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## *Toxoplasma gondii* Infection:

### Relationship With Aggression in Psychiatric Subjects

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#### ABSTRACT

**Objective:** *Toxoplasma gondii* (*T. gondii*), a protozoan parasite that persists in host tissues, including brain, has been associated with several psychiatric disorders and with suicidal behavior. We sought to test the hypothesis that latent *T. gondii* infection, as manifest by circulating immunoglobulin G (IgG) antibodies to *T. gondii*, is associated with both categorical and dimensional measures of aggression.

**Method:** IgG antibodies to *T. gondii* were collected between 1991 and 2008 from 358 adult subjects with DSM-5 intermittent explosive disorder (IED), non-IED psychiatric disorders (psychiatric controls), or no evidence of any psychiatric diagnosis (healthy controls). Assessments of aggression, anger, and impulsivity, as well as state/trait anger, depression, and anxiety were completed. *T. gondii* seropositive status (IgG > 12 IU) was the primary outcome measure for this study.

**Results:** *T. gondii* seropositive status (IgG > 12 IU) was associated with higher aggression ( $P = .022$ ) and impulsivity ( $P = .05$ ) scores. When both aggression and impulsivity scores were controlled for, however, only aggression scores were higher in seropositive subjects ( $P = .011$ ). In addition, *T. gondii* seropositive status and marginal mean  $\pm$  SE aggression scores increased from healthy controls (9.1% and  $-0.66 \pm 0.05$ ) to psychiatric controls (16.7% and  $-0.27 \pm 0.05$ ) to subjects with IED (21.8% and  $1.15 \pm 0.06$ ;  $P \leq .05$ ). These findings were not accounted for by the presence of other syndromal/personality disorders or by states or traits related to depressed or anxious moods.

**Conclusions:** These data are consistent with previous studies suggesting a relationship between *T. gondii* and self-directed aggression (ie, suicidal behavior) and further add to the biological complexity of impulsive aggression both from a categorical and a dimensional perspective.

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*Toxoplasma gondii* (*T. gondii*) is a highly successful neurotropic protozoan parasite, infecting any warm-blooded animal including approximately one-third of all humans.<sup>1</sup> Within the animal world, felids have been identified as the definitive host of *T. gondii* that localizes only in the gastrointestinal tract of any member of the cat family. Humans may be infected by *T. gondii* via ingestion of the parasite's oocysts, which can spread from the feces of infected cats. Other routes of transmission include consumption of undercooked meat that has been infected with *T. gondii* cysts or ingestion of contaminated water<sup>2,3</sup>; congenital infection, occurring if a mother has a primary infection during pregnancy and transmits *T. gondii* to the fetus, is relatively rare. Postnatal chronic "latent" infection is very common, minimally symptomatic in the immune competent host, and with an encephalitic picture in the immunocompromised.<sup>4</sup> When ingested by an intermediate host, the parasite uses leukocytes to travel from the intestine to other organs, finally localizing in muscle and brain. Once in the brain, *T. gondii* hides within neurons and glial cells, forming characteristic cystic intracellular structures under the pressure of the immune system.<sup>5</sup> Although it is thought to be relatively harmless in immunocompetent adults, latent toxoplasmosis has been linked to several psychiatric disorders (eg, schizophrenia,<sup>6,7</sup> bipolar disorder,<sup>7–9</sup> personality disorders<sup>10</sup>) and with suicidal behavior.<sup>11–17</sup>

Given the strong relationship between suicidal behavior and impulsive aggressive behavior,<sup>18</sup> either as a dimension or as a category, and in light of a recent study<sup>19</sup> that reported that *T. gondii* seropositivity status may be associated with high self-reported trait aggression and impulsivity in mentally healthy individuals, we hypothesized that the categorical presence of immunoglobulin G (IgG) antibodies to *T. gondii* would (1) be associated with higher aggression and impulsivity scores in a sample of psychiatric and healthy control subjects and (2) be more frequent in individuals with intermittent explosive disorder (IED), a disorder of recurrent, problematic, and impulsive aggressive behavior, compared with healthy controls. In this study, we used psychometric assessments<sup>20–23</sup> as the dimensional representation of impulsive aggression and the presence of IED as the categorical representation<sup>24</sup> of impulsive aggression.

