Infection with *Toxoplasma gondii* can promote chronic liver diseases in HIV-infected individuals

Ihor H. Hryzhak

Ivano-Frankivsk National Medical University (Ivano-Frankivsk, Ukraine); Ivano-Frankivsk Regional Center for AIDS Prevention and Fight (Ivano-Frankivsk, Ukraine)

Abstract: Liver pathologies and infection with *Toxoplasma gondii* (Nicolle et Manceaux, 1908) are widespread among HIV-infected patients. However, a possible contribution of toxoplasmosis to the development of various forms of liver diseases in HIV-infected individuals has not yet been determined. This research is a retrospective cohort study. Medical cards of 907 HIV-positive patients, including 119 individuals who died, were studied. The patients were divided into two groups: 531 patients were seropositive to *T. gondii* and 376 seronegative. General liver pathology was more widespread among patients seropositive to *T. gondii* than in seronegative patients (63.1 ± 2.1% and 51.9 ± 2.6%, respectively, p < 0.001). The association of seropositive to *T. gondii* with general liver pathology is weak both in the whole cohort (Pearson’s contingency coefficient C = 0.112), and among the deceased patients (C = 0.228). Chronic HBV-HCV coinfection was more common in the seropositive than in seronegative individuals as it was found both in entire cohorts (26.0 ± 1.9% and 18.6 ± 2.0%, respectively, p = 0.010) and in died patients (31.0 ± 5.5% and 14.6 ± 5.1%, respectively, p = 0.041). *Toxoplasma gondii* had a weak role in distributing of HBV-HCV coinfection between cohorts (C = 0.187). In both cohorts in patients with chronic hepatitis, regardless of its etiology, there was no significant difference in alanine transaminase activity (ALT). Cirrhosis of the liver occurred 4.5 times more often in deceased seropositive patients than in the entire seropositive cohort (23.9 ± 5.1 and 5.3 ± 2.0, respectively, p = 0.0006) whereas it no significantly increased in seronegative cohort (10.4 ± 4.4 against 4.8 ± 1.1, p > 0.05). In them *T. gondii* is weakly involved in cirrhosis formation (C = 0.168). Thus, in HIV-infected patients, *T. gondii* is a weak nonspecific adjunct that supports chronic liver inflammation and progression of cirrhosis, regardless of etiology, but does not influence the degree of hepatitis activity. The increased prevalence of HBV-HCV coinfection in patients seropositive for *T. gondii* may be related to their risk factor behaviour associated with uncontrolled blood contacts.

Keywords: *Toxoplasma gondii*, HIV, infection, liver

*Toxoplasma gondii* (Nicolle et Manceaux, 1908) is an obligate intracellular parasitic protozoan eukaryote (an apicomplexan) that causes the infection disease toxoplasmosis (Dardé et al. 2011). It is estimated to affect at least one-third of the planet’s population (Montoya and Liesenfeld 2004). This infection is generally asymptomatic but is associated with the ‘burden’ of different non-infectious human diseases (somatic, psychiatric, oncological and others) (Jones et al. 2001, Flegr et al. 2014, El-Rehim El-Henawy et al. 2015).

There are a few studies of toxoplasmic hepatitis in immunocompetent patients, which is accompanied by hepatomegaly, increasing rates of aspartate transaminase activity (AST), alanine transaminase activity (ALT), alkaline phosphatase activity (ALP) and gamma glutamyl transpeptidase activity (GGT), normal or increased levels of total bilirubin (Doğan et al. 2007, Atilla et al. 2015) and sometimes with symptoms of generalised toxoplasmosis (skin rash, retinochoroiditis, myositis) (Nunura et al. 2010). In many studies, it has been shown that chronic or latent toxoplasmosis makes a negative contribution to various liver diseases. For example, in Hangzhou City (China) the seropositive rate of anti-*T. gondii* IgG antibody among 1,200 patients with liver diseases was 24.0% but it was only 11.42% in healthy people. However, *T. gondii* seroprevalence did not depend on the exact diagnosis of these patients (liver cancer, hepatitis B virus infection, liver fibrosis, fatty degeneration of liver) (Jian-Feng et al. 2018). In non-pregnant healthy women with IgG to *T. gondii* there were higher levels of AST, ALT, ALP and GGT in blood serum compared with the control group of women (Nadwa and Muthear 2012).

