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1 **Title: Learning to trust: Social feedback normalizes trust behavior in first episode psychosis**
2 **and clinical high-risk**

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26

Trust in CHR and psychosis

27 Abbreviations

28 FEP = First episode psychosis

29 CHR = Clinical high-risk

30 CAARMS = Comprehensive Assessment of At-Risk Mental States

31 PANSS = Positive and Negative Syndromes Scale

32 WAIS = Wechsler Adult Intelligence Scale

33 ROI = Region of interest

34 TPJ = Temporo-parietal junction

35 mPFC = medial prefrontal cortex

36

37 Abstract

38 Background: Psychosis is characterized by problems in social functioning that exist well before
39 illness onset, and in individuals at clinical high-risk (CHR) for psychosis. Trust is an essential
40 element for social interactions that is impaired in psychosis. In the trust game, chronic patients
41 showed reduced baseline trust, impaired response to positive social feedback, and attenuated brain
42 activation in reward and mentalizing areas. We investigated whether first episode psychosis
43 patients (FEP) and CHR show similar abnormalities in the neural and behavioral mechanisms
44 underlying trust.

45 Methods: Twenty-two FEP, 17 CHR, and 43 healthy controls performed two trust games, with a
46 cooperative and an unfair partner in the fMRI scanner. Region of interest analyses were performed
47 on mentalizing and reward processing areas, during the investment and outcome phases of the
48 games.

49 Results: Compared to healthy controls, FEP and CHR showed reduced baseline trust, but like
50 controls, learned to trust in response to cooperative and unfair feedback. Symptom severity was not
51 associated with baseline trust, however in FEP associated with reduced response to feedback. The
52 only group differences in brain activation were that CHR recruited the temporo-parietal junction
53 (TPJ) more than FEP and controls during investment in the unfair condition. This hyper-activation
54 in CHR was associated with greater symptom severity.

55 Conclusions: Reduced baseline trust may be associated with risk for psychotic illness, or generally
56 with poor mental health. Feedback learning is still intact in CHR and FEP, as opposed to chronic
57 patients. CHR however show distinct neural activation patterns of hyper-activation of the TPJ.

58 Keywords:

59 First episode psychosis; clinical high-risk; trust; social feedback; fMRI

60 1 Introduction

61 Psychosis is characterized by problems in social functioning (Couture et al, 2006; Fett et al, 2012).
62 Lower social functioning is already present in childhood in individuals who continue to develop
63 psychosis and has also been reported in individuals at high-risk for psychosis (Ballon et al, 2007;
64 Corcoran et al, 2011; Cornblatt et al, 2007; Velthorst et al, 2016a; Velthorst et al, 2016b; Yung et al,
65 2003). Clinical high-risk patients (CHR) are already in care for other psychopathology, reporting
66 psychotic-like symptoms, but have not yet experienced – and maybe never will – full-blown
67 psychosis. In CHR, the developmental course of social functioning is predictive of the conversion to
68 psychosis (Cannon et al, 2008; Jang et al, 2011; Niendam et al, 2007). Understanding the
69 mechanisms underlying deficits in social functioning in at-risk states and first-episode psychosis
70 (FEP) is crucial for understanding transition and outcome prognosis. Intervening at these early
71 stages targeting social functioning can improve outcome and possibly delay (or prevent) transition.

72 Social functioning relates to establishing relationships, both vocational and private
73 (Velthorst et al, 2016a; Velthorst et al, 2016b). Patients show a steep decline in these domains
74 starting about five years before illness onset. The basis of social functioning is the ability to interact
75 in an appropriate way with other people. Previous research studying online social interactions in
76 psychosis has suggested two possible explanatory mechanisms for impairments in social
77 interactions; these are a reduced sensitivity to rewarding effects of social contact (Campellone et al,
78 2016; Fett et al, 2012; Gromann et al, 2013), and an impaired social cognitive ability (Csukly et al,
79 2011; Horat et al, 2017), including impaired mentalizing (Green et al, 2015). Social cognitive skills
80 (Couture et al, 2006; Green & Leitman, 2008) are necessary for the formation and maintenance of
81 relationships and for building trust in other people. Like patients with psychosis, CHR show deficits
82 in a variety of these skills (Bora & Pantelis, 2013; Lavoie et al, 2013; McCleery et al, 2014), albeit to
83 a lesser degree. Research has mainly focused on off-line cognitive skills, without investigating them

84 in real interactions. In the last decade, interactive designs have been widely used, that have the
85 strength to capture social cognitive skills, as well as the rewarding effects of social behavior in an
86 on-line setting. We therefore investigated cooperative and unfair social interactions and the neural
87 correlates of trust, directly comparing FEP and CHR to controls, using an interactive trust game to
88 test whether these groups display similar underlying mechanisms of reduced social interactions.

