

# Different presence of *Chlamydia pneumoniae*, herpes simplex virus type 1, human herpes virus 6, and *Toxoplasma gondii* in schizophrenia: meta-analysis and analytical study

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**Abstract:** In the present study we have performed both a meta-analysis and an analytical study exploring the presence of *Chlamydia pneumoniae*, herpes simplex virus type 1, human herpes virus 6, and *Toxoplasma gondii* antibodies in a sample of 143 schizophrenic patients and 143 control subjects. The meta-analysis was performed on papers published up to April 2014. The presence of serum immunoglobulin G and immunoglobulin A was performed by enzyme-linked immunosorbent assay test. The detection of microbial DNA in total peripheral blood was performed by nested polymerase chain reaction. The meta-analysis showed that: 1) *C. pneumoniae* DNA in blood and brain are more common in schizophrenic patients; 2) there is association with parasitism by *T. gondii*, despite the existence of publication bias; and 3) herpes viruses were not more common in schizophrenic patients. In our sample only anti-*Toxoplasma* immunoglobulin G was more prevalent and may be a risk factor related to schizophrenia, with potential value for prevention.

**Keywords:** meta-analysis, analytical study, *Chlamydia pneumoniae*, herpes simplex virus type 1, human herpes virus 6, *Toxoplasma gondii*, schizophrenia

## Introduction

The word “schizophrenia” refers to a group of disabling mental disorders characterized by alterations in perception, thought, affectivity, and behavior. Its etiology is unknown and there are no curative treatments.<sup>1</sup> Although present worldwide, schizophrenia is more prevalent in developed countries, where the accumulated lifetime prevalence can reach more than 1% and an annual incidence of five cases per 10,000 people.<sup>2</sup> A possible genetic vulnerability exists for schizophrenia, as supported by the increased familial morbid risk for schizophrenia found in first-degree relatives of affected subjects, possibly in conjunction with the influence of environmental factors. Prenatal and perinatal obstetric complications have also been related to schizophrenia, and some authors postulate that environmental factors (such as trauma, cannabis use, or neurotropic infections) might modulate the final impact of such complications on the early genetic alterations described for this illness.<sup>3</sup>

Among other environmental risk factors for schizophrenia, infectious factors have been posed, a hypothesis that would be supported by some studies describing an increased prevalence for the illness among those subjects born during winter and spring months in the northern hemisphere.<sup>4</sup> Indeed, prevalence has been reported as being much higher following flu epidemics. Several infectious agents could harm a fetus.

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Being primarily neurotropic agents or not, acting directly or in an immune-mediated way, or perpetuating themselves or not, these agents could facilitate the occurrence of irreversible neurologic injuries associated with alterations in both neurotransmission expression and sensorial information processing,<sup>5</sup> which may have a role in the development of schizophrenia later in life. These subjects may potentially benefit from antimicrobial treatment, which may improve their outcome.

Among all the infectious agents that have been studied in schizophrenia, *Chlamydia pneumoniae*, herpes simplex virus type 1 (HSV-1), human herpes virus 6 (HHV-6) and *Toxoplasma gondii* are especially interesting because they are neurotropic, they cause illnesses that could share symptoms with schizophrenia, and they can block neural function in a recurrent manner as they produce latent infections with potential reactivations. Although such possibilities are biologically plausible, the association between microbial agents and schizophrenia is not conclusive.<sup>6</sup> The present study aimed to explore such associations via both a meta-analysis and an analytical study exploring the presence of the abovementioned infectious agents in a sample of patients with schizophrenia and controls.

## Materials and methods

### Meta-analysis

The meta-analysis was performed according to the same procedure used in a paper published by our group 3 years ago.<sup>6</sup> Thus, in brief, we performed a systematic search of all articles published in English or Spanish in journals indexed on MEDLINE, psycINFO, ISI Web of Knowledge, and the Cochrane Library up to April 2014. The search terms used were “schizophrenia” and “herpes” or “*Chlamydia pneumoniae*” or “*Chlamydothila pneumoniae*” or “*Toxoplasma gondii*.” After that search, we excluded a group of papers including noncontrolled cohort studies, reviews, studies that did not present their results in a correct and/or explicit manner, animal studies, studies where the control group comprised patients with other psychiatric or neurological disorders, and those that did not assess the infectious processes. On the other hand, included studies were those that examined patients and healthy controls (characterized by absence of psychiatric disorders or any neurological disease), cohort studies of patients and controls, and case series. It was required that schizophrenic patients were included in the studies, and that the aim of the study was the direct or indirect exploration of the possible association between infection and schizophrenia.

