcification and microcephaly (Koppe and Rothova 1989). The second form is the acquired acute toxoplasmosis. It is characteristic by the presence of T. gondii tachyzoites in the blood and other tissues of infected people and by a complex of clinical symptoms which varies from fever and headache to serious neurological and psychiatric malfunctions (Kramer 1966). In the most cases, however, the acute toxoplasmosis is only a mild disease which is often misdiagnosed as a common bacterial or viral infection. Infrequently the acute toxoplasmosis evolves into chronic disease in which the clinical symptoms as well as high titres of specific antibodies persist for many years. Mostly, all symptoms of acute disease quickly fall away, the antibody titres decrease and the toxoplasmosis evolves into the latent form of the infection. During the latent toxoplasmosis the parasite survives in the dormant form of bradyzoites mostly in the neural and muscular tissue of the host. The latent toxoplasmosis probably lasts for the whole life of infected person (Remington and Krahenbuhl 1982) and it can turn into acute toxoplasmosis only after serious violence of integrity of immune systems (AIDS, treatment

with immunosuppressive drugs etc.), (Heitman and Irizarry 1997, Mocsny 1992).

Latent toxoplasmosis is considered to be asymptomatic. Most of the infected people never learn to have met T. gondii in the past and carry the dormant stages of the parasite for the rest of their life. Recently, however, significant differences in personality profiles of men and women with latent toxoplasmosis and normal controls were reported to exist (Flegr and Hrdý 1994, Flegr et al. 1996). Toxoplasma-positive subjects differ by mean value of some psychologic factors measured by 16 PF Cattell's personality questionnaire. The differences in some of these factors increase with the duration of latent toxoplasmosis (Flegr et al. 1996). This suggests that the changes were induced by the toxoplasmosis rather than the personality factor shift influencing the probability of being infected with T. gondii. Under natural conditions the intermediate hosts of Toxoplasma are small rodents and the final host is a cat. Therefore, the changes in the human personality can be a product of manipulation activity of the parasite which is primarily aimed to increase the probability of transmission from intermediate to a definitive host by predation. Such manipulation activity of T. gondii was observed in many experimental systems (Hutchison et al. 1980a,b, Arnott et al. 1990, Webster 1994, Webster et al. 1994, Berdoy et al. 1995,). The mechanism of manipulation activity of Toxoplasma gondii is not known. However, it can probably include the synthesis (or induction of synthesis) of some neuromodulator (Stibbs 1985) or of other biologically active substance (Varela et al. 1956, 1957). In the aberrant host (man) such activity of the parasite is of course nonproductive. However, it can

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**Table 1.** List of sixteen personality factors monitored by Cattell's questionnaire. The names and characteristics in the left column are for persons with low values of the factor, those in the right column for persons with high values of the factor. The traits that increase in young women with latent toxoplasmosis are typed in bold.

SIZOTHYMIA	A	AFFECTOTHYMIA		
reserved, detached, critical		warm-hearted, outgoing, easygoing		
LOW INTELLIGENCE	В	HIGH INTELLIGENCE		
EGO WEAKNESS	C	HIGH EGO STRENGTH		
affected by feelings, emotionally less stable		stable, mature, faces reality, calm		
SUBMISSIVENESS	E	DOMINANCE OR ASCENDANCE		
obedient, mild, easily led, docile		aggressive, competitive, stubborn		
DESURGENCY	F	SURGENCY		
sober, taciturn, serious		enthusiastic, heedless, happy-go-lucky		
LOW SUPEREGO STRENGTH	G	SUPEREGO STRENGTH, CHARACTER		
disregards rules, expedient		conscientious, persistent, moralistic, staid		
THRECTIA	Н	PARMIA		
shy, timid, restrained		adventurous, "thick-skinned", socially bold		
HARRIA	I	PREMSIA		
tough-minded, rejects illusions		tender-minded, sensitive		
ALAXIA	L	PROTENSION		
trusting, accepting conditions, tolerant		suspecting, jealous, dogmatic		
PRAXERNIA	M	AUTIA		
practical, has "down to earth" concerns		imaginative, bohemian		
NAIVETÉ	A	SHREWDNESS		
forthright, unpretentious		astute, worldly, polished		
UNTROUBLED ADEQUANCY	0	GUILT PRONENESS		
self-assured, placed, secured		apprehensive, self-reproaching, insecure		
CONSERVATISM OF TEMPERAMENT	Q1	RADICALISM		
conservative, respecting		experimenting, liberal, analytical		
GROUP DEPENDENCY	Q2	SELF-SUFFICIENCY		
sociably group-dependent, "joiner"		self-sufficient, resourceful, prefers own decisions		
LOW SELF-SENTIMENT INTEGRATION	Q3	HIGH STRENGTH OF SELF SENTIMENT		
uncontrolled, lax, follows own urges		controlled, exacting will power, socially precise		
LOW ERGIC TENSION	Q4	HIGH ERGIC TENSION		
relaxed, tranquil, torpid		tense, frustrated, driven, overwrought		

