

Increased Cooperative Behavior Across Remitted Bipolar I Disorder and Major Depression: Insights Utilizing a Behavioral Economic Trust Game

Desmond C. Ong and Jamil Zaki
Stanford University

June Gruber
University of Colorado Boulder

Mood disorders impact social functioning, but might contribute to experiences—like affective distress—that might result in increased cooperative behavior under certain circumstances. We recruited participants with a history of bipolar I disorder ($n = 28$), major depressive disorder ($n = 30$), and healthy controls ($n = 27$)—to play a well-validated behavioral economic Trust Game, a task that provides a well-controlled experimental scenario, to measure cooperative behavior for the first time across both groups. Both remitted mood-disordered groups cooperated significantly more than the control group, but did not differ from one another. These results suggest that, in some contexts, a history of mood disturbance can produce enhanced cooperation, even in the absence of current mood symptoms. We discuss the clinical significance of enhanced cooperation in mood disorders and point to key directions for future research.

General Scientific Summary

Individuals with mood disorders suffer disruptions in social functioning, but under the right circumstances, might also display *more* prosocial behaviors. We find that individuals in remission from bipolar I disorder and major depressive disorder cooperated more in an economic game than individuals with no such history. The results suggest that in some contexts, a history of clinical mood disturbance generates increases in some forms of prosocial behavior, even in the absence of current mood symptoms.

Keywords: cooperation, trust, bipolar disorder, major depressive disorder

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Mood disorders like bipolar I disorder (BD) and major depressive disorder (MDD) have a debilitating impact on psychosocial functioning (e.g., MacQueen, Young, & Joffe, 2001). Less understood are the mechanisms by which mood disturbance translates to psychosocial dysfunction. Recently, researchers have begun utilizing rigorous behavioral economic approaches to unpack observed social impairments by first examining simple behavioral markers in disorders, through carefully controlled experimental scenarios with clearly defined incentives (e.g., King-Casas & Chiu, 2012). Cooperation—acting in ways that benefits others—is one such behavioral marker crucial to maintaining relationships and coordination

in groups (e.g., Rand & Nowak, 2013; Zaki & Mitchell, 2013). Hence, abnormalities in cooperative behavior could be one mechanism that mediates the impact of mood disorders on broader social dysfunction, and it is critical to study whether and how such abnormalities are expressed in mood disorders. Here we use a game theoretic paradigm to provide an “intermediate” level of analysis between disorder and social impairment, by connecting BD and MDD to abnormalities in specific social behaviors that contribute to broader social functioning. Specifically, we assess cooperation for the first time across both individuals with BD and MDD currently in remission, as compared to healthy non-psychiatric controls, using a well-validated index of cooperative behavior.

Past research suggests that individuals with a history of mood disturbances may exhibit *increased* cooperative behavior when compared with healthy adults. This idea is consistent with classic work in psychology demonstrating that distress—a feature of BD and MDD (O’Connor, Berry, Weiss, & Gilbert, 2002; Shamay-Tsoory, Harari, Szepeswol, & Levkovitz, 2009)—is a strong motivation of prosocial behavior (Batson, 2011; Cialdini, Brown, Lewis, Luce, & Neuberg, 1997). Indeed, two studies using a behavioral economic game (the Ultimatum Game) found that individuals diagnosed with major depression cooperated more than healthy controls, by offering their partner a larger proportion of a pot of money (DeStoop, Schrijvers, De Grave, Sabbe, & De

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Desmond C. Ong and Jamil Zaki, Department of Psychology, Stanford University; June Gruber, Department of Psychology and Neuroscience, University of Colorado Boulder.

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Correspondence concerning this article should be addressed to June Gruber, Department of Psychology and Neuroscience, University of Colorado Boulder, 1905 Colorado Avenue, 345 UCB, Muenzinger D321C, Boulder, CO 80309. E-mail: june.gruber@colorado.edu

Bruijn, 2012), and accepting more unfair divisions from their partner (Harlé, Allen, & Sanfey, 2010). A study of individuals in remission from MDD similarly found that they cooperated more in a Prisoner's Dilemma game than healthy controls (Pulcu et al., 2015, Table 4: Welch's $t(37) = 2.16; p = .038$; see supplemental materials). It is worth noting other studies that find no differences in cooperation between healthy controls and currently diagnosed MDD individuals (Pulcu et al., 2015; Unoka, Seres, Aspán, Bódi, & Kéri, 2009), or BD individuals (Saunders, Goodwin, & Rogers, 2015) on other economic tasks.