The following data were obtained for each serologic or molecular determination in each publication: odds ratios (OR) and their 95% confidence intervals (CI), population weights, and statistical significance of the analyses. The DerSimonian and Laird method was used as it produces overall estimates that are less affected by heterogeneity among studies.<sup>7</sup> Heterogeneity among studies was determined using Cochran’s Q statistic method when the number of papers included was five or more. Additionally, Higgins I<sup>2</sup> was also used as a measure of total OR variability given heterogeneity among studies. Hence, very high values of the latter measure, above 75%, would indicate a strong heterogeneity, suggesting the need to carry out an additional meta-regression using the restricted maximum likelihood method. In such cases, we also performed a more detailed subanalysis of the different subgroups, when the number of studies to be included in each subgroup was big enough (three or more). As a whole, we considered that no relationship existed between exposure to microorganisms and the presence of schizophrenia when the 95% CI included the one unit value.<sup>8</sup>

Begg’s and Egger’s tests were used when the number of studies included was five or more, in order to check for any particular publication biases and their magnitude.<sup>9,10</sup> Study quality was assessed using the Newcastle-Ottawa Quality Assessment Scale (Available from: [http://www.ohri.ca/programs/clinical\\_epidemiology/oxford.htm](http://www.ohri.ca/programs/clinical_epidemiology/oxford.htm); updated 15 June 2014). Data obtained from the various studies were analysed using the STATA Release 10.1 statistical package (StataCorp LP, College Station, TX, USA).

### Analytical study of infectious agents

#### Sample characteristics: patients with schizophrenia

Patients with a diagnosis of schizophrenia and less than 5 years since their first episode (n=143) were invited to participate in the study. Patients were recruited from the Mental Health Services of Jerez de la Frontera and Jaen (south Spain). All patients fulfilled ICD10 (International Statistical Classification of Diseases and Related Health Problems, 10th Revision) criteria for schizophrenia. They were assessed using a Spanish validated version of the Schedules for Clinical Assessments in Neuropsychiatry. Stability in diagnosis during the last year and no changes in treatment for the last 2 months were also taken into account as inclusion criteria. All patients signed an informed consent prior to being included in the study.

#### Sample characteristics: control subjects

The control group comprised 143 unrelated individuals with no personal or family history of mental disorders or suicide

and with no pharmacological or psychological treatment that could interfere with the neuropsychological assessment. A specifically trained psychologist interviewed each participant and used the Schedules for Clinical Assessments in Neuropsychiatry to check the absence of these conditions. They all came from different primary care centers belonging to the Primary Care catchment area of south Granada (Spain). All participants were nonimmunosuppressed and signed an informed consent prior to being included in the study.

Controls were stratified by age and sex with cases. In both groups, sex was equally distributed (62.24% of cases and 62.94% of controls were male subjects; chi-square =0.001,  $P=0.973$ ) and mean age was also comparable between cases and controls (28.69 [standard deviation =5.41] versus 30.42 [standard deviation =5.62] respectively;  $t=1.771$ ,  $P=0.079$ ).

After losing some laboratory samples for different reasons, our final count was 271 blood samples (128 cases and 143 controls) and 284 plasma samples (142 cases and 142 controls). A microbiological study with these samples was made using the following procedures.