manifest its presence by nonspecific (from the point of view of a manipulation hypothesis) changes in human personality.

In the previous studies the personality changes were studied either on population of biologists or on subjects previously diagnosed for acute toxoplasmosis (Flegr and Hrdý 1994, Flegr et al. 1996). Neither of the experimental sets represents a typical population. Therefore, it is difficult to guess whether the obtained results can be generalised for normal population. Moreover, in both cases the subjects were informed that the aim of the research is to reveal possible influence of toxoplasmosis on human personality. Such information could influence the subject's responses on the questionnaire (at least in the second study in which the subjects were aware to be infected by *Toxoplasma gondii* before the questionnaire testing).

The aim of the present study was to find out whether the differences in personality profiles can be detected also in a random sample of normal women previously tested for toxoplasmosis during pregnancies. For ethical reasons (a possible fear of young mothers of congenital toxoplasmosis) the women could not be informed that the research concerns the toxoplasmosis. The absence of *a priori* knowledge about the subject of the research helped to eliminate possible intentional or unintentional data distortion by the experimental subjects. The results showed that personality profiles of women with latent toxoplasmosis significantly differed from those of women without the infection. The character and the amount of these personality changes were similar to those observed in the previous studies.

#### MATERIALS AND METHODS

**Experimental subjects.** The experimental set (young women) consisted of 191 women 18-39 years old who were serologically tested for toxoplasmosis during pregnancy. Among these women 136 tested negative and 55 had the titres of anti-

Toxoplasma antibodies between 1:8 and 1:64 which indicates latent toxoplasmosis. According to the levels of IgG and IgM antibodies no woman involved suffered acute toxoplasmosis. All subjects (824) received the personality questionnaire 6-7 months after the serological examination, mostly 2-5 months after the childbirth. In the cover letter they were only asked to voluntarily participate in an unspecified research project (to avoid a possible fear of young mothers of congenital toxoplasmosis). The return rate was approximately 23.2%.

Personality tests. Cattell's sixteen factor questionnaire (form A) (Cattell 1970) was used for characterisation of personality profiles. This questionnaire is still widely used for personality studies in many countries, including the Czech Republic. It covers sixteen personality factors (Table 1). The main advantage of this traditional questionnaire is that it contains only one hundred and eighty-seven (187) questions. Therefore, it can be completed by most subjects within forty minutes. All subjects obtained the questionnaire by mail and were asked to voluntarily participate in an unspecified research project.

Immunological tests for toxoplasmosis. The latent toxoplasmosis in pregnant women was diagnosed with complement-fixation test (CFT), (Warren and Sabin 1942), in the National Diagnostic Laboratory for Toxoplasmosis, National Institute of Public Health, Prague, Czech Republic. The titre of anti-Toxoplasma antibodies in sera was measured in dilutions between 1:8 and 1:1024. All subjects with the titres from 1:8 to 1:64 were considered latent toxoplasmosis positive, the subject with titre lower then 1:8 latent toxoplasmosis negative. No subjects with titres equal or higher than 1:128 (the indication of recent acute toxoplasmosis) were involved in our study.