89 The trust game investigates real-time social interactions (Berg et al, 1995). In the game, the
90 first player (investor), receives a certain endowment, e.g., €10. He or she can give any amount
91 between €0 and €10 to the second player, the trustee. The given amount is tripled and the trustee
92 then can return any part of this amount to the investor. The best pay-off for the trustee is reached
93 by keeping the money. Thus, investing requires trust that a fair repayment will be made. The
94 iterative game allows for the investigation of baseline trust (i. e. first investment), and the
95 development of trust based on cooperative and unfair social feedback. Key-processes involved in
96 the trust game are thought to be mentalizing (Declerck et al, 2013; Frith & Frith, 2006; Gallagher &
97 Frith, 2003) and reward processing (Fehr & Camerer, 2007; King-Casas et al, 2005; Rilling & Sanfey,
98 2011). Mentalizing appears to be important during both the investment and repayment phase,
99 where estimations of the other's behavior are made. Reward learning signals have been shown to
100 shift from the repayment phase to the investment phase in an iterative trust game (King-Casas et al,
101 2005). Hence, we investigated both the investment and repayment phase (Figure 1).

102 Research in healthy subjects has shown that participants initially invest more than half of
103 their endowment (Berg et al, 1995; Johnson & Mislin, 2011). Studies from our lab have shown that
104 baseline trust tends to be lower in patients than controls (Fett et al, 2016; Gromann et al, 2013).
105 Both positive (Fett et al, 2012) and negative (Fett et al, 2016) symptoms have been associated with
106 lower baseline trust, suggesting that reduced trust may reflect either paranoia or a lack of social
107 motivation. The ability to learn from social feedback seems to depend on context (cooperative or

108 unfair partner's responses) and illness duration: Early psychosis patients were able to adjust their
109 trust to similar levels as controls, whereas chronic patients showed an insensitivity to positive
110 feedback. In unfair interactions, early and chronic psychosis patients responded adequately to
111 negative feedback (Campellone et al, 2016; Fett et al, 2015; Fett et al, 2016; Fett et al, 2012;
112 Gromann et al, 2013). Understanding the mechanisms of trust in early psychosis stages may
113 provide insights in focal points to target in social functioning interventions.

114 At the neural level, reduced caudate activation in chronic patients has been reported in
115 cooperative interactions. Relatives of patients with psychosis, despite behavioral outcomes similar
116 to controls, also showed reduced recruitment of the caudate and insula. These results possibly
117 reflect reduced sensitivity to social reward processing mechanisms in both patients and relatives
118 (Gromann et al, 2013; Gromann et al, 2014), which could account for social impairments.
119 Associations of neural activity with positive symptoms have been reported (Gromann et al, 2013).

120 This study set out to investigate whether CHR and FEP patients, similar to chronic patients,
121 show reduced baseline trust and to explore the neural mechanisms underlying trust behavior in
122 these patient groups. Based on the existing trust game literature, we hypothesized that similar to
123 relatives (Fett et al, 2012) and (chronic) patients (Fett et al, 2015) 1) FEP and CHR will show lower
124 levels of baseline trust, and 2) CHR and FEP are able to learn from positive and negative feedback
125 given by the counterpart and adjust their levels of trust accordingly (Fett et al, 2015; Gromann et al,
126 2013). In both cases, we expected CHR to perform in between FEP and controls. In addition, the
127 associations of symptoms with baseline trust and changes in trust in FEP and CHR were examined.
128 On the neural level we hypothesized to find 3) attenuated activation in brain areas associated with
129 mentalizing and reward (learning) in FEP compared to controls. Based on the trust literature in
130 relatives and imaging research in CHR (Smieskova et al, 2013), we expected intermediate activation
131 in CHR. Based on the findings by Gromann et al (2013), we hypothesized to find 4) positive

132 symptoms related positively to brain activation in mentalizing and negatively to reward areas in
133 both patient groups. In addition, associations of brain activations with negative symptoms were
134 investigated.

135

136 2 Methods

137 2.1 Subjects

138 Twenty-six FEP patients with non-affective psychosis, aged 16-22, and 17 CHR, aged 16-31, were
139 recruited in the Amsterdam and The Hague area. Forty-nine healthy control participants (aged 16-
140 31) were recruited to match both patient populations on age and gender. Patients were contacted
141 through their caregivers at the academic medical center Amsterdam (AMC), the Amsterdam early
142 intervention team psychosis and PsyQ, The Hague. FEP were diagnosed at the AMC, according to the
143 DSM-IV criteria (Association & Association, 2000). FEP were included within 18 months of the
144 diagnosis (mean 5.6 months). CHR were help seeking individuals that were referred to PsyQ by
145 their general practitioners or other mental health institutions. After an initial diagnosis based on
146 their complaints, all new admissions (between age 14-35) were screened for an “at-risk mental
147 state” (ARMS) with the Comprehensive Assessment of At-Risk Mental States (CAARMS) (Yung et al,
148 2005), a semi-structured interview that assesses psychotic experiences in the last year before
149 assessment. Only the positive symptoms sub-scale was used, and both intensity and frequency of
150 the symptoms were assessed. Patients met ARMS criteria for attenuated psychotic symptoms (a)
151 with subthreshold intensity, when scoring 3-5 on severity and 3-6 on frequency of symptoms (all
152 CHR participants), (b) or based on subthreshold frequency, when scoring 5-6 on severity and 3 or
153 above on frequency (N = 2, in combination with intensity), or based on vulnerability, i.e.,
154 schizotypal personality disorder, familial history of psychosis, or a drop in social functioning (N = 3,