### Specific serum immunoglobulins

Different commercial kits were used to detect different microbial immunoglobulin Gs (IgGs): the HERPES SIMPLEX 1 ELISA (enzyme-linked immunosorbent assay) IgG kit system developed by Vircell (Granada, Spain); the TOXOPLASMA ELISA IgG kit, also distributed by Vircell; the human HERPESVIRUS 6 ELISA IgG kit designed by Alere Healthcare (Barcelona, Spain); and the CHLAMYDOPHILA PNEUMONIAE ELISA IgG kit, developed by Savyon Diagnostics (Madrid, Spain). *C. pneumoniae* immunoglobulin A (IgA) was analyzed using a specific kit designed by Vircell (Granada, Spain). As for the rest of the IgA measurements, the assay was previously standardized by using the same solid phase previously used for IgG, a human anti-IgA conjugate (Siemens Lab, Barcelona, Spain), and a commercialized, specific IgA positive control with known concentration

(Lab. Aviva Systems Biology, San Diego, CA, USA). Serum was diluted using anti-IgG. To determine the working dilution for both the conjugate and the positive control, dilutions of both were prepared. These were then doubly processed, comparing the average absorption result against the negative control. The sample was considered positive in detecting IgA when well absorbance was equal to or greater than double the threshold positive value. Samples were processed in duplicate according to our routine laboratory method. The mean absorbance value was used in the evaluation. The coefficients of variation of the ELISA were less than 10%.

### Detection of peripheral-blood microbial DNA

DNA was extracted using ReliaPrep™ Blood gDNA Mini-prep System (Promega Biotech Iberica, Madrid, Spain). Specific DNA was detected with nested polymerase chain reaction (PCR) of the *C. pneumoniae* *pst1* region,<sup>11</sup> HSV-1 *gP-D-1* region,<sup>12</sup> VHH-6 *U67* region,<sup>13</sup> and *T. gondii* *B1* region.<sup>14</sup> We reached sensitivities up to 9 copies/μL (for American Type Culture Collection [ATCC] VR-1356 *C. pneumoniae*), 16 copies/μL (for ATCC VR-735 VHS-1), 6 copies/μL (for ATCC VR-1480 VHH-6), and 10 copies/μL (for ATCC 50174 *T. gondii*). We standardized PCR conditions to improve sensitivity (Table 1). We sequenced the amplified fragment in positive cases to demonstrate specificity. *ACTB* gene DNA was used as a control procedure (Gene-Link, Hawthorne, NY, USA).

### Statistical analyses

Pearson's chi-squared test was used to compare qualitative variables (serum IgG and IgA and blood DNA). A  $P$ -value of 0.05 or less was considered statistically significant.

## Results

### *Chlamydia pneumoniae*

Finally, three studies were included in the meta-analysis<sup>15-17</sup> (see Table 2). All of these studies showed similar quality and

**Table 1** Nested-PCR conditions for characterization of *Chlamydia pneumoniae*, HSV-1, HHV-6, and *Toxoplasma gondii*

Step	<i>Chlamydia pneumoniae</i>		HSV-1		HHV-6	<i>Toxoplasma gondii</i>	
	PCR (external)	PCR (internal)	PCR (external)	PCR (internal)	PCR (external and internal)	PCR (external)	PCR (internal)
Initial denaturalization	94°C 4 min	94°C 4 min	94°C 1 min	94°C 1 min	94°C 2 min	94°C 1 min	94°C 1 min
Denaturalization	94°C 1 min	94°C 1 min	94°C 10 sec	94°C 10 sec	94°C 15 sec	94°C 10 sec	94°C 10 sec
Hybridization	52°C 1 min	54°C 1 min	58°C 20 sec	67°C 20 sec	54°C 20 sec	57°C 10 sec	63°C 10 sec
Extension	72°C 1 min	72°C 1 min	72°C 20 sec	72°C 20 sec	72°C 20 sec	72°C 30 sec	72°C 15 sec
Final extension	72°C 10 min	72°C 10 min	72°C 2 min	72°C 2 min	72°C 2 min	72°C 2 min	72°C 2 min
Number of cycles	40	40	35	30	40	40	40

**Abbreviations:** HHV, human herpes virus; HSV, herpes simplex virus; PCR, polymerase chain reaction; min, minutes; sec, seconds.

