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1 **The association between *Toxoplasma gondii* infection and**
2 **postpartum blues**

3
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26 **Abstract**

27 **Introduction and Aim:** *Toxoplasma gondii* is an intracellular protozoan parasite
28 infecting approximately 30% of the global human population. It has often been
29 suggested that chronic infection with *T. gondii* is related to personality changes and
30 various mental disorders including depression. It is not known whether this includes
31 post-partum blues or depression. In this study, we test the hypothesis that there is a
32 relationship between *T. gondii* infection and post-partum blues by measuring the
33 association between infection and postpartum blues.

34 **Methods:** A total of 475 Chinese women who have just given birth were detected
35 serology for *Toxoplasma* IgG and IgM antibodies, and evaluated the degree of
36 depression by Hamilton Depression Scale (HAMD) score. Data were analyzed by
37 Chi-square or Fisher's Exact tests using SPSS software.

38 **Results:** We found an overall *Toxoplasma* seroprevalence of 5.68% (27/475; 95% CI:
39 3.59 - 7.77) which was broken down into a prevalence of 6.60% (7/106; 95% CI: 1.80
40 – 11.41) in mothers with post-partum blues and 5.42% (20/369; 95% CI: 3.10 – 7.74)
41 in non-affected mothers. There was no significant association between infection and
42 post-partum blues ($p = 0.64$).

43 **Conclusion:** The results suggest that there is no relationship between *T. gondii*
44 infection and postpartum blues, at least in this sample of patients from China.

45
46 **Key words:** *Toxoplasma gondii*, postpartum blues, seroprevalence, newborn, puerpera,
47 Hamilton Depression Scale

48 1. Introduction

49 *Toxoplasma gondii*, which infects 30% to 50% of the global population, is one of
50 the most common parasites affecting both healthy and immunocompromised humans
51 (Furtado et al., 2011; Klaren and Kijlstra, 2002; Prandota, 2014). It is believed that *T.*
52 *gondii* infection, termed toxoplasmosis, in immunocompetent individuals is generally
53 considered asymptomatic (Halonen and Weiss, 2013). However, considering an
54 extended set of disease conditions, evidence is accumulating that strongly suggests
55 that this parasite may be implicated in neurodegenerative diseases and are gradually
56 emerging as a global health threat (Nissapatorn, 2010; Furtado et al., 2011; Torgerson
57 and Mastroiacovo, 2013). For example, chronic infection with *T. gondii* is more
58 frequent in individuals with schizophrenia than in psychiatrically healthy controls, as
59 indicated in several studies from different countries (Torrey and Yolken, 2001, 2003;
60 Torrey et al., 2007). In addition, a number of studies have also demonstrated that *T.*
61 *gondii* seropositivity is related to personality changes and various mental disorders
62 including development of suicidal tendencies, obsessive compulsive disorder, bipolar
63 disorder, and depression (Sutterland et al., 2015; Arling et al., 2009; Hinze-Selch et
64 al., 2010; Ling et al., 2011; Okusaga et al., 2011; Tedla et al., 2011).

65 Depressive disorder, also known as clinical depression, is a mood disturbance
66 characterized by changes in mood, and loss of interest, pleasure, cognitive function, sleep,
67 appetite, or energy level (Pratt and Brody, 2008). Furthermore, clinical depression is
68 commonly associated with significant morbidity and mortality (Hsu et al., 2014). It has
69 been reported that, during 2009-2012, about 7.6% Americans aged 12 and over had