On the other hand, chronic viral hepatitis promotes the activation of toxoplasmosis. Thus, the active co-infection with *T. gondii* was found in 33.3% of patients with chronic hepatitis B virus (HBV) infection and in 31.4% of patients with chronic hepatitis C (HCV) infection. It is suggested that the reactivation of latent infection with *T. gondii*...
Toxoplasma gondii occurs in patients with chronic liver diseases much more frequently (30%) than in patients without it (6%). Furthermore, the high HCV viral load in patients with chronic HCV infection is significantly correlated with parasitemia of T. gondii (Nagwa et al. 2016) and high levels of anti-T. gondii IgG, but a correlation between the stage of liver fibrosis and the presence of antitoxoplasmic-IgM or IgG antibodies has not been found (El-Nahas et al. 2014). However, other researchers think that there is an association between seropositive to T. gondii and fibrosis, inflammation and necrotic process in the liver of patients with chronic HCV infection. In these patients the albumin level, international normalised ratio and platelet count were significantly decreased (El-Rehim El-Henawy et al. 2015). The incidence of non-alcoholic fatty liver disease was higher in individuals seropositive to T. gondii than in seronegative individuals (27.1% and 23.4%, respectively, p < 0.001). However, multivariant analysis revealed that infection with T. gondii was not an independent risk factor in liver diseases (Huang et al. 2018). A study performed among a northern Mexican population found that both seroprevalence of T. gondii and level of specific IgG did not differ significantly between patients with chronic liver diseases (13.3%) and the control group (10.7%). Therefore, researchers concluded that there is no association between seropositivity to T. gondii and any diagnosis of liver disease (Alvarado-Esquivel et al. 2011).

Immunocompromised individuals who are HIV-infected have a high risk of toxoplasmosis manifestation. Typically, these patients might have brain lesions, such as toxoplastic encephalitis and myelopathy (Chaddha et al. 1999, Kung et al. 2011, García-García et al. 2015, Marra 2018) and the rate of cerebral toxoplasmosis in the general HIV-positive population is 3.5% (Gofta et al. 2012). However, the burden of infection with T. gondii in HIV-infected patients is geographically variable and usually follows the prevalence of T. gondii in the general population (Veera-noot 2017, Vidal 2019). A low level of CD4 cells (< 100/ mm³) and a positive T. gondii serology are likely to predict the occurrence of toxoplastic encephalitis within the following 12 months in 25.4% patients (Oksenhendler et al. 1994).

Extracerebral toxoplasmosis occurs in patients with more severe immunosuppression and a CD4+ lymphocyte mean count of 57/mm³. Its localisation includes eyes (50% of patients), lungs (26%), heart and pericardium (3%), bone marrow (3%), blood-(parasitemia with acute febrile syndrome) (3%) and bladder (1%). There were also isolated cases of toxoplasmosis infection of nasopharynx, skin, liver, lymph nodes and conus medullaris. Disseminated toxoplasmosis (at least two extracerebral visceral localisations) occurred in 11.5% of patients (Rabaud et al. 1994). There was a report about a patient with the AIDS and watery diarrhea due to duodenitis, which was associated with toxoplastic hepatitis, elevated serum ALP and lactate dehydrogenase activity (Bonacini et al. 1996).

In general, various liver diseases among HIV-infected patients are very common (Rodriguez-Mendez 2000). In the USA HBV infection (70–90%) and HCV infection (88%) were diagnosed in intravenous drug users (CDC 2017). Also, a wide spectrum of liver pathologies is found in deceased HIV-infected patients, such as chronic portal hepatitis (80.0%), alcoholic hepatitis (23.7%), fatty hepatitis (7.5%), fatty degeneration of the liver (73.7%), and liver cirrhosis (36.3%) (Lytvynenko 2007). Various degrees of fibrosis and hepatocyte dysfunction occur in HIV-infected individuals and contribute significantly to the mortality rate—which is 1.7% per 100 man-years (Quirishi et al. 2003, Castellares et al. 2008). It may be assumed that many opportunistic infections are reactivated before the death of HIV-infected patients. Cytomegalovirus (CMV), mycobacterium and T. gondii were found more frequently in autopsy liver tissues of HIV-positive patients than in biopsy samples of live patients (Trojan et al. 1998, Lang et al. 2005). Therefore, different liver pathologies and T. gondii might be combined in many patients. However, the consequences of interaction between them are not clear.