Table 2 Analysis and evaluation of studies included in the meta-analysis

Microorganism	Reference	Sample	Technique	Determination	Descriptive statistics				Inferential statistics			Quality			Global OR (95% CI; P-value)	
					Cases		Controls		OR	95% CI	Weight (%)	S	C	E		
					Pos	Neg	Pos	Neg								
<i>Chlamydia pneumoniae</i>	15	BL	N-PCR	DNA	4	6	6	109	12.11	2.68–54.75	14.91	**	**	**	5.96 (3.42–10.39; P<0.001)	
	16	BL	PCR	DNA	14	58	9	216	5.79	2.43–13.79	41.01	**	**	**		
	16	BR	PCR	DNA	4	30	1	34	4.53	0.67–30.64	8.45	**	**	**		
	17	BL	PCR	DNA	11	61	8	217	4.89	1.88–12.70	35.64	**	**	**		
	18	BR	PCR	DNA	0	8	0	16	1.94	0.03–106.66	0.50	***	**	*		1.17 (0.88–1.54; P=0.273)
	19	SE	ELISA	IgG	26	21	297	227	0.95	0.52–1.71	15.30	***	**	*		
	20	SE	ELISA	IgG	41	19	70	40	1.23	0.63–2.41	13.00	***	**	*		
	21	BR	HIB	DNA	0	20	0	21	1.05	0.02–55.36	0.50	***	**	*		
HSV-1	22	BR	N-PCR	DNA	0	14	0	26	1.83	0.03–97.01	0.50	***	**	*	1.17 (0.88–1.54; P=0.273)	
	23	SE	IFA	IgG	18	20	27	14	0.47	0.19–1.16	8.00	***	**	*		
	24	SE	ELISA	IgG	7	4	3	6	3.50	0.55–22.30	2.40	***	**	*		
	25	SE	IFA	Ab	41	0	25	0	1.63	0.03–84.59	0.50	**	**	*		
	26	SE	ELISA	IgG	15	15	8	36	4.50	1.58–12.84	6.30	***	**	*		
	27	BR	IPA	Ag	0	25	0	16	0.65	0.01–34.23	0.50	***	**	*		
	28	BR	N-PCR	DNA	0	30	0	23	0.77	0.01–40.28	0.50	***	**	*		
	29	BR	HIB	DNA	0	25	0	31	1.24	0.02–64.45	0.50	***	**	*		
	30	SE	ELISA	IgG	105	87	54	38	0.85	0.51–1.40	19.30	***	**	*		
	31	SE	ELISA	IgG	503	177	192	91	1.35	1.00–1.82	32.30	***	**	*		0.34 (0.49–2.42; P=0.283)
22	BR	N-PCR	DNA	0	14	1	25	0.59	0.02–15.34	38.16	***	**	*			
HHV-6	24	SE	ELISA	IgG	8	3	9	0	0.13	0.01–2.85	42.21	***	**	**	2.50 (1.40–4.47; P=0.002)	
	28	BR	N-PCR	DNA	0	30	0	23	0.77	0.01–40.28	19.63	***	**	**		
	19	SE	ELISA	IgG	16	31	123	401	1.68	0.90–3.16	14.10	***	**	*		
<i>Toxoplasma gondii</i>	22	BR	N-PCR	DNA	0	14	0	26	1.83	0.03–97.01	1.90	***	**	*	2.50 (1.40–4.47; P=0.002)	
	32	SE	ELISA	Ab	5	13	16	164	3.94	1.25–12.48	10.40	***	**	*		
	32	SE	ELISA	Ab	5	15	16	164	3.42	1.10–10.63	10.60	***	**	*		
	33	SE	ELISA	Ab	66	34	11	39	6.88	3.13–15.11	13.00	**	*	**		
	34	SE	ELISA	Ab	60	197	171	511	0.91	0.65–1.27	16.00	***	**	**		
	35	SE	ELISA	Ab	15	165	37	495	1.22	0.65–2.27	14.20	***	**	**		
	36	SE	ELISA	Ab	16	24	5	32	4.27	1.37–13.28	10.50	**	**	*		
	37	SE	ELISA	Ab	14	24	3	24	4.67	1.19–18.35	9.20	***	**	**		

Notes: Asterisks indicate the level of study quality: from low quality (\*) to (\*\*\*\*) high quality.