70 depression (Pratt and Brody, 2014). However, the prevalence of mood disorders in China
71 appears to be lower than that found in North America and Western Europe. Data indicated
72 that the prevalence of major depressive disorders in China was at 1.96% in Kunming,
73 5.3% in Beijing, 3.6% in Shenyang and 1.15% in Jiangsu, respectively (Hu, 2003; Lu et
74 al., 2008; Ma et al., 2009; Qin et al., 2008). Moreover, depression is also considered a
75 common mental health problem among women of childbearing age, with a higher
76 prevalence rate ranging from 10 to 32% (Ertel et al., 2011; Wang et al., 2011). Postpartum
77 nonpsychotic depression is the most common complication of childbirth, affecting
78 approximately 10-15% of women, and represents a considerable health problem affecting
79 women and their families. A mother's ongoing depression can contribute to later
80 emotional, behavioral, cognitive and interpersonal problems in the offspring. Because of
81 these serious consequences, exploring potential factors, early diagnosis and intervention
82 treatment of postpartum illnesses are imperative for the health and well-being of mother
83 and child (Robertson et al., 2004). Previous studies have consistently demonstrated
84 common significant predictors of postpartum depression as follows: experiencing
85 depressed moods or anxiety during pregnancy, life events, no social support and
86 socioeconomic status (Patel et al., 1999; Bartley, 1994; Beck, 2001; Brugha et al., 1998;
87 Neter et al., 1995). Furthermore, studies have shown that latent *T. gondii* infection is
88 associated with symptoms of depression during pregnancy (Groer et al., 2011).

89 Post partum blues are self-limiting depression commonly found within one week
90 soon after delivery, and are considered as important indicators of depression
91 (Maliszewska et al., 2016; Reck et al., 2009). However, it remains unknown whether

92 parasite infection may play an important role in postpartum blues. We are interested
93 in the relationship between chronic latent *T. gondii* infection and the development of
94 postpartum blues in new mothers. Using a cohort of new mothers in China, we aim to
95 investigate this globally important question and to test the hypothesis that *T. gondii*
96 infection may be more frequent and/or more intense in patients with major
97 postpartum depression compared with psychiatrically healthy controls.

98

99 **2. Materials and Methods**

100

101 **Participants and Questionnaires**

102 In this study, blood samples were collected from 475 women one week after
103 delivery (puerpera) in the First Affiliated Hospital of Guangzhou Medical University,
104 China. Mothers were randomly selected for this study and all those selected were
105 tested by a psychiatrist irrespective of any obvious signs of depression. Due to the low
106 prevalence of *Toxoplasma* infection in China, the sample size needed to ensure
107 sufficient statistical power (minimum n=442) was calculated using a previously
108 published prevalence of 7.8% (ONIHPDCSS, 2005). Furthermore, this value
109 concurred with our previous studies (collated from greater than 120000 pregnant
110 women (1990 - 2010)) that recorded the seroprevalence of *Toxoplasma* to be less than
111 10% in this demographic group (Gao et al., 2012). Sera were separated by
112 centrifugation and stored at -80 °C until serological testing. All participant
113 information was obtained (usually 2-3 days postpartum) through questionnaires and

114 recorded. These included participants' age, occupation, relevant eating habits
115 (consumption of raw or undercooked meat – past and current) and cat contact (current
116 and past cat ownership, cats in the same household, playing closely with cats,
117 cleaning cat litter).

118

119 **Ethical Approval**

120 This study was approved by the Medical Science Ethical Committee of Sun-Yet
121 San University and the First Affiliated Hospital of Guangzhou Medical University. All
122 enrolled participants were informed about the objectives of the study, and written
123 informed consent was obtained from all of them.

124

125 **Hamilton Depression Scale (HAMD) score**

126 Psychiatrists evaluated the degree of depression according to the HAMD
127 (Hamilton, 1967) depression scale score, clinical symptoms and the exclusive criteria.
128 A HAMD score of <8, was considered normal; 8-19, indicated mild depression; 20-
129 34, indicated medium depression; and ≥ 35 , was considered as severe depression. All
130 patients with mild, medium and severe depression were considered as postpartum
131 blues cases according to their clinical signs.

132

133 **Serological Tests**

134 All serum samples were tested for anti-*T. gondii* antibodies - both IgM (indicator of
135 acute infection) and IgG (indicator of chronic infection) using a commercially available

136 enzyme-linked immunosorbent assay (ELISA) kit (Modern Gaoda Biotechnology
137 Company, Beijing) according to the manufacturer's instructions (accuracy > 96.7%,
138 detection limit of 5 IU/ml and coefficient of variation < 15%). Positive, negative and
139 cutoff serum controls were included in every plate. The optical densities (ODs) were
140 measured at 450 nm in a microplate reader (Thermo Scientific Multiskan FC, Thermo
141 Scientific, China). All serum samples were run in triplicate. Samples with ODs above the
142 cutoff value were considered as serologically positive.