MATERIALS AND METHODS

Design of the study. A three-year retrospective cohort study was conducted at the Department of Infectious Diseases and Epidemiology of Ivano-Frankivsk National Medical University and at the Ivano-Frankivsk Regional Center for AIDS Prevention and Fight (Ukraine). It is part of a wider study “The course of infectious diseases on the background of concomitant pathology, combined chronic infections and invasions, correction of treatment”, state registration number: 0112U005012 (2019–2023).

The medical data from 2013–2015 for 907 HIV-infected patients (men 58.3% with a mean age of 35.21 ± 0.26 years (18–65 years) were studied. IgG and IgM antibodies to Toxoplasma gondii and markers for hepatitis B and C (HBsAg, anti-HBc IgM and IgG, anti-HCV total Ig) in the blood were determined by enzyme-linked immunosorbent assay (test systems produced by DIAGPROF-MED, Ukraine). Cytomegalovirus (CMV) and Epstein-Barr virus (EBV) DNA were determined by PCR.

According to the test results for IgM and IgG antibodies to T. gondii, the patients were divided into two cohorts: 531 seropositive individuals and 376 seronegative ones. Criteria for inclusion in the seropositive cohort were the presence of anti-Toxoplasma-IgG or IgM or both IgG and IgM in the blood at the first test, regardless of the level of these antibodies. Some previously seronegative individuals were transferred from the seronegative to the seropositive cohort after the appearance of seroconversion (the appearance of specific IgM or IgG, or both IgG and IgM simultaneously). Patients who did not have IgM and IgG antibodies to T. gondii at the first test or during the next 3 years were included into the seronegative cohort. In both cohorts, patients with hepatitis C or HBV-HCV coinfection did not receive antiviral treatment for HCV infection for that period.

Statistical analysis. In both cohorts the percentage of patients with different liver diseases and their average error (P ± m) were estimated. Significance of difference in the frequency of liver pathology in the two cohorts was determined using the Chi-square
test (Pearson’s χ² test). The relationship between seropositive status for toxoplasmosis and specific liver pathology was assessed with the Pearson contingency coefficient (C). Significance of difference between the incidence of liver pathology among patients who died in a cohort, and the frequency in the whole cohort, was assessed by Student’s t-test.

Our study was approved by the Ethics Commission of Ivanovo-Frankivsk National Medical University (Expert Decision No. 96/17 dated 24.05.2017).

RESULTS

A total of 531 individuals (61.4% men and 38.6% women) were included in the cohort of patients-seropositive to Toxoplasma gondii and 376 individuals (53.9% men and 46.2% women) were included in the seronegative cohort. In the seropositive cohort 91 (17.1 ± 1.6%) patients had stage I HIV infection, 101 (19.0 ± 1.7%) had stage II, 80 (15.07 ± 1.55%) had stage III, and 259 (48.8 ± 2.2%) had stage IV. In the seronegative cohort 48 (12.8 ± 1.7%, p > 0.05) patients had stage I of HIV-infection, 78 (20.7 ± 2.1%, p > 0.05) had stage II, 76 (20.2 ± 2.1%, p > 0.05) had stage III, and 174 (46.3 ± 2.6%, p > 0.05%) patients had stage IV. Therefore, the distribution of patients regarding the stage of HIV-infection was similar in both cohorts.

Antiretroviral therapy (ART) was received by 352 patients in the seropositive cohort (66.3 ± 2.05%) and by 287 patients in the seronegative cohort (76.3 ± 2.2%, p = 0.002).