Abbreviations: Ab, antibody; Ag, antigen; BL, blood; BR, biopsy of brain tissue; C, Comparability; CI, confidence interval; E, Exposure; ELISA, enzyme-linked immunosorbent assay; HHV, human herpes virus; HIB, hybridization; HSV, herpes simplex virus; IFA, immunofluorescence assay; IgG, immunoglobulin G; IPA, immunoperoxidase assay; N-PCR, nested-polymerase chain reaction; Neg, negative; OR, odds ratio; PCR, polymerase chain reaction; Pos, positive; S, Selection; SE, serum.

used blood and brain samples to detect bacterial DNA through PCR or nested-PCR. The meta-analysis revealed an association between schizophrenia and *C. pneumoniae* (OR =5.96; 95% CI =3.42–10.39;  $P<0.001$ ). However, our own results found no significant differences between cases and controls for IgG ( $P=0.521$ ), IgA ( $P=0.885$ ), and specific DNA measures because DNA was scarcely represented (see Table 3).

## Herpes simplex virus type 1 and human herpes virus 6

Fourteen studies were finally included in the meta-analysis for these two viruses (Table 2).<sup>18–31</sup> The combined OR estimation involving the 14 studies comparing detection of different markers for infection by HSV-1 in schizophrenic patients and healthy controls was 1.17 (95% CI =0.88–1.54;  $P=0.273$ ). When heterogeneity tests were done, we obtained a value of  $\chi^2_{\text{exp}}=15.4$  ( $df=13$ ;  $P=0.312$ ), after which we could state that the differences found between the studies were due to randomness. This fact was corroborated by an  $I^2$  coefficient of 15.4%, indicating that only 15.4% of the variability of the ORs was due to the heterogeneity among studies.

As shown, in spite of the absence of important and, if any, nonsignificant heterogeneity (<75%), a detailed study was carried out to consider the potential effect of some factors on the value of the overall OR through meta-regression, and it was concluded that the technique used to detect HSV-1 infection did not seem to affect OR value ( $P=0.274$ ). Thus, we found no difference between risks arising from studies that detected DNA in brain tissue (OR =1.15; 95% CI =0.23–5.80) and that from studies in which serum antibodies were detected (OR =1.19; 95% CI =0.81–1.74).

Begg's test was not significant ( $P=0.743$ ), suggesting that there was a potential publishing bias. A similar result was obtained after Egger's test ( $P=0.817$ ). The quality of the studies examined was medium-high, and it was higher in the more-recent studies due to the introduction of quality validation scales, a fact that favored the control of the data included in the articles.

Finally, only three studies comparing infection by HHV-6 were included, all of them of acceptable quality (see Table 2).<sup>22,24,28</sup> The statistical analysis could not confirm the existence of a significant association between infection by this virus and schizophrenia (OR =0.34; 95% CI =0.49–2.42;  $P=0.283$ ). Our results about IgG, IgA, and specific DNA measures are detailed in Table 3. DNA was scarcely present. No significant differences were found between cases and controls for these markers ( $P=1.000$  for HSV-1 IgG;  $P=0.133$  for HHV-6 IgG;  $P=1.000$  for HHV-6 IgA), with the only exception being HSV-1 IgA, which was more frequently detected in controls than in cases ( $P=0.015$ ).

## Toxoplasma gondii

We included eight studies centered on the potential relationship between *T. gondii* infection markers and schizophrenia as compared with healthy controls (Table 2).<sup>19,22,32–37</sup> All these studies had similar weights, except for the one by Conejero-Goldberg et al<sup>22</sup> that used postmortem brain tissue samples, and thus included a smaller number of cases and controls; this study was also of lower quality according to the Newcastle-Ottawa Scale. Due to their large sample size, the studies by Niebuhr et al<sup>35</sup> and Mortensen et al<sup>34</sup> stood out, the former being of the highest quality. The latter obtained the narrowest 95% CI range due to the large amount of participants included. After combining the different studies, we found a significant association between parasitization by *T. gondii* and schizophrenia ( $P=0.002$ ); the combined OR was 2.50 (95% CI =1.40–4.47). We can therefore state that schizophrenia was 2.5 times more frequent in people in whom the marker for *T. gondii* was found when compared to people in whom it was not detected. The study by Conejero-Goldberg et al<sup>22</sup> was the least-precise study and the one with the widest confidence interval. When heterogeneity tests were performed between the studies, a significant value was obtained:  $\chi^2_{\text{exp}}=35.26$  with  $df=8$ ,  $P<0.0001$ . This result parallels an  $I^2$  coefficient equal to 77.3%, indicating that 77.3% of the variability of the ORs is due to the heterogeneity of the studies analyzed.