The present investigation translated a well-validated behavioral economic game paradigm in healthy adults to provide insights into social processes impacted by a history of psychiatric mood disturbance. This work provides broader significance in several key ways. First, as described above, economic games provide tight quantification of social behaviors in psychiatric samples. This literature is small but growing, and more data is required across more contexts and symptomatic states, especially in the remitted mood phase. Second, previous studies have never previously measured both BD and MDD participants together in the same task. Here, we do so, allowing us to examine diagnostically specific, versus transdiagnostic, effects of mood disorders on specific social behaviors like cooperation.

Method

Participants

Participants were recruited as part of a broader study on emotion and mood (Yale University IRB HIC #1309012679; University of Colorado Boulder IRB #14-0390) using posted flyers, online advertisements, and referrals from outpatient mental health centers and psychiatric hospitals. Of the final 85 study participants, 28 were diagnosed with BD Type 1 in remission (i.e., not currently manic, depressed, or mixed; rBD), 30 with MDD in remission (i.e., not currently depressed; rMDD), and 27 were nonpsychiatric controls (CTL) who did not meet current or past criteria for any *Diagnostic and Statistical Manual for Mental Disorders-Fourth Edition-Text Revised (DSM-IV-TR)* Axis I disorder. We chose to examine both BD and MDD groups in a remitted mood phase to first examine potential group differences in prosocial behavior impacted even without mood symptoms (e.g., Gruber, Harvey, & Purcell, 2011). rBD and rMDD participants were not excluded on the basis of comorbid disorders (aside from current substance or alcohol use disorders) given that mood disorders are commonly comorbid with other Axis I disorders (e.g., Kessler, Chiu, Demler, & Walters, 2005), though we verified that BD was the primary, or most severe, diagnosis for the rBD group, and MDD for the rMDD group (Di Nardo, O'Brien, Barlow, Waddell, & Blanchard, 1983). Exclusion criteria for all groups included history of severe head trauma, stroke, neurological disease, severe medical illness (e.g., autoimmune disorder, cardiovascular disease, or HIV/AIDS), or current alcohol or substance abuse in the past 6 months.

Measures of Clinical Functioning

Diagnostic evaluation. A trained postbaccalaureate researcher confirmed all diagnoses using the Structured Clinical Interview for *DSM-IV* (SCID-IV; First, Spitzer, Gibbon, & Williams, 2007). A second reviewer from the research team independently rated a subset

($n = 75, 90%$) of videotaped interviews from the broader study protocol. Ratings across all Axis I disorders matched 100% ($\kappa = 1.00$) of primary diagnoses. During the SCID-IV, the interviewer collected information concerning illness duration and lifetime number of mood episodes.

Mood symptoms. We measured current symptoms of mania using the Young Mania Rating Scale (YMRS; Young et al., 1978), an 11-item, clinician-rated measure with scores ranging from 0 to 60, and current symptoms of depression using the Inventory of Depressive Symptomatology (IDS-C; Trivedi et al., 2004), a 30-item, clinician-rated measure with scores from 0 to 84. We verified current remitted mood status (i.e., neither manic, depressed, nor mixed mood state) for all groups according to both current SCID-IV criteria and cutoff scores on the YMRS (≤ 7), and IDS-C (≤ 11). Intraclass correlations (ICCs) for the same subset of participants were strong for both the YMRS ($= 0.90$) and IDS-C ($= 0.99$).

Global functioning. We used the Global Assessment of Functioning (GAF; *DSM-IV Axis V*) Scale to assess general functioning in the past week on a scale from 1 (lowest level of functioning) to 100 (highest level). The ICC for the same subset of participants ($n = 75, 90%$) was acceptable ($= 0.65$).