Sulfamethoxazole/trimethoprim (400/800 mg twice daily) for the prevention of Pneumocystis pneumonia was given to 134 patients (25.2 ± 1.9%) in the seropositive cohort, and 88 patients (23.4 ± 2.2%, p > 0.05) in the seronegative cohort. Azithromycin (1,200 mg once a week) for the prevention of the Mycobacterium avium complex infection was given to 50 patients (9.4 ± 1.3%) in the seropositive cohort and 34 patients (9.0 ± 1.5%, p > 0.05) in the seronegative cohort. Therefore, sulfamethoxazole/trimethoprim and azithromycin were taken by the same proportion of patients in these two cohorts.

In the seropositive cohort 58 patients (10.9 ± 1.4%) had signs of active Toxoplasma gondii infection and 51 patients had signs of activation of endogenous Toxoplasma gondii infection (either the titre of IgG increasing by more than twice (46 patients) or the appearance of IgM following the previous presence of IgG (five patients). Seven patients had primary Toxoplasma gondii infection with seroconversion, either the appearance of IgM followed by the appearance of IgG after 3–4 weeks (one person) or the initial appearance of IgG without IgM (six people).

There were 24 patients seropositive to Toxoplasma gondii who had a diagnosis of toxoplasmic encephalitis. In addition, toxoplasmosis of the brain was diagnosed in three patients of the seronegative cohort because of the primary infection. Antibodies to Toxoplasma gondii subsequently appeared in two of the patients who were moved to the seropositive cohort. The third patient remained seronegative without any specific antibodies in the blood until her death 15 days later (the patient remained in the seronegative cohort).

There were 49 people with serological signs of active toxoplasmosis. They had a moderate increase in liver and spleen (hepatolienal syndrome) which significantly exceeded the frequency of symptoms in the whole seropositive cohort (84.48 ± 4.75% against 63.1 ± 2.1%, t = 4.1, p < 0.001). The average level of ALT in the group of patients with serological signs of activity was insignificantly higher than the level in the entire cohort at the initial examination (102.2 ± 11.4% against 77.3 ± 9.8%, t = 1.7, p = 0.0986).

Diseases (hepatitis and cirrhosis from every cause) were observed in 530 out of 907 patients examined (58.4 ± 1.6%). However, according to the data presented in Table 1, the frequency of liver pathologies was distributed unequally between the two cohorts with more among seropositive patients (63.1 ± 2.1%) than among seronegative ones (51.9 ± 2.6%), p < 0.001. The total number of patients with chronic hepatitis from all etiological factors (HBV infection, HCV infection, HBV-HCV coinfection, herpesviruses, toxic and cryptogenic origin) was higher in the seropositive cohort than in the seronegative cohort (57.8 ± 2.14% and 47.1 ± 2.6%, respectively, p = 0.002) but only chronic HBV-HCV

Table 1. The distribution of liver diseases in cohorts of HIV-infected individuals with and without antibody to Toxoplasma gondii (Nicolle et Manceaux, 1908).