Statistical tests. The computation of standard residuals by linear regression (for elimination of the effect of age), the t-tests and the Levene tests were performed by the program package Statistica. The program TREEPT, originally called PTPT (Flegr and Záboj 1997), was used for all permutation tests including the permutation tests with elimination of false negative cases. This freeware Windows programme for various permutation tail probability tests is available on the address http://www.natur.cuni.cz/~flegr/programs. The rationale of the permutation test is described in the legend of Table 3. Differences in variances of personality factors between Toxoplasma-negative and Toxoplasma-positive subjects were tested by the same permutation tests, however, the absolute values of age-standardized factors instead of age-standardised factors were entered into the program. The rationale of this approach was based on our previous observation that for certain personality factors both minimal and maximal extremes were absent in Toxoplasma-positive subjects.

### RESULTS

We obtained the personality data from 136 *Toxoplasma*-negative and 55 *Toxoplasma*-positive women. Effect of the age was eliminated be computation of standard residual of linear regression between the age and raw personality factors. The effect of latent toxoplasmosis was tested by t-test and the Levene's test for the homogeneity of variances. The results (Table 2) suggest that no differences in mean values of Cattell's factors exist

between Toxoplasma-negative and Toxoplasma-positive women.

The diagnosis of latent toxoplasmosis was performed by complement fixation test. This simple serological test is very useful for rapid screening for acute toxoplasmosis. However, it often provides false negative results for subjects with low titres of antibodies, for example for subjects with very old T. gondii infections. Therefore, our subset of Toxoplasma-negative women must be contaminated by an unknown percentage of Toxoplasmapositive women. Such women with old infections (and consequently with low levels of specific antibodies) are known to have the highest amount of the personality factor shift (Flegr and Hrdý 1994, Flegr et al. 1996). Therefore, their presence in the Toxoplasma-negative subsets can flatten out any possible difference between the two subsets. The effects of such kind of data contamination can be revealed and eliminated by a permutation test with data reassignment. For this kind of tests we used the program TREEPT which can perform permutation tests analogical to t-test or ANOVA with reassignment of a particular fraction of cases from one group to the other (in this case from the subset of Toxoplasma-negative women to the Toxoplasma-positive subset). The results of this analysis are shown in the Table 3. The results of permutation tests showed that the Toxoplasma-positive women had higher factor B (intelligence), factor O (guilt proneness) and possibly also Q4 (high ergic tension) and Q1 (radicalism). The differences between Toxoplasma-negative and Toxoplasma-positive women in the factors L and O3. and possibly also F and N demonstrated its existence by changes in the character of distribution, rather than changes in the mean values of the factors. The histograms of standardised factors (Fig. 1) suggest that the distribution of values of these factors in Toxoplasma-positive women have lower kurtosis and can be approx-imated by the rectangular while that of Toxoplasma-negative women by the normal distribution.

### DISCUSSION

Our results suggest that the personality profiles of women with latent toxoplasmosis differ in certain factors from those of women without toxoplasmosis. *Toxoplasma*-positive women have higher intelligence, guilt proneness (are more apprehensive, self-reproaching, insecure), possibly also higher ergic tension and radicalism. The changes in other personality factors were manifested by difference in distribution of values of the factor rather than in differences in the mean values of the factor.

These changes, however, cannot be observed directly when the women are sorted into the *Toxoplasma*-negative and positive groups on the basis of routine serological tests (like CFT). These tests are optimised for early diagnosis of acute toxoplasmosis. They are able to prove the existence of high titres of specific anti-*Toxoplasma* 

**Table 2.** Differences in 16 Cattell's factors between *Toxoplasma*-positive and *Toxoplasma*-negative women. The factors are described in Table 1.