#### METHOD

##### Subjects

Three hundred fifty-eight physically healthy subjects participated in this study. All subjects were systematically evaluated in regard to aggressive, anxiety, and other behaviors as part of a larger program that is designed to study correlates of

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**Table 1. Psychiatric and Personality Disorder Diagnoses in Psychiatric Controls and in Subjects With Intermittent Explosive Disorder<sup>a</sup>**

Diagnosis	Psychiatric	Intermittent	P Value
	Controls (n = 138)	Explosive Disorder (n = 110)	
Lifetime syndromal disorders	108 (78.3)	109 (99.1)	<.001*
Any depressive disorder	49 (35.5)	69 (62.7)	<.001*
Any anxiety disorder	32 (23.2)	33 (30.0)	.226
Any substance use disorder	37 (26.8)	57 (51.8)	<.001*
Intermittent explosive disorder	0 (0.0)	109 (99.1)	<.001
Stress and trauma disorders	20 (14.5)	21 (19.1)	.333
Eating disorders	3 (2.2)	12 (10.9)	.004
Obsessive-compulsive disorders	3 (2.2)	3 (2.7)	.778
Other impulse-control disorders	0 (0.0)	4 (3.6)	.226
Any personality disorder	89 (64.5)	109 (99.1)	<.001*
Cluster A	12 (8.7)	27 (24.5)	.001*
Cluster B	19 (13.8)	63 (57.3)	<.001*
Cluster C	22 (15.9)	33 (30.0)	.008
Personality disorder, not otherwise specified	46 (33.3)	31 (28.2)	.384

<sup>a</sup>All values are n (%). Subjects may have more than 1 disorder.

\*Significant after Bonferroni correction for 14 comparisons ( $P < .003$ ).

impulsive aggressive and other personality-related behaviors in human subjects. Subjects were recruited through public service, newspaper, and other media announcements seeking individuals who (1) reported psychosocial difficulty related to syndromal psychiatric and/or personality disorder conditions or (2) had little evidence of any psychopathology. All subjects gave informed consent and signed the informed consent document approved by the first author's (E.F.C.) Institutional Review Board.

### Diagnostic Assessment

Syndromal and personality disorder diagnoses were made according to *DSM-5* criteria.<sup>25</sup> Diagnoses were made using information from (1) the Structured Clinical Interview for *DSM-IV* Axis I disorders (SCID-I)<sup>26</sup> for syndromal disorders and the Structured Interview for *DSM-IV* Personality (SIDP-IV)<sup>27</sup> for personality disorders, (2) a clinical interview by a research psychiatrist, and (3) the review of all other available clinical data. The research diagnostic interviews were conducted by individuals with a master's or doctor's degree in clinical psychology, blinded to the study hypothesis. All diagnostic raters went through a rigorous training program that included lectures on *DSM* diagnoses and rating systems, videos of expert raters conducting SCID-I/SIDP-IV interviews, and practice interviews and ratings until the raters were deemed reliable by the trainer. This process resulted in good to excellent interrater reliabilities (mean  $\pm$  SE  $\kappa = 0.84 \pm 0.05$ ; range, 0.79 to 0.93) across anxiety, mood, substance use, impulse control, and personality disorders. Final diagnoses were assigned by team best-estimate consensus procedures<sup>28,29</sup> involving research psychiatrists and clinical psychologists as previously described.<sup>30</sup> This methodology has been shown to enhance the accuracy of diagnosis over direct interview alone.<sup>31</sup> Subjects with a current history of a substance use disorder or a life history of bipolar disorder, schizophrenia

- While limited information exists on the causes of human aggression, new research is pointing to inflammatory or infectious processes as a possible etiology.
- Patients with significant histories of aggression (ie, intermittent explosive disorder) may have a latent infection with *T. gondii*, a common protozoan, that often goes undetected but is treatable.

(or other psychotic disorder), or mental retardation were excluded from study.

After diagnostic assignment, 110 subjects had no evidence of any psychiatric diagnosis (healthy controls); 138 subjects met criteria for a lifetime diagnosis of a syndromal psychiatric or personality disorder, but not for a lifetime diagnosis of IED (psychiatric controls); and 110 subjects met criteria for a lifetime diagnosis of IED. Of the 248 subjects with any *DSM-5* diagnosis, most (72.4%) reported (1) a history of formal psychiatric evaluation and/or treatment (46.3%) or (2) a history of behavioral disturbance during which the subjects, or others, thought they should have sought mental health services but did not (26.1%). Syndromal psychiatric and personality disorder diagnoses are listed in Table 1.