<table>
<thead>
<tr>
<th>Disease</th>
<th>T. gondii positive (N = 531)</th>
<th>T. gondii negative (N = 376)</th>
<th>Chi-square test</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>P ± m ± %</td>
<td>Number</td>
<td>P ± m ± %</td>
</tr>
<tr>
<td>Chronic HBV infection</td>
<td>43</td>
<td>8.1 ± 1.2</td>
<td>25</td>
</tr>
<tr>
<td>Chronic HCV infection</td>
<td>50</td>
<td>9.4 ± 1.3</td>
<td>42</td>
</tr>
<tr>
<td>Chronic HBV-HCV coinfection</td>
<td>138*</td>
<td>26.0 ± 1.9</td>
<td>70</td>
</tr>
<tr>
<td>Chronic toxonic hepatitis</td>
<td>8</td>
<td>1.5 ± 0.5</td>
<td>1</td>
</tr>
<tr>
<td>Chronic cryptogenic hepatitis</td>
<td>63</td>
<td>11.9 ± 1.4</td>
<td>35</td>
</tr>
<tr>
<td>Chronic EBV-hepatitis</td>
<td>2</td>
<td>0.4 ± 0.3</td>
<td>3</td>
</tr>
<tr>
<td>Chronic CMV-hepatitis</td>
<td>3</td>
<td>0.6 ± 0.3</td>
<td>1</td>
</tr>
<tr>
<td>Total number of patients with chronic hepatitis</td>
<td>307*</td>
<td>57.8 ± 2.1</td>
<td>177</td>
</tr>
<tr>
<td>Liver cirrhosis, class A</td>
<td>12</td>
<td>2.3 ± 0.6</td>
<td>6</td>
</tr>
<tr>
<td>Liver cirrhosis, class B</td>
<td>8</td>
<td>1.5 ± 0.5</td>
<td>8</td>
</tr>
<tr>
<td>Liver cirrhosis, class C</td>
<td>8</td>
<td>1.5 ± 0.5</td>
<td>4</td>
</tr>
<tr>
<td>Total number of liver cirrhosis</td>
<td>28</td>
<td>5.3 ± 0.2</td>
<td>18</td>
</tr>
<tr>
<td>General liver pathology</td>
<td>335*</td>
<td>63.1 ± 2.1</td>
<td>195</td>
</tr>
</tbody>
</table>

* – the difference between the indexes in cohorts of patients with and without T. gondii is statistically significant; χ² - Pearson’s chi-squared test; C - Pearson’s contingency coefficient; P – percentage; m ± – standard error of percentage; p – statistical significance.
However, there were weak associations between infection with *T. gondii* and the total number of chronic hepatitis cases (C = 0.106) and the total number of patients with liver diseases (C = 0.112). The frequency of liver cirrhosis in both cohorts did not differ (see Table 1).

According to ALT activity at initial examination in 307 seropositive patients with chronic hepatitis from any cause there was minimal activity of hepatitis in 275 patients (89.58 ± 1.74%), mild activity – in 25 (8.1 ± 1.6%) and moderate activity in seven patients (2.6 ± 0.9%). At the same time, among 177 seronegative patients with chronic hepatitis from any cause, there was minimal activity in 167 individuals (94.4 ± 1.7%), mild activity in nine patients (5.1 ± 1.7%) and moderate activity in one patient (0.56 ± 0.56%). Therefore, there was no significant difference in hepatitis activity between these two cohorts (p > 0.05).

A total of 119 patients died during the three years, 71 from the seropositive cohort (13.4 ± 1.5%) and 48 from the seronegative cohort (12.8 ± 1.7%). The mortality rate did not differ between the two cohorts (p > 0.05). The causes of death were a variety of severe...

---

**Table 2.** The distribution of liver diseases in deceased HIV-infected patients out of cohorts with and without antibody to *Toxoplasma gondii* (Nicolle et Manceaux, 1908)

<table>
<thead>
<tr>
<th>Disease</th>
<th>Seropositive patients (N=71)</th>
<th>Seronegative patients (N=48)</th>
<th>Chi-square test</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>T. gondii positive</strong></td>
<td><strong>T. gondii negative</strong></td>
<td><strong>n</strong></td>
<td><strong>P ± m_s %</strong></td>
</tr>
<tr>
<td>Number</td>
<td>Number</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chronic HBV infection</td>
<td>4</td>
<td>5</td>
<td>5.6 ± 2.7</td>
</tr>
<tr>
<td>Chronic HCV infection</td>
<td>6</td>
<td>7</td>
<td>8.5 ± 3.3</td>
</tr>
<tr>
<td>Chronic HBV-HCV coinfection</td>
<td>22*</td>
<td>7</td>
<td>31.0 ± 5.5</td>
</tr>
<tr>
<td>Chronic toxic hepatitis</td>
<td>10</td>
<td>6</td>
<td>14.1 ± 4.1</td>
</tr>
<tr>
<td>Chronic cryptogenic hepatitis</td>
<td>5</td>
<td>5</td>
<td>7.0 ± 3.0</td>
</tr>
<tr>
<td>Chronic EBV-hepatitis</td>
<td>1</td>
<td>1</td>
<td>1.4 ± 1.4</td>
</tr>
<tr>
<td>Chronic CMV-hepatitis</td>
<td>1</td>
<td>1</td>
<td>1.4 ± 1.4</td>
</tr>
<tr>
<td>Liver cirrhosis</td>
<td>17</td>
<td>5</td>
<td>23.9 ± 5.1</td>
</tr>
<tr>
<td>General liver pathology</td>
<td>66*</td>
<td>37</td>
<td>93.0 ± 3.0</td>
</tr>
</tbody>
</table>