	Mean	Mean	t separ.	df	p	Levene	р
	Toxo-negat.	Toxo-posit.	var.est.			F(1,189)	Levene
A	12.79	12.80	-0.17	92.51	0.87	0.28	0.60
В	8.32	8.93	-1.80	125.90	0.07	2.65	0.11
C	14.07	13.56	0.99	97.35	0.33	0.40	0.53
E	11.88	12.11	-0.29	93.03	0.77	0.59	0.44
F	13.08	12.87	0.07	88.02	0.95	2.06	0.15
G	11.91	12.00	-0.04	112.59	0.97	0.77	0.38
H	12.31	12.67	-0.42	104.24	0.68	0.14	0.70
I	12.74	12.35	0.69	103.82	0.49	0.38	0.54
L	10.00	10.44	-0.94	87.17	0.35	2.82	0.10
M	11.09	11.05	0.39	111.28	0.70	1.24	0.27
N	10.22	10.29	-0.02	86.95	0.99	2.52	0.11
O	10.75	11.73	-1.89	100.65	0.06	0.19	0.66
Q1	5.76	6.25	-1.15	103.72	0.25	0.00	0.97
Q2	11.45	11.24	0.49	107.12	0.62	0.09	0.77
Q3	11.43	11.55	-0.09	88.05	0.93	3.94	0.05
Q4	13.99	14.84	-1.37	105.26	0.17	0.52	0.47

Table 3. The differences between *Toxoplasma*-negative and *Toxoplasma*-positive women tested by permutation tail probability test with data reassignation. The differences in particular age-standardised personality factor was tested by two-sided permutation test. From zero to 60% of subjects with the highest (or lowest) values of a tested factor were relocated from the group of 136 *Toxoplasma*-negative women to the second group of 55 *Toxoplasma*-positive women. Then, the difference of means of these two groups was calculated. In next 2999 steps the empirical values of analysed factors were arbitrary assigned into two groups of the sizes 136 and 55 cases, the same percentage as before of highest (or lowest) cases from the larger group was relocated to smaller group and the difference of the means of the two groups was calculated. Finally, all 3000 differences (including the one calculated from nonpermutated data) were sorted from highest to lowest. The percentage of differences higher or equal to that calculated on the basis of nonpermutated data was considered as the statistical significance (p), i.e. the probability of obtaining the same or higher difference of means of groups 136 and 55 subjects assigned into these groups randomly. The difference in data distribution within the group was tested by the same approach. In this case, however, the absolute values of age-standardised factors instead of standardised factors were entered into the test.

	Means							Distributions	
	p(0%)	p(5%)	p(10%)	p(15%)	p(20%)	p(30%)	p(60%)	p(5%)	p(10%)
A	0.90	0.46	0.44	0.42	0.44	0.50	0.53	0.39	0.41
В	0.12	0.05	0.03	0.03	0.02	0.03	0.04	0.84	0.72
C	0.33	0.81	0.18	0.79	0.80	0.82	0.89	0.26	0.21
Е	0.75	0.37	0.39	0.42	0.42	0.41	0.54	0.31	0.32
F	0.92	0.53	0.50	0.59	0.65	0.73	0.81	0.08	0.09
G	0.97	0.46	0.47	0.45	0.46	0.44	0.36	0.81	0.80
Н	0.72	0.36	0.32	0.31	0.28	0.30	0.37	0.61	0.61
I	0.50	0.72	0.28	0.64	0.63	0.57	0.56	0.74	0.74
L	0.28	0.16	0.18	0.19	0.23	0.33	0.55	0.04	0.05
M	0.61	0.64	0.30	0.58	0.56	0.55	0.44	0.84	0.84
N	0.98	0.56	0.60	0.61	0.61	0.64	0.74	0.06	0.09
0	0.06	0.02	0.02	0.03	0.03	0.05	0.09	0.32	0.28
Q1	0.26	0.09	0.09	0.09	0.09	0.09	0.17	0.43	0.41
Q2	0.64	0.68	0.29	0.63	0.63	0.66	0.63	0.51	0.44
Q3	0.93	0.47	0.49	0.55	0.58	0.68	0.74	0.01	0.01
Q4	0.20	0.10	0.09	0.07	0.07	0.07	0.09	0.67	0.68

antibodies which are characteristic for patients with an acute phase of acquired toxoplasmosis. In most of clinical laboratories the samples of sera are either measured from the titre 1:8 or the subjects with the titres less then 1:8 are considered as *Toxoplasma*-negative (and are excluded from further analysis by more specific tests, for example

by IgM ELISA). The women in our experimental set were screened for toxoplasmosis during routine clinical examination in pregnancy. Their sera were tested by complement fixation test (Warren and Sabin 1942) and the subjects without positive reaction in 1:8 dilution were considered as *Toxoplasma*-negative. The clinical records