### Psychometric Measures of Aggression, Impulsivity, and Related Behaviors

Aggression was assessed with the Aggression score from the Life History of Aggression (LHA)<sup>20</sup> assessment and the Aggression (Physical and Verbal) score from the Buss-Perry Aggression Questionnaire (BPAQ).<sup>21</sup> The LHA assesses the history of actual aggressive behavior and the BPAQ assesses aggressive tendencies as a personality trait. Impulsivity was assessed with the Barratt Impulsiveness Scale, version 11 (BIS-11)<sup>22</sup> and the Impulsivity scale from the Eysenck Personality Inventory-2 (EPQ-2).<sup>23</sup> Both BIS-11 and EPQ-2 Impulsivity assess a person's disposition to act impulsively as a personality trait. Life history of suicidal and self-injurious behavior was assessed during the diagnostic interviews. An act was considered a suicide attempt if it involved behavior with the conscious (even if ambivalent) intent to die by means that the subject believed could end his or her life; an act was considered self-injurious if it involved behavior with the conscious (even if ambivalent) intent by the subject to physically harm, but not kill, himself or herself. Other assessments included the State-Trait Anger Expression Inventory-2 (STAXI-2)<sup>32</sup> for state and trait anger, Beck Depression Inventory-II (BDI-II)<sup>33</sup> for state depression, Depression scale from the General Behavior Inventory (GBI)<sup>34</sup> for trait depression, Beck Anxiety Inventory (BAI)<sup>35</sup> for state anxiety, and State-Trait Anxiety Inventory (STAI)<sup>36</sup> for trait anxiety. The Global Assessment of Functioning (GAF)<sup>37</sup> scale served as the variable for psychosocial functioning. Racial data, collected by diagnostic assessors, reflected self-identified racial characteristics of subjects. Socioeconomic status was assessed by the Hollingshead method (A. B. Hollingshead, *Four Factor Index of Social Status*, unpublished dissertation, 1975).

**Table 2. Demographic, Psychometric, and Clinical Characteristics of Sample**

Variable	Healthy Controls (n=110)	Psychiatric Controls (n=138)	Intermittent Explosive Disorder (n=110)	P Value
<b>Demographic variables</b>				
Age, mean ± SD, y	31.3 ± 8.7	33.7 ± 8.1	36.1 ± 8.3	<.001 <sup>a</sup>
Male, %	58	59	64 <sup>b</sup>	.748 <sup>b</sup>
Race (white/African American/other), %	64/26/10	66/30/4	65/28/7	.379 <sup>b</sup>
SES score, mean ± SD	37.0 ± 12.8	36.5 ± 14.5	35.6 ± 12.2	.719 <sup>a</sup>
<b>Psychometric variables, mean ± SD</b>				
Aggression (LHA)	4.5 ± 3.6	7.2 ± 5.2	18.3 ± 5.3	<.001 <sup>a</sup>
Aggression (BPAQ)	40.8 ± 13.4	47.7 ± 14.8	74.8 ± 19.7	<.001 <sup>a</sup>
Impulsivity (BIS-11)	56.3 ± 8.4	62.0 ± 9.7	69.3 ± 12.0	<.001 <sup>a</sup>
Impulsivity (EPQ-2)	3.4 ± 3.1	4.4 ± 3.8	8.5 ± 4.4	<.001 <sup>a</sup>
State anger (STAXI-2)	15.2 ± 1.0	16.3 ± 1.4	22.7 ± 1.1	<.001 <sup>a</sup>
Trait anger (STAXI-2)	13.2 ± 2.5	16.3 ± 5.4	26.6 ± 7.2	<.001 <sup>a</sup>
State depression (BDI-II)	2.4 ± 7.4	6.7 ± 7.7	16.5 ± 11.9	<.001 <sup>a</sup>
Trait depression (GBI)	0.1 ± 0.5	8.8 ± 13.6	14.0 ± 11.5	<.001 <sup>a</sup>
State anxiety (BAI)	22.6 ± 2.2	27.9 ± 8.6	32.3 ± 9.4	<.001 <sup>a</sup>
Trait anxiety (STAI)	29.7 ± 6.9	38.5 ± 11.6	47.1 ± 10.2	<.001 <sup>a</sup>
<b>Clinical variables, mean ± SD</b>				
GAF score	84.0 ± 4.9	65.1 ± 10.6	54.6 ± 8.6	<.001 <sup>a</sup>
Suicide attempt history	NA	3.6%	26.4%	<.001 <sup>c</sup>
Self-injurious behavior history	NA	6.5%	15.5%	.023 <sup>c</sup>
Suicidal or self-injury history	NA	7.2%	36.4%	<.001 <sup>c</sup>

<sup>a</sup>By analysis of variance.

<sup>b</sup>By  $\chi^2$ .