155 in combination with intensity) (for details, see (Yung et al, 2005), p. 966). Patients were diagnosed
156 as having a full-blown psychosis when scoring higher than 6. Additionally, patients had to display
157 marked problems in socially useful activities (work and study), relationships, and self-care,
158 indicated by a score below 55 on the Social and Occupational Functioning Assessment Scale
159 (SOFAS; mean score 46.9) (Goldman et al, 1992; Morosini et al, 2000), see also (Rietdijk et al,
160 2012). CHR were included within one year after CAARMS assessment (mean 4.8 months). Fourteen
161 CHR patients also took part in a larger study (EU-GEI), with post-measurements at 6, 12 and 24
162 months using the CAARMS. Of two CHR participants, follow-up data are missing. Symptoms of
163 depression and anxiety are often the primary presenting complaints of CHR patients, rather than
164 the attenuated psychotic symptoms (Modinos et al, 2014). Similar to other CHR samples (Fusar-Poli
165 et al, 2014; Kelleher et al, 2012; Morrison et al, 2012; Wigman et al, 2012; Woods et al, 2009), the
166 current CHR sample had comorbid diagnoses of anxiety (5), personality (3), eating (2) and mood
167 (2) disorders, trauma (2), and ADHD (3). Exclusion criteria for all participants were an IQ < 80 and
168 contraindications for scanning. For FEP additional exclusion criteria were a primary diagnosis of a
169 mood disorder, and comorbidity with autism spectrum disorder. Healthy control participants were
170 excluded if they had a (family) history of psychopathology, which was assessed with self-report,
171 and by a systematic interview with questions regarding past and present mental help seeking,
172 depressed and psychotic symptoms, and intake of medication.

173 We excluded four FEP and six controls due to invalid behavioral data (1 FEP and 2 controls),
174 unusable or missing imaging data (3 FEP and 4 controls). The analysis sample consisted of 22 FEP,
175 17 CHR and 43 controls.

176 2.2 Measures

177 2.2.1 *Trust game*

178 Participants played the role of investor in two multi-round trust games. They were told that they
179 were connected to their anonymous counterpart via the Internet. In reality, they played against a
180 computer, programmed to respond either in a cooperative or in an unfair way. In the cooperative
181 condition, the return was 100%, 150% or 200% of the invested amount, with increasing likelihood
182 of a 200% repayment after each increase of investment. In the unfair condition the return was 75%
183 or 50%, with increasing likelihood of a 50% repayment after increase of investment. The two
184 games were presented in counterbalanced order. Each game consisted of 20 experimental and 20
185 control trials (Figure 1). For a detailed description of the paradigm see Lemmers-Jansen et al
186 (2017). After the trust game, a questionnaire to investigate participants' opinions on the behavior
187 of their counterpart was administered, to check if participants believed that they were playing a
188 real person.

189 *2.2.2 Positive and Negative Syndrome Scale (PANSS; Kay et al, 1987)*

190 The 30-item PANSS semi-structured interview was used for rating symptoms in the two weeks
191 prior to testing. The PANSS distinguishes between positive, negative, and general symptoms (Kay et
192 al, 1987). Items are scaled on a 7-point Likert scale, ratings 3 and higher indicating clinical values.
193 All FEP and 13 CHR completed the interview. Interviews were taken by four researchers, and audio
194 tapes (and if consented video tapes) were made. Responses were rated on the basis of the
195 recordings and notes taken during the interview, by two researchers. Based on the first participants
196 an interrater reliability was calculated ($r = .85$). All PANSS data were rated by the same two
197 researchers.

198 *2.2.3 Wechsler Adult Intelligence Scale (WAIS; Wechsler, 1997)*

199 To control for confounding effects of intelligence, the vocabulary subscale of the WAIS was
200 included, a measure of verbal comprehension, consisting of 33 words that had to be defined by the

201 participants. Answers were coded as fully correct (2 points), partially correct (1) or wrong (0),
202 resulting in a maximum score of 66.

203 2.2.4 Trust Manipulation Check Questionnaire

204 The trust game was followed by a questionnaire to investigate participants' opinions on the
205 behavior of their counterpart, and to check if they believed that they were playing a real person
206 ("What do you think this task was about?", "What were the main causes that influenced your
207 behavior during the task?", "Did you use a strategy during the tasks? If so, which strategy?", and
208 "Did you think that the counterparts made fair choices?"). The last questions required answers
209 specified per counterpart. If a participant referred to the two counterparts as persons in his/her
210 responses, we regarded the manipulation as successful. If participants reported on any of the
211 questions that they had doubts, or did not believe that the counterpart was real, the manipulation
212 was coded as failed. Four controls, three FEP and one CHR did not believe the manipulation.

213

214 2.3 Procedure

215 This research was approved by the Medical Ethical Committee of the VU Medical Center
216 Amsterdam. All participants signed an informed consent, and completed several questionnaires
217 that are not included in the current paper. We then administered the PANSS to FEP and CHR
218 patients. Prior to scanning, participants received oral and written instructions for the trust game,
219 illustrated with screenshots of the game to illustrate the task. Participants played several practice
220 rounds on a computer, accompanied by additional feedback, to ensure understanding of the task.
221 Subsequently participants were scanned for about an hour. First, participants performed the trust
222 game, followed by the structural scan. After this period of relative rest, they performed a second
223 task (the Social Mindfulness Paradigm, to be reported in a separate paper (in preparation)),

224 followed by a resting state scan. Immediately after scanning, a manipulation check for the trust
225 game was administered (see 2.2.4). Participants received an image of their brain, 25€ for
226 participation and reimbursement of their travel costs.