143

144 **Statistical Analysis**

145 Data on the prevalence of anti-*T. gondii* antibodies and depression symptoms in
146 population groups were analyzed by Chi-square or Fisher's Exact tests using SPSS
147 software. Risk factors of occupation, seropositivity for *T. gondii*, raw meat consumption
148 and a significant contact with cats were evaluated by using odds ratios (ORs), together
149 with their corresponding 95% confidence intervals (95% CIs). Bias corrected ORs were
150 obtained by adding 0.5 to each data point.

151

152 **3. Results**

153

154 The collection of 475 serum samples taken from the new mothers was examined
155 for anti-*T. gondii* IgG and IgM antibodies using enzyme-linked immunosorbent assay
156 (ELISA). All 475 puerpera samples were IgM negative, suggesting that no current
157 infection of the parasite was occurring. Anti-*Toxoplasma* positive IgG was detected in

158 27 (5.68%; 95% CI: 3.59 - 7.77) of the 475 serum samples as shown in Table 1.

159 Hamilton Depression Scale scores in the same set of new mothers were

160 determined. All patients with mild, medium and severe depression were considered as

161 true cases having postpartum blues according to their clinical signs. There was no

162 significant association between *T. gondii* seropositivity and postpartum blues ($p =$

163 0.643; $n = 475$). When broken down, the seroprevalence of *T. gondii* was slightly

164 higher in the new mothers with postpartum blues (6.60%, 7/106; 95% CI: 1.80 –

165 11.41) than that in the remaining normal group (5.42%, 20/369; 95% CI: 3.10 – 7.74)

166 but this was not significant ($p = 0.643$, OR = 1.234[0.507-3.002]) – see Table 1.

167

168 To further explore the hypothesis that *T. gondii* infection is related to postpartum

169 depression, it is possible that there is a quantitative influence on the relationship. For

170 example, *T. gondii*-positive new mothers might show higher scores on the depression

171 scale when compared to their uninfected counterparts. To examine that, we further

172 analyzed the relationship between *T. gondii* immunoglobulin G (IgG) optical densities

173 (ODs) and the Hamilton Depression Scale (HAMD) scores. We found that depression

174 scores were not significantly different ($p = 0.873$) between the positive and negative

175 *T. gondii* optical density (ODs) groups in the entire population of new mothers

176 ($n=475$) (Figure S1A). In addition, as shown in Figure S1B, the *T. gondii* positive

177 group ($n = 27$) did not show a positive correlation of *T. gondii* ODs (serointensity)

178 with the Hamilton Depression Scale (HAMD) scores ($p = 0.214$). Although we

179 recognize that the sample sizes in this breakdown are small, these analyses further

180 support the prevalence data that suggests there is no significant association between
181 infection and postpartum blues.

182

183 According to previous reports, the probability of becoming infected with *T.*
184 *gondii* increases with age because the infection is ubiquitous and the probability of
185 being exposed to infection increases with age (Hinze-Selch et al., 2007). Therefore,
186 we analysed the seroprevalence of *T. gondii* in the 475 pregnant women in different
187 age groups (Table 2) and the results demonstrated that there was no significant
188 correlation between *T. gondii* seroprevalence and age ($p = 0.872$). Therefore, the
189 results from our current study demonstrate that anti-*T. gondii* seropositivity is not age
190 dependent in this sample of mothers. One possible reason for the results is that the
191 age range of pregnant women is too narrow to reveal this effect and we also recognize
192 that the sample sizes in this breakdown are small. We also found that the incidence of
193 postpartum blues did not show a significant relationship with age of the mothers ($p =$
194 0.610) (Table 2) although again our numbers of individuals were small.