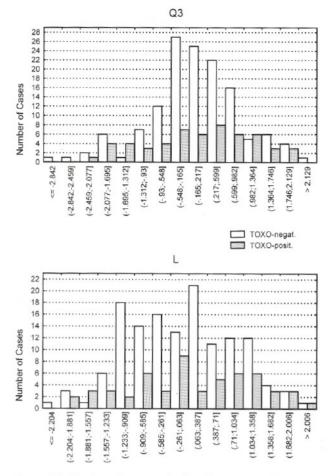


Fig. 1. Distribution of age-standardised factors L and Q3.

as well as the comparison of results of CFT and IgG ELISA tests suggest that the titres of specific antibodies estimated by CFR can decrease below 1:8 within a few years after the end of acute toxoplasmosis. Therefore, our group of 136 Toxoplasma-negative women must be contaminated by an unknown percentage of women with old Toxoplasma gondii infection. The results of our observation of toxoplasmosis patients during the interval of 13 years after infection (Flegr et al. 1996) showed that at least some of the Toxoplasma gondii-induced personality changes are very slow. Therefore, the personality of subjects with longest infections (who might be often misdiagnosed as Toxoplasma-negative) can express the most prominent changes in our experimental set. Even a small percentage of such subjects in the Toxoplasma-negative set can flatten out the possible differences between the two subsets of women. In the first study (Flegr and Hrdý 1994) in which prominent differences between Toxoplasma-negative and Toxoplasma-positive were observed, the toxoplasmin intradermal delayed hypersensitivity test (Feldman 1954) was used for diagnosis of latent toxoplasmosis. This old test monitors cellular rather than humoral immunity. The

clinical experience shows that the positive reactions in the intradermal delayed hypersensitivity test persists for a very long time, probably for the whole life of an infected person (Remington and Krahenbuhl 1982). When the toxoplasmosis is diagnosed with a common serological test like CFT the presence of unknown percentage of false negatives must be anticipated and the existence of any differences between *Toxoplasma*-negative and *Toxoplasma*-positive subjects must be tested by proper statistical method (such as the permutation test with data reassignation).

In our study the permutation tests revealed the existence of differences in mean values of factors B, O, possibly also O4 and O1. In the factors F. L. N and O3 the latent toxoplasmosis affected the distribution of personality factor's values, rather then their mean values. The lover kurtosis of the distribution as well as the absence of distinct peak in the histogram of the distribution can be explained by the fact that the Toxoplasma-induced personality shift is probably a long-lasting process. Our results (Flegr et al. 1996) suggest that the process is triggered by the infection, however, it takes many years to fully evolve. Consequently, the women with a relatively fresh infection are Toxoplasma-positive on the basis of serological tests, however, their personality profile has still the Toxoplasma-negative character. As our experimental set contains the women with toxoplasmosis of various length no distinct peak but rather a flat plateau can be observed on the histograms of personality factors of Toxoplasma-positive women. As discussed above, the absence of significant difference of mean values of the factor can be explained by misdiagnosis of the most changed subjects (with lowest antibody titres) as Toxoplasma-negative.

The results of present analysis of the set of young women slightly differ from those obtained on the set of students and teachers of biology of the Charles University (Flegr and Hrdý 1994, Flegr et al. 1996). They are, however, similar to those obtained on the set of acute toxoplasmosis patients and on the subset of 55 Toxoplasma-positive women (submitted). In the set of biologists the Toxoplasma-positive women had higher A (affectothymia), Q3 (strength of self sentiment), and lower L (protension). In the set of 230 women with clinical history of acute toxoplasmosis the duration of infection positively correlated with G (superego strength), and Q3 (strength of self-sentiment) while in the set of 55 young women with latent toxoplasmosis the length of infection (estimated on the basis of antibody titres) positively correlated with A (affectothymia), F (surgence), G (superego strength), H (parmia) and L (protension). The differences between the results obtained on particular experimental sets can be probably explained by special nature of some of these sets. Especially the biologists represent a highly atypical population sample which differs in many personality traits from a general