Cognitive functioning. We assessed cognitive functioning using the Mini-Mental Status Examination, a brief objective measure of cognitive status and impairment (MMSE; Folstein, Folstein, & McHugh, 1975). Raw scores (range: 0–30) were calculated as the total number of trials correct. All participants exceeded the eligibility cutoff score (≥ 24). We also assessed executive functioning using the letter-number sequencing subtest of the Wechsler Adult Intelligence Scale-III (WAIS-III; Wechsler, 1997). Participants are read aloud a series of increasingly long strings of randomly ordered numbers and letters, and have to verbally repeat back all numbers in ascending numerical order, followed by all letters in alphabetical order. Raw scores (range: 0–21) were calculated as the total number of trials correct, from which WAIS-III age-normed scaled scores were computed.

Behavioral Economic Cooperation Game

To objectively measure cooperative behavior, we used a standardized version of the Trust Game (TG; Figure 1). The TG is widely used in behavioral economics as an index of both trusting and trustworthy behaviors, and as a proxy of cooperative tendencies (Berg, Dickhaut, & McCabe, 1995). The task involves two players: the Investor and the Trustee. On each round, the Investor receives an endowment of \$10, and decides to invests \$X. This investment is then tripled and given to the Trustee, who decides how much (\$Y) of \$3X to return to the Investor. If the Investor chooses to invest \$X and the Trustee chooses to return \$Y, the Investor receives $\$(10 - X + Y)$, while the Trustee receives $\$(3X - Y)$. This procedure is repeated for multiple rounds.

In the present study, participants played the role of the Trustee, and believed the investor with whom they played was another participant. In fact, we designed a computer algorithm to simulate investor decisions. Previous studies using the TG and other games have similarly used computer programs to standardize conditions that participants were exposed to (e.g., McCabe, Houser, Ryan, Smith, & Trouard, 2001; Sripada et al., 2009). In iterated paradigms with repeated rounds of interactions, humans tend to react to the outcomes of previous rounds, and reinforcement-learning (RL) models serve as a good

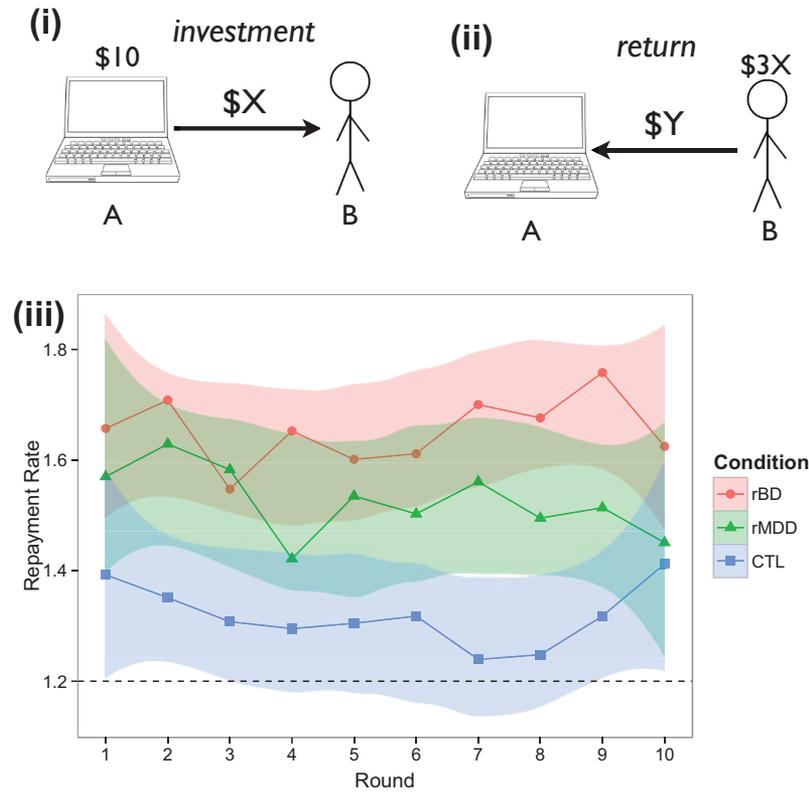


Figure 1. (i) Illustration of the investment portion of a round, where the computer chooses to invest $\$X$. (ii) Return portion: participant chooses to return $\$Y$ out of $\$3X$. (iii) Mean repayment rates ($\$Y/\X) over rounds, colored by condition. Shaded regions indicate LOWESS-smoothed 95% confidence intervals. The dashed line at 1.2 indicates the expected repayment rate (details in text).

approximation to such behavior (Dayan & Niv, 2008). Intuitively, a human investor would increase their investment if the trustee returned a decent amount the previous round, and would decrease their investment if they felt cheated. The average human trustee, from a meta-analysis of over 20,000 participants, returns an average of 37% of the total sum they were entrusted with (Johnson & Mislin, 2011). Hence, we designed a simple RL investor that “expected” participants to behave like the average trustee and return 40% of the total amount that they were entrusted with, which corresponds to 120% of the investment. If a participant returned more/less than this expectation, the investor will increase/decrease its investment on the subsequent round. Additional details are included in the supplemental materials, and code can be found at <https://github.com/desmond-ong/CooperationInMoodDisorders>.