227 2.4 fMRI Data Acquisition.

228 Imaging data were obtained at the Spinoza Center Amsterdam, using a 3.0 T Philips Achieva whole
229 body scanner (Philips Healthcare, Best, The Netherlands) equipped with a 32 channel head coil. A
230 T2* EPI sequence (TR = 2.31, TE = 27.63, FA = 76.1°, FOV 240mm, voxel size 2.5 x 2.5 x 2.5, 40
231 slices, 0.3 mm gap) was used, which resulted in 325 images per condition. A T1-weighted scan was
232 obtained for anatomical reference (TR = 8.2, TE = 3.8, FA = 8°, FOV 240*188mm, voxel size 1 x 1 x 1,
233 220 slices).

234 2.5 Data Analysis

235 2.5.1 Behavioral data

236 Demographic and behavioral data were analyzed using Stata 13 (StataCorp, 2013) with regression
237 analyses and chi-square tests. We analyzed group differences in first investment (baseline trust),
238 and the development of investments (changes in trust) across repeated interactions (indicated by
239 trial number) in each game (cooperative and unfair). We used multilevel random regression
240 analyses to account for multiple observations [investments (level 1); within participants (level 2)].
241 All analyses were controlled for age and WAIS score.

242 2.5.2 Imaging data

243 Imaging data were analyzed using Statistical Parametric Mapping 8 (SPM, 2009). Functional images
244 for each participant were preprocessed as follows: realign and unwarp, coregistration with
245 individual structural images, segmented for normalization to an MNI template and smoothing with
246 a 6 mm Gaussian kernel (FWHM). At fist-level, a general linear model was used to construct

247 individual time courses for the investment and repayment phase per condition, using an event-
248 related design. For each trial we defined the investment as the period of stimulus onset to the
249 moment of investment, and the repayment phase as the period during which the partner's return
250 was displayed (Figure 1). Trials from both the cooperative and unfair conditions were contrasted
251 with control trials. Additionally, cooperative trials were contrasted with unfair trials, to directly
252 compare the differences in response to cooperative and unfair feedback.

253 A priori ROI analyses were performed. ROIs were derived from Gromann et al (2014).
254 Talairach coordinates were converted to MNI space (tal2mni under MatLab), resulting in the
255 following ROIs: right caudate (MNI coordinates 16, 17, 7), superior temporal sulcus (STS; 62, -58, 5)
256 and TPJ (51, -57, 26), left insula (-33, 14, 0), and medial prefrontal cortex (mPFC; -3, 65, 25). We
257 tested group differences using MarsBaR (version0.43; <http://marsbar.sourceforge.net>). An adjusted
258 *p*-value was calculated, taking the correlation between the β -values into account by using the
259 Simple Interactive Statistical Analysis Bonferroni tool
260 (<http://www.quantitativeskills.com/sisa/calculations/bonfer.htm>), resulting in adjusted *p*-values
261 (see Table 2) (Li et al, 2014; Woudstra et al, 2013). Additional whole-brain analyses were
262 performed, to investigate activation outside the predefined ROIs.

263 2.5.3 Associations with symptoms

264 Group differences in the association of first investment and development of investments with
265 symptoms (paranoia item, positive and negative PANSS subscales) were investigated. The
266 persecution item (P6) and the depression item (G6) were used as an additional index for paranoid
267 ideation, as previously reported (Gromann et al, 2013), and depression, based on CHR comorbidity.
268 Second, beta weights of the ROIs (average over all voxels) were associated with symptoms. To
269 further explore the association of symptoms with behavior and brain activation, additional post-hoc
270 analyses within patient groups were performed.

271 3 Results

272 3.1 Participant characteristics

273 Participant characteristics and baseline trust are displayed in Table 1. CHR were significantly older
274 than controls ($\beta = .40, p < .001$) and FEP ($\beta = .57, p < .001$), and FEP scored significantly lower on
275 the WAIS vocabulary scale than controls ($\beta = -.39, p = .003$) and CHR ($\beta = -.31, p = .01$). The time
276 between diagnosis and inclusion in the study did not differ significantly between FEP and CHR ($\beta =$
277 $-.35, p = .21$).

278 3.2 Behavioral results

279 Based on the group differences, all analyses were controlled for age and WAIS vocabulary score.
280 Four controls, three FEP and one CHR did not believe they played against a human counterpart.
281 Analyses without these subjects yielded similar results. Below the results of the complete sample
282 are reported.

283 Group differences in baseline trust – the first investment of the first game – were found ($\beta =$
284 $-.27, p = .02$; Table 1), with FEP and CHR showing lower baseline trust than controls (FEP: $\beta = -0.24,$
285 $p = .04$; CHR: $\beta = -0.25, p < .05$). CHR and FEP did not differ significantly from each other ($p = 0.8$).

286 To investigate the development of trust over trials we performed a three-way interaction
287 “trial number-by-group-by-condition” on investment. This interaction was not significant and
288 therefore removed from the model. Significant trial number-by-group, condition-by-group and trial
289 number-by-condition interactions on investment were found ($b = .03, 95\% \text{ CI } [.031, .05], p = .001$; b
290 $= .49, 95\% \text{ CI } [.28, .71], p < .001$; $b = -.20, 95\% \text{ CI } [-.24, -.17], p < .001$, respectively), indicating that
291 the development of trust differed between groups (Figure 2).

