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- 13
- 14 Abstract

15 Social norms are a cornerstone of human society. When social norms are violated (e.g. fairness) 16 people can either help the victim or punish the violator in order to restore justice. Recent research has shown that empathic concern influences this decision to help or punish. Using functional magnetic 17 resonance imaging (fMRI) we investigated the neural underpinnings of third-party help and 18 19 punishment and the involvement of empathic concern. Participants saw a person violating a social 20 norm, i.e. proposing unfair offers in a dictator game, at the expense of another person. The participants could then decide to either punish the violator or help the victim. Our results revealed 21 that both third-party helping as well as third-party punishing activated the bilateral striatum, a region 22 23 strongly related with reward processing, indicating that both altruistic decisions share a common neuronal basis. In addition, also different networks were involved in the two processes compared 24 25 with control conditions; bilateral striatum and the right lateral prefrontal cortex (IPFC) during helping 26 and bilateral striatum as well as left IPFC and ventral medial prefrontal cortex (vmPFC) during 27 punishment. Further we found that individual differences in empathic concern influenced whether people prefer to help or to punish. People with high empathic concern helped more frequently, were 28 faster in their decision and showed higher activation in frontoparietal regions during helping 29

30 compared with punishing. Our findings provide insights into the neuronal basis of human altruistic

31 behavior and social norm enforcement mechanism.

32 1. Introduction

33 Humans have an intriguingly complex social norm system, which is unique in the animal kingdom and essential for the functioning of human society (Fehr & Rockenbach, 2004). Self-interests are 34 often in conflict with these social norms. When allocating resources our self-interests might lead us 35 to favor an unequal distribution at the expense of others, violating fairness or equal distribution 36 norms. When observing another person violating a social norm, e.g. treating another person unfair, 37 38 we have at least two options of how to react to this norm violation, namely to either punish the offender, or to help (compensate) the victim. Punishing the offender is referred to as retributive 39 justice (Hogan & Emler, 1981) and helping the victim is referred to as compensatory justice (Darley 40 41 & Pittman, 2003). Usually people have to choose whom they want to focus on (i.e. the offender or the victim) and then decide whether they want the offender to pay for what he or she did, or whether 42 43 they want to restore the harm done to the victim (Schroeder, Steel, Woodell, & Bembenek, 2003). It was shown that people's first reaction to norm violations of high severity is to punish the offender. 44 However, people have a desire to help the victim after norm violations of low severity or when asked 45 to focus on the victim (Gromet & Darley, 2009). Furthermore, victims themselves attach importance 46 to being helped or compensated (Umbreit, 1998). Thus, both punishing the offender as well as 47 48 helping the victim are conceivable reactions to norm violations and might help to restore social 49 equity.

Helping a victim as well as punishing a norm violator as a third person (outside observer) can be 50 51 regarded as altruistic acts. Both cost people at least time and effort but provide no direct benefits. Nevertheless people show altruistic helping (Leliveld, Van Dijk, & Van Beest, 2012) as well as 52 altruistic punishment (Fehr & Fischbacher, 2004; Fehr & Gächter, 2002). Both behaviors reduce 53 54 inequality between offender and victim. Recent neuroimaging studies suggest that altruistic behavior is intrinsically rewarding as it was found to be correlated with activity in the striatum, an area known 55 56 to be involved in reward processing (Haber & Knutson, 2010). Specifically, the ventral striatum was 57 shown to be activated when people invest their own money to reduce their teammates' physical pain (Hein, Silani, Preuschoff, Batson, & Singer, 2010) and when helping an African orphan (Genevsky, 58 Västfjäll, Slovic, & Knutson, 2013). Although the first party is not explicitly mentioned in these 59 studies, helping in this context can be regarded as a form of third party helping. Participants were not 60 involved in the unfair situation themselves (they were neither victims nor violators) but helped 61 another victim. De Quervain and colleagues found that the striatum was also involved in second party 62 punishment, namely when participants punished the untrustworthy opponent in a trust game 63 64 paradigm (de Quervain et al., 2004). In this case the participant was the victim of unfair behavior. So

far, there are only two studies on the neural correlates of third-party punishment (Buckholtz et al., 65 2008; Strobel et al., 2011). In the study by Buckholtz and colleagues, participants were asked to rate 66 the appropriate punishment for crimes they were not involved in. However, decisions in this study 67 68 were not incentivized and therefore not costly for participants. In another study by Strobel and colleagues, a modified paradigm of dictator game was adopted in which participants played the role 69 of either the recipient (i.e. second-party) or the observer (i.e. third-party) and they could punish the 70 71 dictator with their own money. They found that both second-/third-party punishment (vs. no punishment) elicited stronger activation in ventral striatum. Thus, up to now neuroimaging studies 72 73 show that second-party punishment and third party help involve similar neuronal processes, namely activity in reward areas. Reward might be an underlying mechanism for both third party help and 74 punishment decisions, thus both might involve activity in the striatum 75