Procedure

Study design and procedures were reviewed and approved by the University of Colorado Boulder and Yale University Institutional Review Boards. Participants arrived at the laboratory, provided written and verbal informed consent and completed diagnostic and cognitive assessment measures. Participants then completed the TG task along with other unrelated laboratory tasks as part of the larger study protocol.

Participants participated in the TG task using a computer in an individual testing room. The experimenter remained in an adjacent

room and communicated via microphone. Participants learnt that they were about to play a two-player Investment Game, and saw two stick figures representing players A and B. On each round, A receives \$10 and chooses an amount $\$X$ to invest, which is automatically tripled and given to B. B decides how much of $\$3X$ to return to A. Participants saw an example round where A invests \$7 out of \$10 (that becomes \$21), and B returns \$12 out of \$21: A ends the round with $\$10 - \$7 + \$12 = \15 , and B ends the round with $\$21 - \$12 = \$9$. Participants could repeat the instructions until they understood. Next, participants were assigned to the role of Player B, and learnt that they would be playing 10 rounds with another participant in a separate room. Participants saw a screen that claimed to be connecting their computer to their “partner.” In reality, they played with a computer program (described above). After 10 rounds, participants were debriefed, and compensated for their participation at a standard hourly rate. We did not include additional financial compensation as a function of the amount earned in the game.

Results

Demographic and Clinical Characteristics

We report demographic and clinical characteristics in Table 1. The groups did not differ with respect to age, sex, and ethnicity ($ps > 0.47$). Although participants differed with respect to income,

Table 1
Demographic and Clinical Characteristics

Sample characteristics	rBD (<i>n</i> = 28)	rMDD (<i>n</i> = 30)	CTL (<i>n</i> = 27)	Statistic
Demographic				
Age (years)	29.9 (6.8)	27.9 (6.4)	28.4 (6.5)	<i>F</i> = .765
Female (%)	64%	73%	63%	$\chi^2 = .837$
White (%)	79%	77%	74%	$\chi^2 = .156$
Education (Yrs)	14.23 (2.02)	15.95 (1.79)	15.96 (2.20)	<i>F</i> = 6.93**
Employed (%)	53.6%	70.0%	70.4%	$\chi^2 = 2.26$
Partnered (%)	42.9%	63.3%	63.0%	$\chi^2 = 1.81$
Number children	1.5 (.7)	1.1 (.4)	1.3 (.7)	<i>F</i> = 2.57
Annual income				$\chi^2 = 28.9^{**}$
<\$10K	17.9%	13.3%	25.9%	
\$10K–\$25K	46.4%	30.0%	3.7%	
\$26K–\$50K	28.6%	23.3%	33.3%	
\$51K–\$75K	0%	6.7%	11.1%	
\$76K–\$100K	3.6%	20.0%	0%	
>\$100K	3.6%	6.7%	25.9%	
Cognitive				
MMSE	29.0 (1.7)	28.4 (1.8)	29.1 (1.4)	<i>F</i> = 1.53
WAIS-III	10.8 (3.1)	11.1 (2.9)	10.4 (2.8)	<i>F</i> = .35
Clinical				
YMRS	1.32 (1.49)	1.07 (1.20)	.56 (.97)	<i>F</i> = 2.70 ^a
IDS-C	3.43 (2.33)	4.77 (3.04)	2.30 (2.05)	<i>F</i> = 6.83 ^{bc}
GAF	70.46 (7.22)	76.97 (7.11)	85.85 (6.41)	<i>F</i> = 34.08 ^{abc}
Remission duration (months)	36.95 (36.03)	31.40 (33.36)	—	—
Age at onset (years)	18.10 (5.34)	14.50 (3.04)	—	<i>F</i> = 10.2 ^b
Illness duration (years)	12.16 (5.85)	13.3 (5.17)	—	<i>F</i> = .50
No. comorbid disorders	.14 (.36)	.43 (.63)	—	<i>F</i> = 4.63 ^b
% comorbid disorders	14%	36.7%	—	$\chi^2 = 4.44$
No. depressive episodes	13.48 (17.53)	17.23 (24.10)	—	<i>F</i> = .45
No. manic episodes	13.63 (21.53)	—	—	—
No. medications	1.46 (1.26)	.60 (.77)	—	<i>F</i> = 10.1 ^b
Anticonvulsants	39.3%	3.33%	—	
Lithium	10.7%	0%	—	
Neuroleptics	35.7%	0%	—	
Stimulants	0%	3.33%	—	
Antidepressants	35.7%	43.3%	—	
Benzodiazepines	10.7%	3.33%	—	
Sedative-hypnotics and other anxiolytics	14.3%	6.67%	—	