292 In the *cooperative condition* there was a significant group-by trial number interaction on
293 investment ($b = .03, 95\% \text{ CI } [.006, .05], p = .01$), with FEP showing significantly stronger increase
294 than controls ($b = .08, 95\% \text{ CI } [-.07, .08], p = .03$). Controls and CHR did not differ significantly from
295 each other ($p = .9$). Analysis by group showed that all groups increased investments significantly

296 (all p 's $<.01$; see Figure 2a). In the *unfair condition*, analysis revealed a significant group-by trial
297 number interaction on investment ($b = .03$, 95% CI [.01, .06], $p = .005$), with FEP showing
298 significantly less decrease than the other groups ($b = -.97$, 95% CI [-1.89, -.06], $p = .04$). All groups
299 decreased investments significantly (all p 's $<.001$; see Figure 2b).

300 All analyses were also conducted with medication type (no medication; atypical anti-
301 psychotics; combination of typical and atypical; other psychotropic medication) as a grouping
302 variable. No differences in baseline trust, nor in adjustment of trust were found between the
303 medication groups, and no interactions with symptoms on trust were found.

304

305 3.3 Imaging results

306 ROI analyses revealed significant group differences in the right TPJ only, showing more
307 activation in CHR compared to controls and FEP during the investment phase in the unfair
308 condition (Table 2). Furthermore, CHR activated the TPJ and mPFC more than controls, when
309 investing in an unfair partner compared to a cooperative partner. Other ROIs showed no group
310 differences. During cooperative investment, cooperative and unfair repayment no significant group
311 differences in ROI activation were found. All ROI analyses were also conducted between medication
312 groups. No significant differences in activation were found.

313 Additional exploratory whole-brain analyses, based on a significance level of $p < .05$ FWE
314 cluster corrected, did not reveal significant group differences. Results with a more lenient threshold
315 are presented in Supplementary Table S1.

316 3.4 Symptoms

317 CHR and FEP did not differ significantly from each other in terms of overall and positive symptoms,
318 and on the depression item. Only on the paranoia item, CHR scored significantly higher than FEP (β
319 = .37, $p = .03$). There was a trend towards significance indicating that FEP had higher negative
320 symptoms than CHR ($p < .07$).

321 3.4.1 Associations between behavioral outcomes and symptoms

322 No group-by-symptoms interactions on first investment were found (all $ps > .24$). After removing
323 the interaction from the model, no main effects of symptoms on baseline trust were found (all $ps >$
324 $.16$).

325 In the *cooperative condition*, the group-by-trial number-by-symptoms models showed a
326 significant interaction for negative symptoms only ($b = -.02$, 95% CI $[-.03, -.001]$, $p = .03$), indicating
327 that negative symptoms impacted upon the development of trust differentially in the three groups.
328 Post-hoc analyses showed a significant association between symptoms and changes in investments
329 over trials in FEP ($b = -.01$, 95% CI $[-.02, -.003]$, $p = .004$), but not in CHR. To visualize this
330 association, we divided the negative symptoms in three levels (Figure 3). Analysis indicated that the
331 only highest level of negative symptoms interfered with increasing investments. No significant
332 interactions with positive symptoms, paranoia, or depression were found.

333 In the *unfair condition* the group-by-trial number-by-symptoms models did not show
334 significant interactions. After removing the 3-way interaction from the model, the interaction of
335 positive symptoms with trial number became significant ($b = .01$, 95% CI $[.0002, .012]$, $p = .04$).
336 Higher positive symptoms were associated with less decrease in investments in FEP and CHR.
337 Associations of decreasing investment with negative symptoms, paranoia, and depression showed
338 no significant group differences. Analyses within each patient group revealed a significant
339 association between depression and investment over trials in FEP ($b = .03$, 95% CI $[.007, .056]$, $p =$
340 $.01$), showing that FEP with a more severe depression score adjusted their investment less to the
341 negative feedback than FEP with milder depression scores.

342 3.4.2 Associations of ROI beta weights and symptoms

343 Beta weights of the ROIs showing group differences (Table 2) were correlated with the positive and
344 negative subscales, and PANSS paranoia and depression score. No group differences were found in
345 the association between symptoms and ROI activation. After removing the group-by-symptoms
346 interaction from the model there was a positive association at trend level for TPJ activation and
347 paranoia in the unfair investment phase ($\beta = .32, p = .07$), indicating that in both patient groups the
348 TPJ was increasingly activated in patients with higher paranoia.

349 Beta weights of the TPJ were unevenly distributed. Therefore, Spearman rank correlation
350 was used for the exploratory analyses per group. This analysis revealed significant associations
351 between symptoms and beta weights of the TPJ for CHR, but not for FEP. Specifically, CHR showed a
352 significant positive association between the paranoia item, positive, and negative symptoms, and
353 TPJ activation during unfair investments ($\rho = .57, p = .04$; $\rho = .70, p = .008$; $\rho = .64, p = .02$,
354 respectively). No associations between symptoms and mPFC activation were found.