195 To investigate whether there was any association between the prevalence of
196 either post-partum blues or *T. gondii* seropositivity, analyses were conducted on a
197 breakdown of occupational status of the participants based on the questionnaire data
198 (Table 3). The participants were categorized into 10 occupational type groups:
199 farmers (most are less well educated and live in rural areas), workers (born in the city,
200 most are less well educated and tend to be employed in factories and companies),
201 medical staff (well-educated, with a good understanding of health and hygiene),

202 managers (born in the city and most are well-educated but may not have an
203 understanding of health issues), teachers (well-educated, have health awareness but
204 may not have detailed knowledge), business staff (may or may not be well-educated,
205 most of them do not have an understanding of health issues), self-employed (may or
206 may not be well-educated, most of them do not have an understanding of health
207 issues), housewives (most are less well educated and are responsible for the routine
208 work in the family), unemployed (most are less well educated and have been out of
209 work for the past few years) and others (no details of their working status). There was
210 no significant difference in anti-*T. gondii* seroprevalence ($p = 0.971$) and incidence of
211 postpartum blues ($p = 0.918$) between the different occupational groups. We again
212 recognize that there are small numbers of individuals sampled in this analysis.

213 We also investigated traditional risk factors associated with the seroprevalence
214 of *T. gondii* and prevalence of postpartum blues in new mothers (Table 4).
215 Surprisingly, our analyses showed that contact with cats and consumption of raw or
216 uncooked meat, two generally well established risk factors, were not associated with
217 *T. gondii* seropositivity ($p = 0.766$ and 0.357 , respectively). By contrast, consumption
218 of raw or uncooked meat was significantly associated with increased prevalence of
219 postpartum blues among new mothers ($p = 0.007$) albeit based on small numbers. The
220 prevalence of postpartum blues in the group of participants that consumed raw meat
221 was significantly higher than the group that did not consume raw meat (32.97%
222 [30/91] vs 19.79% [76/394], OR: 1.993, 95% CI: 1.204-3.229). In addition, the
223 association between cat contact and prevalence of postpartum blues in new mothers is

224 close to significance at $p = 0.061$ suggesting that a larger sample size would be useful
225 to explore this further. The prevalence in the cat-contact group (29.47%, 28/95) is
226 slightly higher than that of the non-cat-contact group (20.53%, 78/380; OR: 1.618;
227 95%CI: 0.975-2.685).

228

229 **4. Discussion**

230 Depression during the perinatal period does not only have a significant impact on
231 quality of life of the mother (Darcy et al., 2011), but also influences the
232 developmental outcomes of their children (Ertel et al., 2011; Turney, 2012). Many
233 previous studies have tried to identify the role of *T. gondii* in general neurological and
234 psychiatric conditions in humans (Torrey et al., 2007; Arling et al., 2009; Groer et al.,
235 2011; Hinze-Selch et al., 2010; Prandota, 2014; Sutterland et al., 2015). Clear
236 associations seem to be emerging for schizophrenia (Torrey et al., 2007, Sutterland et
237 al., 2015) and probable associations with attempted suicide (Arling et al., 2009) and
238 personality disorders (Hinze-Selch et al., 2010). A major detailed meta-analysis
239 provided convincing evidence of an association between *T. gondii* infection and
240 schizophrenia but also suggested links with bipolar disorder, obsessive-compulsive
241 disorder, addiction but not for major depression (Sutterland et al., 2015). Several
242 other studies also demonstrated no link with major depression (Gale et al 2014; Gale
243 et al 2016; Suvisaari et al 2017). Some studies have focused specifically on prenatal
244 depression. Groer et al (2011) conducted a study on women during pregnancy and
245 showed that there was no significant difference in *T. gondii* seroprevalence between

246 mothers with prenatal depression and controls. However, they observed that higher
247 titers of *T. gondii* IgG antibody were positively correlated with increased measures of
248 depression and anxiety (as measured by the Profile of Mood States –POMS –
249 method). Furthermore, other studies have shown no association between the
250 seroprevalence of *T. gondii* and prenatal depression in both low (5.25%) (Alvarado-
251 Esquivel et al., 2017) and in high (59%) (Nourollahpour Shiadeh et al., 2016)
252 seroprevalence of infection. The latter study demonstrated that there was an
253 association of increasing *T. gondii* IgG titer with increasing depression score (as
254 measured by the Iranian version of the Edinburgh Post-Partum Depression Scale
255 (EPDS) (Nourollahpour Shiadeh et al., 2016) however, no such association was found
256 in the former one (EPDS, Mexican version) (Alvarado-Esquivel et al., 2017). The
257 consensus seems to be emerging that there is no association of *T. gondii*
258 seroprevalence and prenatal depression although conflicting evidence exists with
259 regard to association between serointensity and depression severity score. To our
260 knowledge, no studies have been conducted that investigate any possible association
261 between *T. gondii* infection and postpartum depression or postpartum blues, the latter
262 being considered as a self-limiting mild depression commonly found in new mothers
263 a few days after delivery (Maliszewska et al., 2016). In our study, we report the first
264 investigation addressing this question. Using a cohort of 475 participants, we
265 analyzed the association between *T. gondii* infection and postpartum blues. Our
266 results show clearly that overall there was no significant association between
267 seropositivity of *T. gondii* and new mothers with postpartum blues. This lack of