Note. rBD = remitted bipolar disorder group; rMDD = remitted major depressive disorder group; CTL = healthy control group; employed = employed full-time or part-time; partnered = married or in a relationship; YMRS = Young Mania Rating Scale; IDS-C = Inventory of Depression Symptomatology-Clinician Rating; GAF = Global Assessment of Functioning; age at onset = age of first depressive or manic episode; No. comorbid disorders = the number of current *Diagnostic and Statistical Manual for Mental Disorders-Fourth Edition-Text Revised (DSM-IV-TR)* Axis I comorbidities; % comorbid disorders = at least one *DSM-IV-TR* Axis I comorbidity; No. medications = the number of psychotropic medications currently taken (including anticonvulsants, lithium, neuroleptics, stimulants, antidepressants, benzodiazepines, and sedative-hypnotics and other anxiolytics). Mean values are given with *SDs* in parentheses.

^a *p* < .05 for rBD and CTL. ^b *p* < .05 for rBD and rMDD. ^c *p* < .05 for rMDD and rCTL.

** *p* < .01.

($\chi^2(10) = 28.9, p = .001$), income did not affect any of the behavioral results (*ps* > 0.50), and our main results regarding group differences still hold after excluding those with more than \$100,000 annual income. All groups also scored well below standardized cutoffs on the YMRS (≤ 7) and IDS-C (≤ 11). There was a marginally significant group main effect suggesting differences in YMRS scores (*p* = .073). The rMDD group scored higher than the rBD (*p* = .047) and CTL (*p* < .001) groups on subsyndromal depressive symptoms (IDS-C), and rBD and CTL groups did not differ from each other (*p* = .10). The rBD and rMDD groups scored lower on global functioning (GAF) than the CTL group (*p* < .001 for both), and the rBD group also scored lower than the rMDD group (*p* < .001) on GAF. Participants did not differ on either MMSE scores (*p* = .22), or WAIS-III scores (*p* = .70).

Additionally, the rBD and rMDD groups did not differ in illness duration (*p* = .45), though the rMDD group had an earlier average age of onset (*p* = .002).

Cooperative Behavior on the Trust Game

To compare returns across rounds with different investments, we calculated a *repayment rate*, the proportion of the investment the participant returns (i.e., \$Y/\$X). There were no significant trends of repayment rates over rounds ($\chi^2(3) = 2.24, p = .52$). To account for the repeated nature of the data, we ran linear mixed-effects models with condition as fixed effects, and with random intercepts and random slopes (on rounds) by participant (Table 2). Across all rounds, rBD participants displayed a significantly

Table 2
Multiple Regression Analyses of Behavior in Trust Game

Model (1)	Repayment rate (amount repaid/amount invested)				
	Coefficient	SE	95% CI	<i>t</i>	<i>p</i>
rBD (> CTL)	.303	.125	[.059, .548]	2.43	.017*
rMDD (> CTL)	.241	.123	[.0002, .481]	1.96	.053
rBD (> rMDD)	.063	.122	[-.176, .301]	.52	.608
Marginal R^2			.049		
Conditional R^2			.677		