<sup>c</sup>By Fisher exact test.

Abbreviations: BAI=Beck Anxiety Inventory; BDI-II=Beck Depression Inventory-II; BIS-11=Barratt Impulsivity Scale, version 11; BPAQ=Buss-Perry Aggression Questionnaire; EPQ-2=Eysenck Personality Inventory-2; GAF=Global Assessment of Functioning; GBI=General Behavior Inventory; LHA=Life History of Aggression; SES=socioeconomic status; STAXI-2=State-Trait Anger Expression Inventory-2; STAI=State-Trait Anxiety Inventory.

### Assessment of *T. gondii* Seropositivity Status

Subjects were free of all medications for at least 4 weeks. Whole blood, anticoagulated with EDTA (ethylenediaminetetraacetic acid), was obtained between 9 AM and 11 AM through venipuncture of a forearm vein. Plasma was processed after centrifugation, placed in a polypropylene tube, and stored at  $-80^{\circ}\text{C}$  until assay. These frozen plasma samples were collected between 1991 and 2008 and were tested for IgG antibodies to *T. gondii* in 2014 by solid-phase enzyme-linked immunosorbent assay (ELISA) with kits from IBL (Hamburg, Germany). All samples were run in duplicate, and quality controls were used. The coefficient of intra-assay variation was less than 10%. A subject with plasma *T. gondii* IgG antibodies > 12 IU was considered to be seropositive for *T. gondii*. Equivocal samples (8–12 IUs) were reanalyzed to accurately classify them as negative or positive. The laboratory technician was not aware of the diagnostic status of the subject. *T. gondii* seropositive status, rather than serointensity, was used because the latter cannot be measured in seronegative subjects. Despite the 18-year duration of sample collection, no association was observed between *T. gondii* seropositive status and time from the first to last study year (Spearman  $\rho=0.08$ ,  $P=.12$ ). Finally, plasma levels for the proinflammatory cytokine interleukin 6 (IL-6) were available in 176 of these subjects as part of a previously published study.<sup>38</sup>

### Statistical Analysis

Comparisons of between-group variables were performed by  $\chi^2$ , univariate (ANOVA/ANCOVA), and multivariate analysis of variance/covariance (MANOVA/MANCOVA), followed by Tukey honestly significant difference post hoc testing. Other analyses included binary logistic regression using age as a covariate. The primary biological variable in this study was *T. gondii* seropositive status as in previous studies.<sup>19</sup> The primary dimensional variables included composite scores for aggression and impulsivity; these variables were created by z-transforming each of the 2 sets of aggression (LHA/BPAQ) and impulsivity (BIS-11/EPQ-2) variables and taking the mean z score of the source variables. The primary categorical variables included diagnostic status (healthy controls/psychiatric controls/subjects with IED), positive history of suicide attempt, and positive history of self-injurious behavior. A 2-tailed  $\alpha$  value of .05 was used to denote statistical significance for all analyses. The primary analysis tested the relationship between *T. gondii* seropositive status and composite aggression scores, composite impulsivity scores, and history of self-directed aggression in all subjects. This was followed by analyses to determine if (1) *T. gondii* seropositive status was more frequent among IED subjects (subjects characterized by high levels of impulsive aggressive and suicidal behaviors), (2) *T. gondii* seropositive status was greater among subjects with other psychiatric and personality disorders, and (3) *T. gondii* seropositive individuals had higher levels of state and/or trait depression, anxiety, or anger.

## RESULTS

### Demographic Characteristics of the Sample

Healthy controls, psychiatric controls, and IED subjects differed only in age, with a less than 5-year age difference between IED subjects and healthy controls and a less than 3-year age difference between IED subjects and psychiatric controls. *T. gondii* seropositive status did not differ as a function of sex, race, or socioeconomic status (Table 2). As expected, the groups differed as a function of aggression, impulsivity, and history of suicidal and self-injurious behavior (Table 2). The rate of *T. gondii* seropositive subjects in this study was 15.9%, which is comparable with the most recent estimate of 14.1% for the United States.<sup>9</sup>

### *T. gondii* Seropositive Status as a Function of Aggression and Impulsivity

One-way ANCOVA, with age as a covariate, revealed that Composite Aggression scores were significantly higher in *T. gondii* seropositive subjects ( $F_{1,342}=5.32$ ,  $P=.022$ ; Figure 1, left). Composite Impulsivity scores were also higher in *T. gondii* seropositive subjects ( $F_{1,325}=3.83$ ,  $P=.05$ ; Figure 1, right). Composite