**Table 3** Positive samples for IgG, IgA, and DNA in patients with schizophrenia and in controls

		IgG			IgA			DNA*	
		Cases (n=142)	Controls (n=142)	P-value	Cases (n=142)	Controls (n=142)	P-value	Cases (n=128)	Controls (n=143)
<i>Chlamydia pneumoniae</i>	Samples (%)	41 (28.9)	47 (33.1)	0.521	29 (20.4)	31 (21.8)	0.885	2 (1.6)	0 (0)
HSV-1	Samples (%)	125 (88.0)	124 (87.3)	1.000	12 (8.5)	27 (19.0)	0.015	1 (0.8)	2 (1.4)
HHV-6	Samples (%)	112 (78.9)	100 (70.4)	0.133	3 (2.1)	4 (2.8)	1.000	1 (0.8)	2 (1.4)
<i>Toxoplasma gondii</i>	Samples (%)	56 (39.4)	29 (20.4)	0.001	1 (0.7)	2 (1.4)	0.624	1 (0.8)	0 (0)

**Note:** \*DNA was scarcely present in samples, so the  $P$ -values were not calculated.

**Abbreviations:** IgA, immunoglobulin A; IgG, immunoglobulin G; HHV, human herpes virus; HSV, herpes simplex virus.

Such heterogeneity is explained by significant differences between reported risk at the only publication that detected *T. gondii* DNA in brain biopsies (OR =1.83; 95% CI=0.03–97.01;  $P=0.001$ ) and risk reported by studies using detected serum antibodies (OR =2.74; 95% CI =1.33–5.62). Finally, even though Begg's test was not significant ( $P=0.711$ ), Egger's test was ( $P=0.03$ ), indicating a tendency to publish studies where results were significant, according to this last test.

Our own sample results on IgG, IgA, and specific DNA measures are detailed in Table 3. DNA was scarcely present. No significant differences were found between cases and controls for IgA measures ( $P=0.624$ ). However, IgG was significantly more frequently detected in cases than in controls ( $P=0.001$ ).

## Discussion

### *Chlamydia pneumoniae*

Previous studies describing a significant association between *C. pneumoniae* infection and the origin of schizophrenia have been published<sup>15–17</sup> (Table 2). Thus, Fellerhoff et al<sup>16,17</sup> found DNA of *C. pneumoniae* in 15.3% of mononuclear blood cells from patients with schizophrenia and 11.8% in brain biopsies of these subjects. Finally, these authors described a significant improvement in psychotic symptoms in schizophrenic patients treated with azitromicine. Moreover, a study showed an improvement in cognitive functioning among schizophrenic patients treated with clozapine plus minocycline.<sup>38</sup> The meta-analysis carried out in the present study reveals that *C. pneumoniae* is present in some patients and may be a potential etiological agent in schizophrenia.

This bacteria enters the organism via the respiratory system and it spreads using monocytes and lymphocytes reaching the central nervous system.<sup>39</sup> Th1 lymphocytes are activated by exposure to bacterial antigens (in glial cells, for example), causing secretion of proinflammatory cytokines (eg, interferon- $\gamma$ ), which, in turn, activate macrophages. Such macrophages increase indoleamine 2,3-dioxygenase (IDO) production, which converts tryptophan into kynurenic acid, hence diminishing tryptophan availability. In such a context, the pathogen cannot replicate itself. Intracellular kynurenic acid inhibits glutamine and nicotinic receptors, which are responsible for cognitive impairment. An increased activity of IDO, which may be caused by the infection, has been described in genetically vulnerable patients with schizophrenia.<sup>40</sup> The failure of this defensive mechanism produces persistent and chronic infections with intermittent replication,<sup>41</sup> along with both maintained and

insufficient levels of kynurenic acid. Prenatal administration of lipopolysaccharide seemed to decrease dopamine concentration in rats' frontal cortex whilst enhancing dopaminergic system activity in the striatum.<sup>42</sup>