268 association, between seroprevalence and postpartum blues, follows the same pattern
269 revealed by studies on prenatal depression during pregnancy, described above, and
270 supports a consensus that there may be no general association with perinatal
271 depression. Furthermore, when we broke our data down further, we found no
272 evidence of association between increasing IgG titer and depression severity score (as
273 measured by the Hamilton Depression Scale). Although we do recognize that the low
274 prevalence of *T. gondii* infection and prevalence of postpartum blues in our cohort
275 reduces the power of such a detailed analysis. Again, though, our data, on the
276 quantitative association between *T. gondii* IgG titer and depression score, is consistent
277 with some other studies (Alvarado-Esquivel et al., 2017). However, both our results
278 and those of Alvarado-Esquivel et al. (2017) conflict with those of Nourollahpour
279 Shiadeh et al. (2016) on this quantitative association. This could be due, in part, to
280 differences in either depression scoring methods or due to differing backgrounds of
281 the prevalence of parasite infections. The former is unlikely, since both the studies of
282 Alvarado-Esquivel et al (2017) and Nourollahpour Shiadeh et al (2016) used variants
283 of the EPDS. However, the background of prevalences in each study differed
284 considerably (5.25% and 59%) with the former corresponding to our reported
285 prevalence in our cohort (5.68%) suggesting that this could be a factor. In China, the
286 seroprevalence of *T. gondii* in the Chinese population (~10%) is much lower than that
287 in some parts of South America and Europe (50%-80%) (Fromont et al., 2009;
288 ONIHPDCSS, 2005). In pregnant mothers in China, the prevalence is also low
289 (~10%) and corresponds to the general population levels (Gao et al., 2012). In this

290 study, the prevalence followed the typical low level found in China (5.68%). The
291 correspondence between the background prevalence in the study of Alvarado-
292 Esquivel et al (2017) and our study, both finding no quantitative association between
293 *T. gondii* IgG titer and depression scores, supports a view that the conflict with the
294 study of Nourollahpour Shiadeh et al (2016) could be related to prevalence. This also
295 raises the issue that studies conducted in areas of low parasite prevalence may need
296 substantially larger samples sizes to achieve the required power to conduct more
297 detailed breakdowns of interactions. The question is clearly complex and it is
298 possible that other interacting factors may confound any role that *T. gondii* may
299 possess in perinatal depression.

300 If *T. gondii* infection is not the main factor causing postpartum blues nor
301 depression in new mothers, are there any other factors, such as age, which could
302 influence the seroprevalence of *T. gondii* or baby blues. Seroprevalence of *T. gondii*
303 has been shown to that increase with age (Hinze-Selch et al., 2007). For instance, in
304 the Israeli population, the seroprevalence rate of *T. gondii* is 7.6% in the 10-19 years
305 group, 31.4% in the 50-59 years group, followed by a sharp increase to 58.1% in the
306 ≥ 60 years group (Markovich et al., 2014). In addition, similar results have been
307 observed in pregnant women in Poland, where mean prevalence of IgG antibodies
308 was seen at 40.6% and increased with age with a yearly seroconversion rate of 0.8%
309 (95% CI: 0.6-1.0, $p < 0.001$) (Nowakowska et al., 2014). However, in an older study,
310 Ye and Zou (1993) reported that seroprevalence of *T. gondii* in new mothers in China
311 did not increase with age. In our study of Chinese new mothers, no age prevalence

312 increase was observed ($p = 0.610$). These latter studies may be explained by bias due
313 to a relatively narrow age window found in cohorts of new mothers/women in
314 pregnancy.