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Aggression scores adjusted for Composite Impulsivity scores continued to be higher among *T. gondii* seropositive subjects ( $P = .011$ ; Figure 1, left), although the reverse was not true for Composite Impulsivity scores adjusted for Composite Aggression scores ( $P = .984$ ; Figure 1, right). Similarly, when placed in the same binary logistic regression, with age as covariate, Composite Aggression ( $B = 0.44 \pm 0.20$ ,  $Wald_1 = 4.77$ ,  $Exp(B) = 1.55$ ,  $P = .029$ ), but not Composite Impulsivity ( $B = 0.02 \pm 0.20$ ,  $Wald_1 = 0.08$ ,  $Exp(B) = 1.06$ ,  $P = .773$ ), scores were associated with *T. gondii* seropositive status.

### T. gondii Seropositive Status as a Function of Self-Directed Aggression

Binary logistic regression, with age as covariate, revealed that *T. gondii* seropositive status did not predict history of suicide attempt (20.6% vs 15.4% *T. gondii* seropositive:  $B = -0.26 \pm 0.46$ ,  $Wald_1 = 0.31$ ,  $P = .577$ ), history of self-injurious behavior (26.9% vs 15.1% *T. gondii* seropositive:  $B = -0.75 \pm 0.47$ ,  $Wald_1 = 2.48$ ,  $P = .116$ ), or history of either type of self-directed aggressive behavior (24.0% vs 14.6% *T. gondii* seropositive:  $B = -0.58 \pm 0.37$ ,  $Wald_1 = 2.42$ ,  $P = .119$ ).

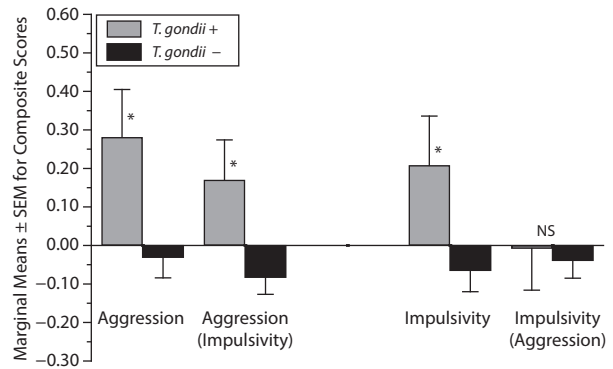
### T. gondii Seropositive Status as a Function of IED

Next, we examined the relationship between *T. gondii* seropositive status as a function of IED, a disorder of recurrent, problematic, impulsive aggressive behavior, compared with healthy controls and psychiatric controls. A significant difference in *T. gondii* seropositive status was noted among the groups (linear by linear association  $\chi^2_1 = 6.06$ ,  $P = .014$ ; Figure 2). Binary logistic regression, with age as a covariate, yielded the same result with IED subjects ( $B = 0.89 \pm 0.41$ ,  $Wald_1 = 4.68$ ,  $P = .030$ ) significantly associated with *T. gondii* seropositive status compared with healthy controls. Psychiatric controls were not significantly associated with *T. gondii* seropositive status compared with healthy controls ( $B = -0.63 \pm 0.41$ ,  $Wald_1 = 2.42$ ,  $P = .120$ ). Increasing *T. gondii* seropositive status paralleled mean Composite Aggression (and Composite Impulsivity) scores, adjusted for age, across the groups (Figure 2). Adding Composite Aggression scores to the model eliminated the relationship for *T. gondii* seropositive status for IED subjects, compared with healthy controls ( $B = 0.26 \pm 0.63$ ,  $Wald_1 = 0.18$ ,  $P = .675$ ).

### T. gondii Seropositive Status as a Function of Non-IED Syndromal Disorders

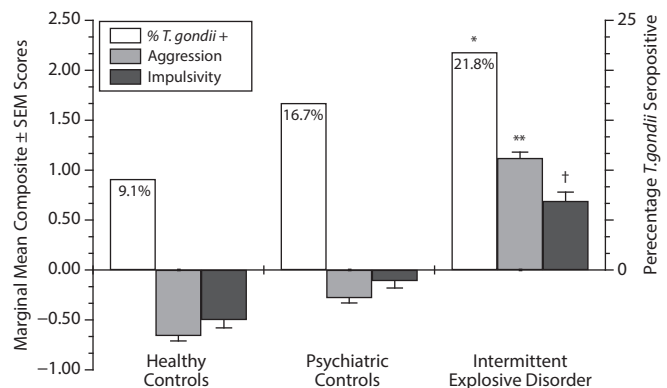
A significant difference in *T. gondii* seropositive status was also noted as a function of lifetime depressive disorder and lifetime anxiety disorder, but not lifetime substance use disorder. Binary logistic regression, with age as covariate, yielded similar results for subjects with depressive disorder ( $B = 0.79 \pm 0.42$ ,  $Wald_1 = 3.51$ ,  $P = .061$ ) and subjects with anxiety disorder