Our own sample results do not show an association between *C. pneumoniae* infection and schizophrenia. The increased prevalence of *C. pneumoniae* infection in subjects without a respiratory illness may explain the lack of differences in antibody levels when comparing cases and controls.<sup>43</sup> Moreover, because the infection is focused on the central nervous system, it seems reasonable to detect microbial DNA in brain (that could be responsible for the illness) but not in peripheral blood.<sup>16</sup> Nonetheless, limited sample power cannot be ruled out for this negative finding.

### Herpes simplex virus type I and human herpes virus 6

Different studies have explored the association between HSV-1/HHV-6 infections and schizophrenia, showing inconclusive results.<sup>6,18–29,34,35,44–52</sup> Some of these studies have been included in our meta-analysis (Table 2). The lack of association found in the meta-analysis may reflect a real absence of association or a type 2 error due to the use of indirect measurements on brain samples. Only two previous studies have been able to find a significant association between HSV-1 and schizophrenia.<sup>26,31</sup> In such studies, authors postulate that maternal HSV-1 antibodies may cause fetal brain damage even in the absence of a fetal infection.

However, HSV-1 is neurotropic and mainly spreads in frontal and temporal brain regions in which certain impairments could produce cognitive and memory alterations similar to those found in patients with schizophrenia.<sup>26,53</sup> Moreover, a controlled trial showed an improvement in cognitive functioning among seropositive schizophrenic patients treated with one antipsychotic plus valacyclovir when compared to those treated with the antipsychotic agent only.<sup>54</sup> Nevertheless, these results may be due to a simple clinical coincidence<sup>30</sup> due to the high prevalence of this infection, as shown in other studies (Table 2). Also, regarding the possible role of the glutamate system, the association between mutations in the specific GluN2B N-methyl-D-aspartate (NMDA) receptor subunit and maternal herpes simplex virus infection and schizophrenia,<sup>55</sup> a disorder where brain activation and cortical neurotoxicity of NMDA receptor antagonists, has been extensively reported.<sup>56,57</sup> In addition, schizophrenia has also been related to an interaction of prenatal immune activation and subsequent peripubertal stress.<sup>58,59</sup> Furthermore, schizophrenia has also been linked with the effect of stress

on hippocampal neurogenesis,<sup>60</sup> apoptosis, and parvalbumin expression.<sup>61,62</sup>

Our own sample results also failed to find any association, and the clinical meaning of the increased frequency of IgA anti-HSV-1 among controls is not easily explainable. However, the cause of medical consultation may be associated with a herpetic reactivation. This fact is difficult to demonstrate in a retrospective design and it is something that we did not take into account during our control recruitment. Because such herpes virus associations do not have to involve a primary effect, and they cannot be detected from cross-sectional studies<sup>63</sup> due to the high prevalence of the infection, future longitudinal studies are needed to conclude if there are causal associations between virus infections and psychiatric illness.

### *Toxoplasma gondii*

In the last years, several kinds of studies (serological, pharmacological, epidemiological, and behavioral studies) have been published on the association between *T. gondii* infection and changes in human behavior and neuropsychiatric disorders, including schizophrenia.<sup>19,22,32–35,37,44,46,51,64–74</sup> There is clinical evidence supporting the interest of studying the role of *T. gondii* on such phenotypes: 1) higher titers of IgG anti-toxoplasma have been found in the serum and cerebrospinal fluid of patients with early-onset and late-onset diagnostics; 2) some antipsychotics (haloperidol, risperidone, fluphenazine) show anti-*T. gondii* activity; and 3) seropositive patients showed a predominance of positive symptoms.