355

356 4 Discussion

357 This study investigated the behavioral and neural mechanisms associated with trust and the
358 association with symptoms in a high-risk (CHR) and first episode psychosis (FEP) sample using an
359 interactive trust game. Participants played two trust games, with a cooperative and an unfair pre-
360 programmed partner. Behaviorally, FEP and CHR only differed from controls, and not from each
361 other, showing reduced basic trust, that is initial trust before partner feedback is revealed. No
362 impairments in the development of trust in response to feedback over trials were found in either
363 patient group, compared to controls. Only in FEP associations between trust development and
364 symptoms were found. On the neural level, CHR recruited the TPJ more than the other groups

365 during investment in the unfair condition, suggesting differential processing as compared to
366 healthy controls and FEP.

367 4.1 Behavioral mechanisms of trust

368 Importantly, and in line with previous research both FEP and CHR showed reduced baseline trust
369 toward unknown others (Fett et al, 2015; Fett et al, 2016; Fett et al, 2012; Gromann et al, 2013).

370 Reduced baseline trust has been found in individuals at genetic risk for psychosis, (Fett et al, 2012),
371 but contradicted by another (Gromann et al, 2014). Contrary to previous research (Fett et al, 2016;
372 Fett et al, 2012) baseline trust in FEP and CHR was not associated with symptom severity. The
373 association with positive symptoms found by Fett et al (2012) was at trend level (.09), providing
374 only tentative support. The fact that reduced baseline trust has also been found in individuals at
375 genetic risk for psychosis and now in CHR in combination with the lack of an association with
376 symptoms, tentatively suggests that reduced baseline trust is linked to the risk for psychosis trait,
377 rather than a consequence of the illness (a state marker) that would be associated with (temporal
378 fluctuations of) symptoms.

379 Feedback learning in cooperative and unfair interactions is still intact in CHR and FEP, as
380 opposed to chronic psychosis (Campellone et al, 2016; Fett et al, 2015). The development of trust in
381 response to positive feedback by the game partner showed, as predicted, that FEP and CHR
382 increased their levels of trust significantly. The same pattern was found in response to negative
383 feedback: over game rounds FEP and CHR decreased their trust to the same level as controls. FEP
384 showed steeper increase in positive interactions than controls, possibly because they were more
385 sensitive to the effects of the positive feedback given they initially had lower expectations, as
386 reflected in lower baseline trust. Furthermore, a ceiling effect might result in a less steep increase
387 for controls, with 23% of the control participants investing the maximum of 10 in 75% or more of

388 the trials. The slightly reduced response to negative feedback in FEP, resembling results of Fett et al
389 (2016), might be explained by the differences in first investment (FEP starting significantly lower).

390 4.1.1 Symptoms

391 Symptom severity on average was similar in the two patient groups. FEP showed substantial
392 variability in symptom severity, which reflects the fact that we included both hospitalized patients
393 and ambulant patients who were in a rehabilitation trajectory. CHR experienced more paranoia
394 than FEP, and FEP tended to have more negative symptoms than CHR. These differences might be
395 explained by medication effects: 64% of the FEP were on atypical antipsychotic medication,
396 probably dampening positive symptoms, whereas 47% of the CHR was on other psychotropic
397 medication (see Table 1). First episode patients with highest negative symptoms, as opposed to
398 milder symptoms, showed almost no adjustment of trust in response to positive feedback. Intact
399 feedback learning mechanisms in FEP were associated with milder negative symptom severity. The
400 association between negative symptoms and problems in social functioning and responding to
401 feedback has been well established in psychosis (Addington & Addington, 2005; Campellone et al,
402 2016; Milev et al, 2005; Strauss et al, 2013; Voges & Addington, 2005; Waltz et al, 2011), possibly
403 reflecting a lack of (social) motivation or depression. In the unfair condition positive symptoms
404 were associated with less decrease in trust in both FEP and CHR. This suggests that positive
405 symptoms interfere with learning from negative social feedback, contradicting earlier findings that
406 found no associations of positive symptoms with learning to trust (Fett et al, 2016).

407 4.2 Neural mechanisms of trust

408 On the neural level CHR activated the TPJ significantly more than the other two groups. The TPJ
409 forms part of the mentalizing system (Fletcher et al, 1995; Frith & Frith, 2006; Van Overwalle,
410 2009), and was previously found to be activated in the trust game (King-Casas et al, 2005; Krueger

411 et al, 2007; Saxe & Kanwisher, 2003; van den Bos et al, 2011). In chronic patients reduced TPJ
412 activation was associated with more positive symptoms (Gromann et al, 2013). In our sample, TPJ
413 activation did not differ between FEP and controls, suggesting a decline in TPJ response with longer
414 illness duration. CHR however, showed increased activation in this area compared to FEP and
415 controls during unfair investment. CHR also showed more TPJ and mPFC activation than controls
416 during investments towards the unfair counterpart, as compared to cooperative counterpart. Since
417 both areas form part of the mentalizing system, this could suggest that unfair interactions elicit
418 increased mentalizing in CHR. Gromann et al (2013), in contrast, found the mPFC to be activated
419 more in cooperative interactions in both patients and controls. Increased neural activation in
420 patients at-risk for psychosis during mentalizing and emotion processing areas despite similar
421 behavioral performance was previously found (Brüne et al, 2011; Derntl et al, 2015; Marjoram et al,
422 2006). The elevated TPJ activation in CHR was associated with higher symptoms in all domains,
423 thus associating higher illness severity with greater neural activity. This association was not found
424 in FEP, possibly suggesting different underlying mechanisms between groups. In combination with
425 the behavioral data, showing that CHR adapted adequately to negative social feedback, the
426 increased TPJ activity could indicate a cognitive mechanism by which increased mentalizing helps
427 to respond adequately to negative feedback, indicating more effort, or an inefficient use of the TPJ.
428 The data do not point to compensating mechanisms, since they would suggest deficiencies or
429 reduced processing in other parts of the brain. These were not found in the ROI analyses, nor in the
430 additional whole brain analyses (see supplementary Table S1). The results show no evidence for
431 reduced sensitivity to social reward in FEP and CHR, and suggest that altered mentalizing might be
432 associated with reduced baseline trust. However, due to the small sample size, this result must be
433 interpreted with caution.