population. Despite the differences a similar set of personality factors (usually incorporating the factors A, F, G, L, O, Q3 and Q4) seems to by influenced by latent toxoplasmosis in all examined sets of women. The results of discrimination analysis and of logistic regression (Flegr and Hrdý 1994, Flegr et al. 1996) suggest that only some of these factors are influenced by the parasite directly while the others are changed because of their correlation with other (toxoplasmosis influenced) factors. As the correlation matrices of Cattell's factors often differ among various social and professional groups, the changes in secondary-influenced factors might vary among different experimental sets.

The results of the current study are very important in two respects. First, in the comparison with all previous studies the experimental set represents a homogeneous sample of normal population of young women. The experimental set of biologists (Flegr and Hrdý 1994) represents highly untypical population of university students and teachers. The second experimental set (Flegr et al. 1996) is a relatively good representation of Czech population. However, the age-structure of the sample is highly heterogeneous. Moreover, all men in this sample experienced relatively severe acute toxoplasmosis disease (in contrast with most of other infected people they were diagnosed for acquired toxoplasmosis and received a medical help). In both cases it is not clear whether the results and conclusions can be extrapolated for normal population.

Secondly, in all previous studies the subjects were informed about the aim of the projects before the experiment. In this study neither toxoplasmosis nor *T. gondii* were mentioned in the cover letter (to avoid possible fear of young mothers of congenital toxoplasmosis). For the first time we fully excluded any possible intentional or unintentional distortion of data by the experimental subjects.

It is not possible to decide on the basis of present data whether the toxoplasmosis induces the changes in personality or whether the certain combination of the personality traits increases the probability of contact with T. gondii infection. There are, however, several reasons to believe that former possibility is the correct one: (1) Significant positive correlation exists between the duration of latent toxoplasmosis and the amount of personality shift in the sets of men (Flegr et al. 1996) and women (submitted) diagnosed with acute toxoplasmosis during the period of 13 and 14 years, respectively in various Prague hospitals. (2) Significant negative correlation exists between the level of anti-Toxoplasma antibodies (which indirectly reflects the duration of latent toxoplasmosis) and the amount of personality shift in the sets of biology students (Flegr and Hrdý 1994) and Toxoplasma-positive young women (submitted). (3) No correlation exists between the major toxoplasmosis-risk factors (consumption of raw meat and contact with cat) and the personality factors influenced by toxoplasmosis (Flegr et al. 1998). (4) All but one (factor O) personality factors influenced by toxoplasmosis are shifted in the opposite direction in Toxoplasma-positive men and women. It is difficult to explain how the oppositely changed factors can promote the same effect in men and women, namely the increase of probability of T. gondii infection.

It can be only speculated whether the induction of personality changes in men is connected with an attempt of the parasite to manipulate the host behaviour or whether it is only a byproduct of a non-specific decline in the quality of host life (connected, for example, with *Toxo-plasma*-induced general immunosuppression).

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## REFERENCES

- ARNOTT M.A., CASSELLA J.P., AITKEN P.P., HAY J. 1990: Social interactions of mice with congenital *Toxoplasma infection*. Ann. Trop. Med. Parasitol. 84: 149-156.
- BERDOY M., WEBSTER J.P., MACDONALD D.W. 1995: Parasite-altered behaviour: is the effect of *Toxoplasma gondii* on *Rattus norvegicus* specific? Parasitology 111: 403-409.
- CATTELL R.B. 1970: Handbook for the Sixteen Personality Factors Questionnaire (16PF). Institute for Personality and Ability Testing, Champain, 115 pp.
- FELDMAN H.A. 1954: Laboratory methods in current use for the study of toxoplasmosis. Bibl. Ophthalmol. 39: 1-11.
- FLEGR J., HRDÁ Š., TACHEZY J. 1998: The role of psychological factors in questionnaire-based studies on routes of human toxoplasmosis transmission. Centr. Eur. J. Public Health 6: 45-50.