Different meta-analyses on experimental studies focused on infection for *T. gondii* have been published. Results are congruent enough with the existence of an association between infection for *T. gondii* and schizophrenia. Our current meta-analysis OR values are very similar to those reported earlier, namely: OR =2.73 (95% CI =2.10–3.60;  $P<0.001$ ) in the study by Torrey et al<sup>75</sup> and OR =2.70 (95% CI =1.34–4.42;  $P=0.005$ ) in our group's previous meta-analysis.<sup>6</sup> According to these results, we could say that schizophrenia was 2.7 times more frequent in people infected by *T. gondii* than in people with no signals of this infection. Thus, *T. gondii* may indeed play a key role in the etiology of schizophrenia.

In our present meta-analysis, which includes the most-recent publications on the field, schizophrenia was found to be 2.5 times more frequent in people with a previous *T. gondii* infection than in subjects with no signals of having been under the effects of a *T. gondii* infection. However, we have to point out that heterogeneity tests and publication bias

were also significant. Studies supporting this positive finding were: 1) larger studies, where it is more difficult to detect differences, and 2) studies involving soldiers who showed an increased seroprevalence, probably due to their lifestyles. But the significant OR may also be due to a higher prevalence of antibodies acquired after the onset of the disease, due to, for instance, bad hygienic habits.<sup>69</sup>

In hypothesis, a previous neurological injury caused by the parasite, and the consequent inflammatory response,<sup>76</sup> may activate astrocyte function, increasing levels of intracerebral kynurenic acid the toxicity of which acts to inhibit glutamine and nicotinic receptors. These changes are believed to be related to serotonin and melatonin deficits, which would contribute to increased cognitive impairment.<sup>67,77</sup> Such mechanism has also been suggested for *C. pneumoniae*. Increased levels of dopamine in the central nervous system is another consequence of *T. gondii* infection.<sup>78</sup> This effect would be modulated through genes involved in tyrosin-hydroxylase synthesis, which metabolizes L-tyrosine into dihydroxyphenylalanine (a dopamine precursor) and could explain some behavioral and motor symptoms.<sup>79</sup> Also, *T. gondii* has been shown to induce elevated levels of central nervous system dopamine in experimentally infected animals.<sup>80</sup>

Since most of the previous studies have not detected an increase of anti-toxoplasma immunoglobulin M, it is thought that manifestations are a consequence of a parasite reactivation. However, the study of parasitism is especially important because the first infection occurs in the fetus and produces a congenital toxoplasmosis with neurodevelopmental consequences.<sup>81</sup> These alterations might be evident only later in life, maybe coinciding with the start of gray matter degeneration after puberty.<sup>82</sup> Some disorders, such as chorioretinitis, also appear in subjects around 20–30 years old and could have a similar pathogenesis.<sup>77</sup> This fact is supported by serological studies<sup>34,65</sup> showing increased levels of IgG in mothers and newborns who are at higher risk for schizophrenia in adult life. Finally, Xiao et al<sup>74</sup> described an association between *T. gondii* genotype I fetal infection and schizophrenia. This genotype is responsible for up to 75% of congenital toxoplasmosis in Spain.<sup>83</sup> Thus, there seems to be a plausible connection between primary maternal parasitism and increased risk for schizophrenia in the offspring.<sup>69</sup>

We also found that anti-*T. gondii* IgG was more frequent in patients with schizophrenia than in controls, supporting the possibility of a previous infection with a possible intracerebral inflammatory response. DNA studies using blood samples were not significant. This would be in agreement with poor systemic consequences in the process.

## Conclusion

The present meta-analysis shows that *C. pneumoniae* DNA (measured both in blood and brain) is significantly more frequent in patients with schizophrenia than in controls. A significant association was also detected between *T. gondii* parasitism and schizophrenia, even after controlling for publication bias. In addition, in the analysis performed in our own sample, we found an important relationship between the illness and the presence of IgG anti-*T. gondii*, which may suggest the potential interest of detecting such parasitism to develop potential prevention strategies. Finally, we think that new studies are needed to have robust, and more conclusive results. We need prospective and comparative studies, with larger patient and control samples and using combined microbiological techniques for the analysis of the same subject and sample. Future studies should also take into account the illness phase and analyze blood, cerebrospinal fluid, and brain tissues simultaneously using standardized, sensitive, and appropriate techniques, including, if possible, pregnant women and their offspring as subjects under analysis.

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## Disclosure

The authors report no conflicts of interest in this work.

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