Figure 1. Composite Aggression and Impulsivity (age as covariate) in Subjects Seropositive (+) and Seronegative (-) for *T. gondii*<sup>a</sup>



<sup>a</sup>Aggression (impulsivity) refers to Composite Aggression scores with Composite Impulsivity scores as a covariate; impulsivity (aggression) refers to Composite Impulsivity scores with Composite Aggression scores as a covariate.  
\* $P \leq .05$ .  
Abbreviation: NS = not significant.

Figure 2. Percentage of *T. gondii* Seropositive (+) Subjects and Composite Aggression and Composite Impulsivity Scores (adjusted for age) as a Function of Healthy Controls, Psychiatric Controls, and Intermittent Explosive Disorder Status



\* $P < .05$ , based on  $\chi^2$ .  
\*\* $P < .025$ , based on analysis of covariance (ANCOVA).  
† $P = .051$ , based on ANCOVA.

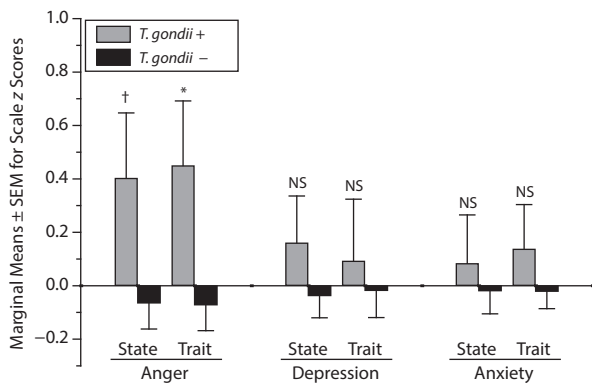
( $B = 1.05 \pm 0.47$ ,  $Wald_1 = 5.06$ ,  $P = .025$ ), compared with healthy controls. Adding Composite Aggression scores to the models eliminated these relationships for lifetime depressive disorder ( $B = 0.56 \pm 0.50$ ,  $Wald_1 = 1.26$ ,  $P = .263$ ) and lifetime anxiety disorder ( $B = 0.76 \pm 0.521$ ,  $Wald_1 = 2.13$ ,  $P = .144$ ).

### T. gondii Seropositive Status as a Function of Personality Disorder

A significant difference in *T. gondii* seropositive status was also noted as a function of the presence of borderline and/or antisocial personality disorder compared with healthy controls (healthy controls, 9.1%; psychiatric controls, 18.2%; borderline and/or antisocial personality disorder, 20.9%; linear by linear association:  $\chi^2_1 = 5.10$ ,  $P = .024$ ); the difference with psychiatric controls was not significant. Binary logistic regression, with age as covariate, yielded a similar result that approached, but did not

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**Figure 3. State and Trait Anger, Depression, and Anxiety (z) Scores (age as covariate) as a Function of *T. gondii* Seropositive (+) and Seronegative (-) Subjects<sup>a</sup>**



<sup>a</sup>To place all symptom measures on the same scale, z scores were used.

\* $P < .05$ .

† $P < .10$ .

Abbreviation: NS = not significant.

reach, statistical significance ( $B = 0.85 \pm 0.45$ ,  $Wald_1 = 3.47$ ,  $P = .062$ ). Adding Composite Aggression scores to the model eliminated this relationship ( $B = 0.32 \pm 0.60$ ,  $Wald_1 = 0.29$ ,  $P = .592$ ). Expanding this analysis to include all personality disorders resulted in the same observation with respect to psychiatric controls having a higher rate of *T. gondii* seropositive status compared with healthy controls; this finding was nonsignificant after the addition of Composite Aggression scores to the model.

### ***T. gondii* Seropositive Status as a Function of State and Trait Dysphoric Moods: Anger, Depression, and Anxiety**

ANCOVA, with age as covariate, revealed higher state and trait anger scores as a function of *T. gondii* seropositive status (Figure 3, left). Similar analysis using state and trait depression and anxiety scores, however, revealed no significant differences as a function of *T. gondii* seropositive status (Figure 3, center and right).