Summary measures	Total amount invested in participant (2)	Total amount repaid by participant (3)	Total amount kept by participant (4)	Total amount kept by investor (5)
rBD (> CTL)	12.41 (6.01)*	34.7 (16.3)*	2.52 (8.28)	22.3 (11.3)
rMDD (> CTL)	5.94 (5.91)	22.2 (16.0)	-4.37 (8.14)	16.3 (11.1)
R^2	.050	.054	.009	.049

Note. Model (1) is a linear mixed-effects model predicting repayment rate with clinical history as fixed effects, and random intercepts and slopes (on round number) by participant. The rBD > rMDD contrast was calculated by re-running the model with rMDD as the base group. Marginal and Conditional R^2 are reported, after Johnson (2014). Models (2–5) predict summary measures calculated after the end of all rounds; they are fixed effects only linear models, as there are no repeated measurements. The units on the unstandardized coefficients for Models (2–5) are in terms of hypothetical dollars, with SE in parentheses. Note these summary measures are partly correlated with the repayment rate, but they also provide an estimate of the behavioral effect size across the whole task. rBD = remitted bipolar disorder group; rMDD = remitted major depressive disorder group; CTL = healthy control group; CI = confidence interval.

* $p < .05$.

greater repayment rate, repaying 30.3% more of the investment, than control participants ($b = 0.303$, 95% confidence interval [0.059, 0.548], $t(82) = 2.43$, $p = .017$). rMDD participants similarly repaid more than control participants, with an effect almost reaching significance ($b = 0.241$ [0.0002, 0.481], $t(82) = 1.96$, $p = .053$). There was no difference between the repayment rates of rBD and rMDD participants ($p = .61$). Overall, clinical history explained a significant proportion of variance (marginal $R^2 = .049$ for fixed effects variance only, and conditional $R^2 = .677$ for fixed and random effects variances; after Johnson, 2014).

Next, we calculated several key measures that represent meaningful summaries of behavior: (a) the total amount invested in each participant after all 10 rounds, (b) the total amount repaid by each participant, (c) the total amount kept by each participant, and (d) the total amount kept by the investor. Note that these quantities are all partly correlated with the participant's repayment rate, as the simulated investor followed a deterministic algorithm, up to random variations in learning rates. We regressed these summary measures on condition using simple linear models (Table 2). Recall that the investor can invest up to \$10 per round for 10 rounds for a potential maximum of \$100. On average, rBD participants encouraged the investor to invest \$12.41 more as compared with control participants ($b = 12.41$ [0.64, 24.19], $t(82) = 2.07$, $p = .04$), and repaid \$34.72 more to the investor ($b = 34.72$ [2.87, 66.57], $t(82) = 2.14$, $p = .04$). The investor earned marginally more if they were playing with a rBD participant than a control participant ($b = 22.31$ [0.24, 44.38], $t(82) = 1.98$, $p = .051$); however, rBD participants did not earn more than control participants ($p = .76$). Thus, the investor that played against the average rBD participant would have invested more, and “earned” slightly more after 10 rounds, than if they had played against the average CTL participant. rMDD participants did not differ significantly from controls along any of these four measures ($ps > 0.15$). In summary, not only did participants with a history of BD differ

in their own behavior as compared with controls, they also encouraged more trusting behavior for “people” who play with them, resulting in better outcomes for their partners.

Discussion

Utilizing a behavioral economic trust game, we found that a history of mood disturbance is associated with *increased* cooperative behavior when individuals are trusted with an investment. Specifically, both rBD and rMDD groups in our study repaid more to a simulated investor in the Trust Game than a healthy control group. These results are consistent with the idea that although mood disturbance, such as affective distress, generates hardship for individuals, they might, in some contexts, also lend themselves to other-oriented, cooperative behavior (Batson, 2011; Batson, O'Quin, Fultz, Vanderplas, & Isen, 1983; Cialdini et al., 1997; O'Connor et al., 2002). Our findings are consistent with previous work showing increased cooperation in current and remitted MDD on other economic tasks (DeStoop et al., 2012; Harlé et al., 2010; Pulcu et al., 2015).