* – the difference between the indexes in cohorts of patients with and without *T. gondii* is statistically significant; χ^2 – Pearson’s chi-squared test; C – Pearson’s contingency coefficient; P – percentage; m_s – standard error of percentage; p – statistical significance.

---

Fig. 1. Frequency of different liver pathologies in deceased patients from cohorts seropositive and seronegative to *Toxoplasma gondii* (Nicolle et Manceaux, 1908).

* – difference between the indices for the entire cohort of seropositive patients and for the group of deceased patients from this cohort (p < 0.05); ^ – difference between the indices for the entire cohort of seronegative patients and for the group of deceased patients from this cohort (p < 0.05).
opportunistic diseases including tuberculosis in 36 patients, meningoencephalitis (of different origins) in 23 patients, including toxoplasmosis – in five patients, bacterial lung destruction in two patients and oncology diseases in two patients (Kaposi’s sarcoma in one, non-Hodgkin lymphoma in the other). Decompensated liver cirrhosis as the cause of death was in 22 patients.

According to the data in Table 2, the general liver pathology in deceased patients was more common in seropositive individuals than in seronegative ones (93.0 ± 3.04% and 77.1 ± 6.1%, respectively, p = 0.013). Chronic HBV-HCV coinfection was twice as common in the seropositive cohort than in the seronegative cohort (31.0 ± 5.5% and 14.6 ± 5.1%, respectively, p = 0.041). In deceased patients the frequency of liver cirrhosis showed an insignificant increase in the seropositive cohort (23.9 ± 5.1% and 10.4 ± 4.4%, respectively, p = 0.063).

In deceased patients there were weak associations between seropositivity to T. gondii and chronic HBV-HCV coinfection (C = 0.187), liver cirrhosis (C = 0.168) and the general liver pathology (C = 0.228) (see Table 2). The activity of hepatitis, regardless of etiology, was minimal in both deceased seropositive patients and deceased seronegative patients, except one with cryptogenic hepatitis who had pronounced activity with ALT activity 16-fold higher than the upper limit of the normal level (658 U/ml). Thus, no significant difference of hepatitis activity was observed between cohorts.

Among deceased patients seropositive to T. gondii the number with total liver pathologies increased by 30% compared to the entire seropositive cohort (92.96 ± 3.04% and 63.09 ± 2.09%, respectively, t = 8.1, p < 0.0001). The number of patients with toxic hepatitis was 9.3-fold higher (14.08 ± 4.13 and 1.51 ± 0.53, respectively, t = 3.02, p = 0.0026). For deceased patients with cirrhosis it was 4.5 times higher (23.94 ± 5.06 and 5.27 ± 1.97, respectively, t = 3.44, p = 0.0006), (see Fig. 1).

Among the deceased seronegative patients, general liver pathology increased by 25% compared to the total seronegative cohort (77.08 ± 6.07 and 51.86 ± 2.58, respectively, t = 3.82, p = 0.00015) due to the increase of the proportion of patients with toxic hepatitis (12.50 ± 4.77 and 0.27 ± 0.27, respectively, t = 2.56, p = 0.011). However, among seronegative deceased patients the proportion with liver cirrhosis did not increase (see Fig. 1).