- FLEGR J., HRDÝ I. 1994: Influence of chronic toxoplasmosis on some human personality factors. Folia Parasitol. 41: 122-126.
- FLEGR J., ZÁBOJ P. 1997: PTPT, the freeware program for permutation testing concordance between phylogeny and the distribution of phenetic traits. Acta Soc. Zool. Bohem. 61: 91-95.
- FLEGR J., ZITKOVÁ S., KODYM P., FRYNTA D. 1996: Induction of changes in human behaviour by the parasitic protozoan *Toxoplasma gondii*. Parasitology 113: 49-54.
- HEITMAN B.B., IRIZARRY A.F. 1997: Recognition and management of toxoplasmosis. Nurse. Pract. 22: 79-86.
- HUTCHISON W.M., AITKEN P.P., WELLS B.W.P. 1980a: Chronic *Toxoplasma* infection and familiarity-novelty discrimination in the mouse. Ann. Trop. Med. Parasitol. 74: 145-150.

- HUTCHISON W.M., BRADLEY M., CHEYNE W.M., WELLS B.W.P., HAY, J. 1980b: Behavioural abnormalities in *Toxoplasma*-infected mice. Ann. Trop. Med. Parasitol. 74: 337-345.
- KOPPE J.G., ROTHOVA A. 1989: Congenital toxoplasmosis. A long-term follow-up of 20 years. Int. Ophthalmol. 13: 387-390.
- KRAMER W. 1966: Frontiers of neurological diagnosis in acquired toxoplasmosis. Psychiatr. Neurol. Neurochir. 69: 43-64.
- MOCSNY N. 1992: Toxoplasmic encephalitis in the AIDS patient. J. Neurosci. Nurs. 24: 30-33.
- NĚMEC R., NEČAS S. HOLKOVÁ R., ČATÁR G., PORUBENOVÁ. A. 1988: Titre of toxoplasmic antibodies in workers with potentially higher probability of toxoplasmic infection. Bratisl. Lek. Listy 89: 779-783.
- REMINGTON J.S., KRAHENBUHL J.L. 1982: Immunology of Toxoplasma gondii. In: A.J. Nahmias, J. O'Reilly (Eds.), Immunology of Human Infection, Part II. Plenum Publishing Corporation, New York, pp. 327-371.
- SENGBUSCH H.G., SENGBUSCH L.A. 1978: *Toxoplasma* antibody prevalence in veterinary personnel and a selected population not exposed to cats. Am. J. Epidemiol. 103: 595-597.

- STIBBS H.H. 1985: Changes in brain concentrations of catecholamines and indoleamines in *Toxoplasma gondii* infected mice. Ann. Trop. Med. Parasitol. 79: 153-157.
- VARELA G., PALENCIA L., VAZQUEZ A. 1957: Utilization del pez Lebistes reticulatus (guppy) en el diagnostico de la toxoplasmosis. Rev. Inst. Salubr. Enferm. Trop. 17: 75-80.
- VARELA G., VAZQUEZ A., TARROELLA J. 1956: Probable existencia de la dietilamida del acido D-lisergico en la infeccion por *Toxoplasma gondii*. Rev. Inst. Salubr. Enferm. Trop. 16: 29-32.
- WARREN J., SABIN A.B. 1942: The complement fixation reaction in toxoplasmic infection. Proc. Soc. Exp. Biol. Med. 51: 11-16.
- WEBSTER J.P. 1994: The effect of *Toxoplasma gondii* and other parasites on activity levels in wild and hybrid *Rattus norvegicus*. Parasitology 109: 583-589.
- WEBSTER J.P., BRUNTON C.F.A., MACDONALD D.W. 1994: Effect of *Toxoplasma gondii* upon neophobic behaviour in wild brown rats, *Rattus norvegicus*. Parasitology 109: 37-43.
- YAMAOKA M., KONISHI E. 1993: Prevalence of antibody to Toxoplasma gondii among inhabitants under different geographical and climatic conditions in Hyogo Prefecture, Japan. Jpn. J. Med. Sci. Biol. 46: 121-129.