315 In our study, we investigated the influence of occupation, association with cats
316 and consumption of raw meat on the relationship between *T. gondii* infection and
317 postpartum blues. Our data shows no significant association between these risk
318 factors and infection or postpartum blues. However, we recognise that once our data
319 was broken down to these levels of detail, sample sizes were small and future studies
320 may be needed with increased statistical power to dissect these questions.

321 To our knowledge, this is the first study that has investigated the association of
322 *T. gondii* infection with postpartum blues in new mothers. We found no association
323 between the seroprevalence of *T. gondii* and clinically depressed participants. Other
324 studies, that have focused on prenatal depression during pregnancy, have also failed
325 to detect significant association with *T. gondii* seroprevalence. We propose, therefore,
326 that, the current studies are tending towards a consensus that shows little support for
327 the involvement of this parasite in perinatal depression in general. However, perinatal
328 depression has a complex etiology and future larger scale studies may be required to
329 unpick further detail and to investigate specific epidemiological interactions.

330

331 **Authors' contributions**

332 Jiang-Mei Gao and Zhao-Rong Lun they were responsible for designed research;

333 Jiang-Mei Gao and Yi-Ting Xie conducted research; Zhi-Hui He provided essential

334 materials; De-Hua Lai conducted analyzed data or performed statistical analysis;

335 Jiang-Mei Gao, Geoff Hide, De-Hua Lai and Zhao-Rong Lun wrote paper.

336

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341

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345

346 **Conflict of interest**

347 The authors report no conflict of interest.

348

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493 **Table 1.** Detection of anti-*Toxoplasma gondii* IgM and anti-*T. gondii* IgG antibodies in new mothers with baby blue and in the control group.

494

| | <i>Toxoplasma gondii</i> IgM | | <i>Toxoplasma gondii</i> IgG | | |
|-----------------------|------------------------------|-------|------------------------------|--------------------|-----------------|
| | | Total | Total | postpartum blues | Normal |
| No. examined | | 475 | 475 | 106 | 369 |
| No. positive | 0 | | 27 | 7 | 20 |
| Prevalence(95% CI), % | | 0 | 5.68(3.59-7.77) | 6.60(1.80-11.41) | 5.42(3.10-7.74) |
| OR (95% CI) | — | | Ref. | 1.234[0.507-3.002] | |
| <i>P</i> -value | — | 0.64 | | | |

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<25 57 2 (0) 3.51(Ref. 0.872 57 14 24.56 Ref. 0.610
0- (13.0
8.43) 4-
36.08
)

510

25- 274 17 6.20(1.819 274 63 22.990.917
30 (5) 3.33-(0.40 (17.9(0.47
9.08) 8- 8- 1-
8.102 28.011.784
)))

511

| | | | | | | | | |
|-----------|-----|--------|---------------------|---------------------|-----|-----|------------------------|--------------------|
| 30- 35 | 113 | 6 (2) | 5.31(1.11-9.51) | 1.542(0.301-7.894) | 113 | 25 | 22.12(14.35-29.90) | 0.873(0.413-1.845) |
| >35 | 31 | 2 (0) | 6.45(0-15.61) | 1.897(0.254-14.167) | 3 4 | | 12.90(0.40-25.40) | 0.455(0.136-1.527) |
| All | 475 | 27 (7) | 5.68(3.59- 7.77) | | 475 | 106 | 22.32(18.56- 26.07) | |

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519 **Table 3.** Occupation demographics associated with seropositivity of *Toxoplasma gondii* and prevalence of postpartum blues in new mothers.