**DISCUSSION**

This study found that T. gondii could have a non-specific additional impact on the development of some liver diseases. The total number of liver diseases occurred more often in seropositive individuals than in seronegative ones. This can be explained by the negative influence of T. gondii on liver parenchyma. In the literature there are reports of the development of disorders of basic metabolic processes in mouse cells under the influence of T. gondii. This parasite is able to block the peroxisome of proliferator-activated receptors of the host cells and through this mechanism to suppress the activity of many genes participating in the metabolism of xenobiotics, fatty acids and lipids, it can weaken the mitochondrial respiratory chain in mice liver cells (He et al. 2016). Due to negative reprogramming of metabolism of liver cells, the toxic effects of medicines, alcohol and other chemical compounds might be increased. *Toxoplasma gondii* can affect liver cells not only independently but also through the mechanism of cytokine imbalance (increased level of pro-inflammatory cytokines, especially INF-γ, TNF-α, IL-12) (Masek and Hunter 2013).

The increased frequency of HBV-HCV coinfection in seropositive to *T. gondii* individuals has remained without proper explanation. However, it is known that the parasite is able to contribute to the replication of HCV and also contribute to its chronisation (Nagwa et al. 2016). On the other hand, HCV infection has been shown to promote the reactivation of latent toxoplasmosis (El-Nahas et al. 2014, Nagwa et al. 2016).

HBV-HCV coinfection in seropositive to *T. gondii* persons more common than in seronegative ones, that probably indicated on a high risk of infection through injection drug using or sexual abuse with traumatisation of mucous membranes or neglecting the contact with menstrual blood, tattoo practice, etc. It could be assumed that such risky behavior of some patients is stimulated by *T. gondii*, which persists in the brain and it is able to do remodelling of the human mental health (Flegr et al. 2002, Lafferty 2006, Flegr 2007).

In patients with a fatal outcome, the general liver pathology as well as the frequency of toxic hepatitis increased in each cohort regardless of their serological status. Here *T. gondii* had a little importance because of low coefficient of contingency with general liver pathology (C = 0.228). Chronic HBV-HCV coinfection was more common in the seropositive cohort than in the seronegative cohort as it was found both in entire cohorts and in deceased patients. *T. gondii* had a weak role in distributing HBV-HCV coinfection among died individuals (C = 0.187). Also, a liver cirrhosis was more common in seropositive deceased patients than in seronegative, and in them *T. gondii* is weakly involved in a formation of cirrhosis (C = 0.168).

These findings are important for healthcare workers because they indicate the necessity for the diagnosis and treatment of active infection with *T. gondii* in HIV-positive individuals with liver diseases.

However the results of the present study may have been influenced by the higher percentage of patients receiving ART in the seronegative cohort than in the seropositive cohort. The use of prophylactic doses of biseptol (sulfamethoxazole/trimethoprim) and azithromycin in patients with low CD4+ T-lymphocyte counts could prevent infection or reactivation of *T. gondii*. Although the same number of people in both cohorts received these medicines, the duration of and their compliance to the treatment could be different.

In conclusion, in HIV-infected patients, *T. gondii* is a weak non-specific additional factor that supports the inflammatory process in the liver and the progression of liver cirrhosis, regardless of their etiology and the degree of activity, but does not influence the degree of hepatitis activity. The increased prevalence of chronic HBV-HCV
co-infection in *T. gondii* carriers may be related to their risk factor behaviour towards uncontrolled blood-borne infections.

**REFERENCES**


Jia-Feng H., Jing-Ping X., Ru-Jin X., Chun-Yan Q. 2018: [Investigation on *Toxoplasma gondii* infection in liver disease patients in Hangzhou City.] Zhonggou Xue Xi Chong Bing Fang Zhi Za Zhi 30: 322–324. (In Chinese.)


Lytvyenyko M.V. 2007: [Etiological structure and peculiarities of liver damage at HIV infection.] Patologiia 4: 43–46. (In Ukrainian.)


Acknowledgements. We would like to thank the doctors, nurses, and other medical staff of Ivano-Frankivsk Regional Center for AIDS Prevention and Fight for their professional assistance throughout the study.

Hryzhak: *Toxoplasma gondii* promotes liver diseases
Toxoplasma gondii promotes liver diseases


Cite this article as: Hryzhak I.H. 2020: Infection with Toxoplasma gondii can promote chronic liver diseases in HIV-infected individuals. Folia Parasitol. 67: 034.