434 4.3 Clinical high-risk

435 Following the procedure of previous CHR investigations (Fusar-Poli et al, 2010a; McGorry &
436 van Os, 2013; Phillips et al, 2009; Rietdijk et al, 2012; Shim et al, 2008; Thompson et al, 2012;
437 Valmaggia et al, 2013; van der Gaag et al, 2012; Wood et al, 2011), we included participants
438 assessed with the CAARMS, and with a score below 55 on the SOFAS. Our sample was comparable
439 to other samples in terms of comorbidities (Corcoran et al, 2011; Fusar-Poli et al, 2014; Ising et al,
440 2016; Modinos et al, 2014; Morrison et al, 2012; Woods et al, 2009). One year after testing, e.g.
441 around two years after initial assessment, CHR participants were re-assessed with the CAARMS, to
442 investigate their current status. One of the CHR had made the transition to psychosis. Of two CHR
443 transition data were missing. In this aspect, our high-risk group differed from other high-risk
444 groups. Variant transition rates have been reported in comparable samples with regard to
445 assessment and age range (Broome et al, 2012; Broome et al, 2005; Demjaha et al, 2010; Fusar-Poli
446 et al, 2010b; Nelson et al, 2011). Transition rates in similar referred samples are under 10%
447 (Rietdijk et al, 2012; Yung et al, 2011). Patients already received treatment for their primary
448 problems, including cognitive behavioral therapy (CBT) for their CHR status (psychotic symptoms).
449 This has shown to be an adequate strategy to reduce symptoms, increasing their social functioning
450 skills, to reduce the transition rates (by 46%), and to increase chances for remission (Cannon et al,
451 2008; Ising et al, 2016; Jang et al, 2011; Niendam et al, 2007; van Os & Murray, 2013).

452 In a recent discussion on CHR it has been argued that the presence of psychotic symptoms is
453 possibly more important than transition in the assessment of CHR (van Os & Reininghaus, 2016).
454 Many patients in care for anxiety and depression report psychotic symptoms (van Os & Linscott,
455 2012; van Os & Reininghaus, 2016; Velthorst et al, 2009; Wigman et al, 2012; Woods et al, 2009),
456 but do not transition to psychosis. The current sample fits previous descriptions, making it a
457 representative sample of patients with psychotic symptoms and generally poor mental health. The
458 addition of psychotic symptoms renders these patients at-risk for developing psychopathology,
459 without the direct consequence of developing a psychotic disorder (Fusar-Poli et al, 2013; Yung et

460 al, 2012). In many cases subclinical psychotic experiences are transitory (van Os & Reininghaus,
461 2016). However, the presence of psychotic symptoms is associated with a poorer prognosis,
462 showing that these patients are certainly in need of special care (McGorry & van Os, 2013;
463 Ruhrmann et al, 2010; Valmaggia et al, 2013; van Os & Linscott, 2012; van Os & Reininghaus, 2016).

464 4.4 Limitations and future directions

465 Several limitations should be considered. First, current results should be interpreted with caution,
466 due to the small sample size, especially of the CHR sample. Our CHR results should therefore be
467 considered as a first step in research on real-time social interaction in high-risk individuals, that
468 warrants replication and extension in future research (see also (Broome et al, 2010; Fusar-Poli et
469 al, 2011; Juckel et al, 2012). With a larger CHR sample, the number of converters will increase,
470 allowing for an investigation of converters vs. non-converters. Additionally, a larger sample would
471 provide the possibility to subdivide the CHR sample on the basis of different symptomatology
472 (Fusar-Poli et al, 2014; Valmaggia et al, 2013), yielding more insight in the factors causing social
473 problems, and explaining transition trajectories. Transition in these studies was not explained by
474 comorbid anxiety and depression, but by the severity of CAARMS score, social dysfunction and
475 increased negative symptoms. Direct comparison between at-risk subjects with and without
476 progression into a diagnosis of schizophrenia could also elucidate whether greater activation of the
477 mentalizing network in CHR is serving as a compensatory mechanism, or could also be linked to
478 transition to psychosis. Larger samples could have revealed group differences that were not
479 apparent in this sample. Further, FEP symptom severity was rather mild, possibly due to
480 responsiveness to antipsychotic treatment. Similar symptom severity has been found in stable and
481 medicated patients (Möller et al, 2005). Including a broader range of symptom severity would
482 increase the validity of the sample. Methodologically, a limitation is that participants were not paid
483 on performance. There is some evidence that real payment has a different effect on decisions and