520

| Occupation | <i>Toxoplasma gondii</i> IgG | | | | post-partum blues | | | | |
|------------|------------------------------|--------------|----------------------|------------|-------------------|--------------|--------------|----------------------|------------|
| | No. examined | No. positive | Prevalence(95% CI),% | OR(95% CI) | P-value | No. examined | No. positive | Prevalence(95% CI),% | OR(95% CI) |

521

| | | | | | | | | | | |
|------|----|---|-------------|---------------|----|---|-------------------|------------|--|--|
| Farm | 22 | 2 | 9.09(22.14) | (0.971-41.75) | 22 | 5 | 22.73(3.71-41.75) | Ref. 0.918 | | |
|------|----|---|-------------|---------------|----|---|-------------------|------------|--|--|

522

| | | | | | | | | | | |
|--------|----|---|------------|---------------|----|----|--------------|-------|--|--|
| Worker | 51 | 2 | 3.92(9.44) | (0.408-3.101) | 51 | 11 | 21.57(33.25) | 0.935 | | |
|--------|----|---|------------|---------------|----|----|--------------|-------|--|--|

523

Medi 36 2 5.56(0.588 36 6 16.670.680
 cal 0- (0.07 (3.88(0.18
 staff 13.42 7- - 0-
) 4.507 29.462.565
))

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| | | | | | | | | |
|---------|----|---|---------------|--------------------|----|---|-------------------|--------------------|
| Manager | 14 | 1 | 7.14(0-22.57) | 0.769(0.063-9.371) | 14 | 2 | 14.29(0-35.25) | 0.567(0.094-3.423) |
| Teacher | 18 | 1 | 5.56(0-17.28) | 0.588(0.049-7.067) | 18 | 4 | 22.22(0.95-43.50) | 0.971(0.218-4.323) |

525

Busi 66 3 4.55(0.476 66 14 21.210.915
 ness 0- (0.07 (11.0(0.28
 staff 9.71) 4- 9- 7-
 3.055 31.342.916
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Self- 69 6 8.70(0.952 69 18 26.091.200
 empl 1.88- (0.17 (15.4(0.38
 oyed 15.51 8- 6- 7-
) 5.097 36.713.725
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 es 0.00) 3- 92.29 2-
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12.75 15.40
 0)* 6)

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| | | | | | | |
|------|----|---|------------|----|---|------------|
| Une | 47 | 3 | 6.38(0.682 | 47 | 9 | 19.150.805 |
| mplo | | | 0- (0.10 | | | (7.47(0.23 |
| yed | | | 13.64 6- | | | - 4- |
| | | |) 4.405 | | | 30.832.765 |
| | | |) | | |)) |

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|------|-----|---|------------|-----|----|------------|
| Othe | 145 | 7 | 4.83(0.507 | 145 | 34 | 23.451.041 |
| rs | | | 1.30-(0.09 | | | (16.4(0.35 |
| | | | 8.36) 8- | | | 7- 8- |
| | | | 2.615 | | | 30.433.032 |
| | | |) | | |)) |

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536 **Table 4.** Risk factors associated with the seropositivity of *Toxoplasma gondii* and prevalence of postpartum blues in new mothers.

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IgG

538

| No. examined | No. positive (with depression) | Prevalence(95% CI),% | OR(95% CI) | P-value |
|--------------|--------------------------------|----------------------|------------|---------|
| Cat Contact | No 380 21 (4) | 5.53(3.22-7.83) | Ref. | 0.766 |

539

| | | | | |
|----------|--------------|-----------------|--------------------|-------|
| Raw Meat | Yes 95 6(3) | 6.32(1.33-11.3) | 1.152(0.452-2.940) | |
| | No 384 20(2) | 5.21(2.98-7.44) | Ref. | 0.357 |

540

Yes 91 7(5)7.691.51
(2.17(0.
1- 621
13.2 -
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4)

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550 **Figure legends.**

551

552 **Figure S1A.** Quantitative comparison between *Toxoplasma gondii* immunoglobulin
553 (IgG) optical density (OD) and Hamilton Depression Scale (HAMD) score in all new
554 mothers (n=475). There is no significant correlation between IgG optical density and
555 HAMD score ($p = 0.872$).

556

557 **Figure S1B.** Quantitative analysis of all *Toxoplasma gondii* seropositive new mothers
558 (n=27) by comparison of immunoglobulin (IgG) optical density (OD) and Hamilton
559 Depression Scale (HAMD) score in the same mothers. There is no significant
560 correlation between IgG optical density and HAMD score ($p= 0.214$).

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