### **Circulating IL-6 Levels as a Function of *T. gondii* Seropositive Status**

ANCOVA, with age as covariate, revealed no difference in circulating levels of IL-6 as a function of *T. gondii* seropositive status in the subgroup in which these data were available (*T. gondii* seropositive:  $0.98 \pm 1.64$  pg/mL vs *T. gondii* seronegative:  $1.41 \pm 1.67$  pg/mL;  $F_{1,172} = 1.27$ ,  $P = .262$ ). As previously reported, IL-6 (log IL-6:  $r = -0.39$ ,  $P < .001$ ) levels were inversely correlated with Composite Aggression scores in the subjects in this study.<sup>39</sup>

## **DISCUSSION**

The results of this study suggest a relationship between latent infection with *T. gondii* and impulsive aggression from both a dimensional and categorical perspective. Specifically, *T. gondii* seropositive status was associated with higher

scores on the psychometric measures for both Aggression and Impulsivity. Between aggression and impulsivity, these data suggest that *T. gondii* seropositive status is primarily related to aggression than to impulsivity in that the variance associated with impulsivity overlaps with the variance associated with aggression. In addition, the rate of *T. gondii* seropositivity in IED subjects was significantly greater than that in healthy controls, though not significantly greater than that in psychiatric controls without IED. The nonsignificant difference in seropositivity rate between IED subjects and psychiatric controls may be due to the fact that Aggression scores in psychiatric controls were intermediate between healthy controls and IED subjects, that this sample did not have the power to detect this difference to a statistically significant degree, or that other behaviors, such as depression or anxiety, are also associated with latent toxoplasmosis. These results are supported by findings from animal studies<sup>39</sup> that showed a relationship between *T. gondii* infection and elevated aggression-related behaviors and a recent study<sup>19</sup> of 1,000 psychiatrically healthy subjects that documented elevated trait aggression and impulsivity as a function of *T. gondii* seropositivity.

Typically, other-directed aggression is strongly associated with self-directed aggressive behavior in psychiatric subjects,<sup>18</sup> and greater rates of *T. gondii* seropositive status have been reported among those with a history of suicidal behavior.<sup>11-15</sup> Despite these previous findings, we did not find an association between *T. gondii* seropositive status and self-directed aggression in our sample. The proportion of subjects with lifetime histories of suicidal or self-injurious behavior was small, however, and the present study had limited statistical power to detect relationships reported from previous, and larger, samples. It is also possible that the psychiatric diagnostic composition of the sample (psychiatric controls) and tendency to direct aggression outward (IED) reduced an association with suicidal self-directed aggression. Consistently, in the largest study<sup>14</sup> on *T. gondii* seropositive status and suicidal self-directed violence, performed on a cohort of Danish women, the association was significantly weaker in those women who had a concurrent diagnosis of mood disorder, psychotic disorder, or personality disorder.

Individuals with a lifetime history of depressive and anxiety disorder also had higher rates of *T. gondii* seropositive status compared with healthy controls. While higher depression and anxiety scores should be observed as a function of *T. gondii* seropositive status, no significant differences in state or trait scores for depression or anxiety were observed. The observed effect size for depression or anxiety scores was modest ( $d = 0.10$  to  $0.20$ ), and it is possible that a larger sample would have yielded different results. However, studies in much larger samples report no significant association between *T. gondii* seropositive status and unipolar major depression or dysthymia,<sup>9</sup> generalized anxiety disorder, panic disorder, or posttraumatic stress disorder,<sup>39</sup> suggesting that mood and anxiety disorders are not accounting for the findings in our study. In contrast, composite aggression and state and trait anger scores were

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significantly elevated as a function of *T. gondii* seropositive status and, in every case, eliminated all differences as a function of *T. gondii* seropositive status. Thus, we posit that the higher *T. gondii* seropositive rates observed in individuals with depressive/anxiety disorder, compared with healthy controls, were due to their comorbidity with IED or a correlation between aggression, depression, and anxiety scores. In the current sample, IED was highly comorbid with lifetime depressive disorder (64% vs 35%,  $P < .001$ ), and aggression scores correlated with both trait depression ( $r = 0.38$ ,  $P = .001$ ) and anxiety ( $r = 0.52$ ,  $P < .001$ ) scores, though not as strongly as depression correlated with anxiety ( $r = 0.74$ ,  $P < .001$ ).