These data generate an interesting dilemma: If mood disorders are associated with increased cooperation in the lab, why do they also track strained social functioning outside the lab? One possibility is that economic games simplify social interactions, emphasizing interpersonal expectations more than real-world encounters, and individuals with MDD and BD might respond to these clear social cues. In our study, another player trusted participants with an investment, and so our participants might feel more compelled to repay that trust by cooperating. This is consistent with another examination of compliance in cooperation in MDD: Using a modified Trust Game in which the investor added an explicit request for a specific amount of repayment, Zhang, Sun, and Lee (2012) found that currently depressed patients repaid the exact requested amount more frequently than healthy controls, even when they had

the freedom to repay more or less than the requested amount. By contrast, real-world contexts are much more nuanced than economic games. People have to rely on complex cues about others' emotions when deciding how to direct their cooperative behavior (Zaki & Ochsner, 2012). To the extent that mood disorders hamper these individuals' ability to understand complex social signals (Devlin, Zaki, Ong, & Gruber, 2016; Samamé, Martino, & Strejlevich, 2012), such individuals may be unable to express their cooperative motives.

More broadly, increased cooperative behavior observed in bipolar and major depressive disorders may have downstream maladaptive consequences. For example, individuals may cooperate with good intentions but produce negative consequences for themselves or those they intended to help (e.g., starving oneself to reduce one's burden on caregivers; Oakley, Knafo, Madhavan, & Wilson, 2011). Being excessively cooperative and/or trusting may also leave such individuals more vulnerable to exploitation by others; this could, in turn, generate life stressors that are known contributors to exacerbated depressive and manic symptom levels. Notwithstanding, these clinical risks, it is possible that increased cooperative behavior might afford important opportunities. For example, therapists could work with patients to scaffold more structure in patients' social relationships to channel this cooperative motivation adaptively into group settings (e.g., at work or with family). Much work remains to unpack the potentially deleterious aspects of excessive cooperation in mood disorders, as well as ways to capitalize upon its strengths.

The present investigation should be carefully interpreted within the confines of several caveats. First, although the TG task is a well-validated task, it represents only one form of cooperation, and does not capture all the nuances of cooperation in daily life. Future work should obtain parallel assessments of real-world social functioning to provide direct links between experimental outcomes and social functioning in daily life. Second, it is worth noting that game theoretic tasks operationalize social phenomena as *behaviors*, irrespective of the underlying psychological mechanisms (Camerer, 2003). Cooperative behaviors can reflect many sources, including distress (Batson et al., 1983) but also reputation management (Harbaugh, 1998), reciprocity (Trivers, 1971), and compliance with social norms (Nook, Ong, Morelli, Mitchell, & Zaki, 2016). Future work should more closely examine these mechanisms and their effects on prosociality in mood disorders. Third, to obtain ecologically valid samples, we did not exclude participants from the BD or MDD clinical groups on the basis of comorbidities. Future studies with larger sample sizes should examine whether the effects of disorders on cooperation varies as a function of such comorbidities to rule out the possibility that the observed findings are not better attributed to comorbidity versus a history of mood disturbance.

Fourth, both BD and MDD participants were remitted at the time of testing. The use of a remitted sample enables the identification of areas of potential preservation during remission, but it also leaves open critical questions about how cooperative behavior covaries as a function of current mood symptom severity. Future work should carefully examine the relationship between current symptom status and cooperation. Fifth, the present sample sizes are respectable given the severe nature of the psychiatric groups recruited and complexity of the measured variables. However, there may still have been insufficient statistical power to assess

within-group affective correlates of cooperative behaviors, between-groups differences between BD and MDD groups, and other more complex interaction effects. Future studies (or meta-analyses) should also examine cooperative behavior in larger sample sizes and across psychopathology more broadly, such as in borderline personality disorder (King-Casas et al., 2008; Unoka et al., 2009), or anxiety (Rodebaugh, Heimberg, Taylor, & Lenze, 2016; Sripada et al., 2009). Sixth, given the challenges of accessing an unmedicated clinical sample, we were unable to investigate the influence of medication status on cooperation. Future studies should explore larger sample sizes to assess both functional correlates of cooperation within and across diagnoses and the effects of medication status. Finally, although our study was cross-sectional, it is important for future prospective studies to examine the potential protective effects of increased cooperative behavior on symptom remission and mood relapse.

In conclusion, although BD and MDD are characterized by disruptions in social behavior and relationships, here, we document increased cooperation in an economic game among individuals in remission from these disorders. We hope that this work, and future studies, will provide a deeper understanding of the social consequences of affective disorders, and inform treatment options that leverage cooperation to improve psychosocial functioning.

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