484 related brain activity than hypothetical payment (Hertwig & Ortmann, 2001; Johnson & Mislin,
485 2011; Vlaev, 2012), but other studies have found no differences (Locey et al, 2011; Madden et al,
486 2003). Plausibly, hypothetical payment may influence the strength, but not the direction of the
487 effect (Derks, 2015). Furthermore, eight participants did not believe they were playing against a
488 human counterpart. This might have influenced their behavior. However, analyses without these
489 participants did not change the results. Adequate increase and decrease of investments in response
490 to cooperative and unfair feedback showed that overall the experimental manipulation of the
491 counterpart was effective. Additionally, including an online mentalizing task in the scanner could
492 establish direct links with the current outcomes, pointing in the direction of differential mentalizing
493 processes.

494

495 5 Conclusion

496 Summarizing, baseline trust is impaired in FEP and CHR, indicating that reduced baseline
497 trust is associated with the risk for psychotic illness trait, or with poor mental health in general,
498 rather than a consequence of psychotic disorder. CHR performed in between controls and FEP
499 (Pukrop et al, 2006; Thompson et al, 2012), resembling most the patient group. In contrast to
500 chronic patients (Fett et al, 2015), reward learning was not impaired in FEP and CHR (see also
501 (Juckel et al, 2012). This suggests that with positive social feedback, the lack of initial trust in FEP
502 and CHR can be restored, at least in the context of the trust game. The neural results pointed to
503 globally intact neural mechanisms associated with trust, except for changes in brain areas
504 associated with mentalizing in CHR. However, these findings should be considered as preliminary,
505 and more research in the field is needed to replicate and extend our findings.

506

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772

773

774 Table 1

775 Participant Characteristics and Baseline Trust in the Trust Game

776

	Controls <i>N</i> = 43	CHR <i>N</i> = 17	FEP <i>N</i> = 22	
Gender, <i>N</i> male (%)	22 (51.16 %)	7 (41.18 %)	14 (63.64 %)	777
Age, mean (<i>SD</i>)	21.06 (2.74)	23.78 (2.42) **	19.88 (1.54)	778
WAIS vocabulary, mean (<i>SD</i>)	41.77 (11.39)	41.71 (12.16)	32.5 (10.53) **	779
PANSS total (<i>SD</i>)		58.92 (11.84)	61.32 (14.51)	
mean (<i>SD</i>)		1.95 (.37)	2.09 (.51)	780
- Positive total (<i>SD</i>)		13.38 (2.59)	13.23 (5.72)	781
mean (<i>SD</i>)		1.91 (.39)	1.89 (.82)	
- Negative total (<i>SD</i>)		13.69 (3.73)	17.18 (5.85)	782
mean (<i>SD</i>)		1.96 (.53)	2.45 (.84) * [^]	783
- General total (<i>SD</i>)		31.85 (6.07)	30.91 (7.66)	784
mean (<i>SD</i>)		2.0 (.38)	1.93 (.48)	785
- Paranoia mean (<i>SD</i>)		3.38 (1.50)	2.23 (1.41) *	
- Depression mean (<i>SD</i>)		4 (1.73)	2.89 (1.49)	786
Medicated <i>N</i> (%)		8 (47%) *	16 (73%)	787
- Atypical anti-psychotics (%)			13 (81.5%)	788
- Typical and atypical antipsychotics (%)			1 (6.25%)	789
- Anti-depressant (%)		3 (37.5%)	-	
- SSRI (%)		3 (37.5%)	-	790
- Benzodiazepine (%)		2 (25%)	1 (6.25%)	791
- Sertraline (%)		-	1 (6.25%)	792
Baseline trust, mean (<i>SD</i>)	7.02 (1.81) **	5.82 (2.32)	5.52 (2.02)	793

794 * = significant difference between FEP and CHR at $p < .05$ 795 ** = significantly different from both other groups at $p < .05$ 796 *[^] = FEP > CHR at $p < .07$

797 Note: CHR = clinical high-risk; FEP = first episode psychosis; SD = standard deviation; WAIS =

798 Wechsler Adult Intelligence Scale; PANSS = Positive and Negative Syndrome Scale; SSRI = selective

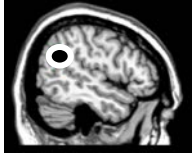

799 serotonin reuptake inhibitors. The paranoia item forms part of the positive subscale (P6), the

Trust in CHR and psychosis

800 depression item forms part of the general subscale (G6). For analyses, these item were investigated
801 separately.
802

803 Table 2

804 Region of Interest (ROI) Analyses Outcome

Condition	ROI	<i>p</i>	Contrast	Location
Unfair Investment *	TPJ	.019	CHR > FEP	
	TPJ	<.001	CHR > Controls	
Unfair investment > cooperative investment **	TPJ	.005	CHR > Controls	
	mPFC	.007	CHR > Controls	

805 *Note:* * = adjusted significance level for multiple comparisons of *p* = .027.

806 ** = adjusted significance level of *p* = .012.

807 Montreal Neurological Institute (MNI) coordinates: TPJ = temporo-parietal junction, 51, -57, 26;

808 mPFC = medial prefrontal cortex, -3, 65, 25. CHR = clinical high-risk; FEP = first episode psychosis

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