Several factors may account for these findings. First, chronic latent infection with *T. gondii* may lead to a low-grade chronic immune activation within the brain, with (or without) downstream effects on neurotransmitter systems involved in aggressive behavior.<sup>40</sup> Second, chronic *T. gondii* infection may alter the structure and function of corticolimbic circuits that are known to modulate impulsive aggressive behavior.<sup>41</sup> Specifically, persistent *T. gondii* infection in mice is associated with neuronal tissue lesions, altered neuronal function, ventricular dilation, and neuroinflammation.<sup>42</sup> In addition, several, though not all, studies suggest that *T. gondii*-containing cysts localize primarily in the prefrontal cortex and amygdala<sup>43,44</sup> and that latent infection with *T. gondii* induces dendritic retraction in the basolateral amygdala.<sup>45</sup> Third, as shown experimentally in rats,<sup>46</sup> *T. gondii* infection increases testicular expression of genes involved in the production of testosterone. In addition, there is evidence that *T. gondii*-infected males, though not females, have higher circulating levels of testosterone compared with controls.<sup>47</sup> However, while a number of studies report a relationship between elevated levels of testosterone and aggression,<sup>48</sup> the magnitude of this relationship is small. Thus, it is unlikely that testosterone plays any more than a modest role in this regard.

Neurotransmitter mechanisms by which *T. gondii* may affect behavior include effects on serotonergic and glutaminergic transmission, both of which have been shown to play a role in aggressive behavior in human studies.<sup>49,50</sup> Relevant to serotonin, conversion of tryptophan to kynurenine is controlled by indoleamine 2,3-dioxygenase ([IDO]; IDO-1 and IDO-2).<sup>51</sup> Since IDO can be activated by inflammatory cytokines, levels of kynurenine can rise while levels of serotonin decline. In addition, increased levels of kynurenine lead to increased levels of its active metabolite quinolinic acid, a potent *N*-methyl-D-aspartate receptor agonist, which may increase the risk for aggressive behavior in humans.<sup>50</sup> While this hypothesis is partially supported by reported elevations of kynurenine and quinolinic acid levels in mice with chronic *T. gondii* infection,<sup>52</sup> we did not find differences in circulating levels of proinflammatory cytokines (ie, IL-6) as a function of *T. gondii* seropositivity. It is possible that the proinflammatory processes that keep *T. gondii* in a latent state are confined to the brain and are not reflected in the periphery. It is also possible that impulsively aggressive

individuals engage in behaviors that increase their own risk of infection with *T. gondii* or that latent toxoplasmosis changes behavior, as in felids,<sup>1</sup> so that the expression of aggression is increased. In addition, *T. gondii* is known to increase risk-taking behavior in rodents, evolutionarily benefiting the parasite (ie, transforming natural aversion in cats to attraction).<sup>53,54</sup> This is an example of the general phenomenon of host manipulation by parasites, documented in nature,<sup>55</sup> and proposed as a model with some explanatory potential for alterations in human behavior associated with parasitic infections.<sup>56,57</sup>

The strengths of this hypothesis-driven study include a well-characterized sample of healthy and psychiatric controls as well as validated measures of aggression, impulsivity, depression, and anxiety. Limitations to our study are present, as well. First, we used a cross-sectional design, and no causal, or directional, conclusions can be made from these analyses. Second, ascertainment of subjects may limit the generalizability of these findings in that these involved subjects who volunteered for a research study, rather than for clinical treatment. However, nearly three-quarters of the psychiatric subjects reported a past history of psychiatric treatment (or of having episodes of behavioral disturbance for which they, or others, thought they should have sought mental health services but did not), and, thus, most of these subjects are likely similar to individuals who would have been recruited from a clinical setting. Third, it is possible that the presented associations are nonspecific and, instead, due to other common latency-establishing neurotropic pathogens such as herpes viruses or cytomegalovirus. However, recent studies have documented that associations between *T. gondii* and self-directed<sup>58</sup> and other-directed<sup>19</sup> aggression in human subjects are not due to the presence of these other potential pathogens. Finally, because immunoglobulin M antibodies to *T. gondii* were not assessed, there is a possibility that a small number of seropositive subjects had an acute, rather than a chronic latent, infection at time of study.

In summary, we report a greater rate of *T. gondii* seropositive status in subjects with DSM-5 IED compared with healthy controls and a positive relationship with aggression and anger, but not with depression or anxiety. These findings are consistent with previous *T. gondii* seropositive status data, suggesting a relationship with self-directed aggression (ie, suicidal behavior) and a relationship involving schizophrenia or mania—disorders in which many individuals are often aggressive.<sup>59,60</sup> Our results further add to the biological complexity of impulsive aggression, from both a categorical and a dimensional perspective.

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