76 Despite some similarities, behavioral studies suggest that third-party altruistic punishment and help 77 seem to be driven by different motives. On the one hand people feel sympathy/empathy with the victim triggering a desire to restore the person (Gromet & Darley, 2009). On the other hand norm 78 violations induce strong negative affect which lead people to punish the offender (Egas & Riedl, 79 2008; Fehr & G ächter, 2002). One additional motive of punishment is deterrence; punishment has the 80 additional function to prevent offenders from future norm violations (Carlsmith, Darley, & Robinson, 81 82 2002). Taken together, behavioral studies suggest that third party help and punishment are differently motivated and might therefore involve different processes. Intriguingly, people differ in their 83 84 responses when asked to choose between punishing the offender and helping the victim of a norm violation. A recent behavioral study found that when witnessing an unfair case of monetary 85 allocation, people as third parties with low empathic concern preferred punishment, whereas those 86 87 high in empathic concern preferred helping (Leliveld et al., 2012). This indicates that empathic 88 concern plays an important role in influencing people's choice either to help or to punish. Empathic concern is defined as an other-oriented altruistic motivation congruent with the perceived welfare of 89 90 another person; namely a feeling of concern for other people who are in need or suffer from an unfortunate case (Batson et al., 1988; Coke, Batson, & McDavis, 1978). More crucially, previous 91 92 studies have shown empathic concern is a reliable indicator for helping behavior (Batson et al., 1988; Coke, Batson, & McDavis, 1978). As a stable disposition variable, empathic concern was measured 93 by one subscale of the Interpersonal Reactivity Index (IRI_EC, German version; Davis, 1983). The 94 95 IRI_EC was also used in previous neuroimaging studies to investigate correlations between empathic concern and empathic neural responses, however the results are inconclusive. One of the main 96 97 reasons is that different approaches were used in those studies to assess the neural correlates of empathy, which makes it difficult to compare the results (Decety, 2011; Lamm, Nusbaum, Meltzoff, 98 99 & Decety, 2007; Singer et al., 2004). For example, Singer and colleagues (2004) adopted a cue-based paradigm, in which participants' empathy was elicited by abstract visual information about their 100

partner's affective state. They found stronger positive relation between IRI_EC scores and empathic
neural activities in anterior cingulate cortex and left anterior insula. In the study by Lamm *et al.*(2007), a picture-based paradigm was used, in which participants' empathy was elicited by viewing
other's body parts in painful situations (e.g. the painful needle injection on someone's hand).
However, no correlation was found between IRI_EC and empathic neural activities in those regions.

Although third-party help and punishment have been extensively investigated in behavioral studies, 106 the neuronal basis of third-party help and punishment has not been examined simultaneously in one 107 108 study using the same paradigm so far, allowing for a direct comparison. Furthermore, the association 109 between empathic concern and brain responses to third-party help or punishment is still unclear. Adapting the paradigm of Leliveld et al. (2012) we investigated the neural correlates of third-party 110 help and punishment simultaneously in one study by using fMRI. Our aim was to examine the neural 111 processes underlying third-party help and punishment and their relation to individual differences in 112 113 empathic concern. Based on previous neuroimaging research we hypothesize that both third-party help as well as punishing activates the striatum (de Quervain et al., 2004; Genevsky et al., 2013). 114 However, since behavioral studies showed that the motives to punish and help are different we 115 predict that help and punish elicit activity in separate brain regions connected to the striatum. 116 Furthermore, we assume that individual differences in empathic concern correlate with both the 117 118 frequency of help decisions and brain activity related to help (vs. punishment). Given that previous studies do not report any consistent results about possible target regions for the connectivity analyses, 119 120 we refrain to make strong predictions but rather choose to present exploratory results.

121

122 2. Materials and methods

123 2.1. Participants

Thirty-six German participants (12 males; mean age = 22.72 ± 2.85) were tested in the fMRI experiment. All participants reported no history of psychiatric or neurological disorders. They were recruited via the Online Recruitment System for Economic Experiments (ORSEE). Written consent was given by all participants according to the Declaration of Helsinki (BMJ 1991; 302: 1194) and the study was approved by the ethics committee of the University of Bonn. Additional 84 participants (30 males; mean age = 23.58 ± 6.13) were recruited for the behavioral experiment from the same subject pool as used for the fMRI experiment.

131

132 2.2. Stimuli and design

The experiment consisted of two parts: a behavioral and an fMRI part. Participants of the behavioral 133 part were asked to play a Dictator Game. During ten rounds half of them played the role of the 134 "proposer" (i.e. first-party) and the other half the role of the "recipient" (i.e. second-party). We used a 135 136 perfect stranger matching to allocate participants for each round. The "proposer" received an endowment of 100 monetary units (MUs; 1MU=20 Cents) per round and could decide how to 137 distribute these between him-/herself and the recipient (i.e. 0, 10, 20, 30, 40, 50). Participants were 138 informed that some of their decisions were forwarded to a third party (i.e. the fMRI participants). In 139 case of an unfair allocation the third party could decide to either help the recipient by transferring 140 141 MUs to increase his/her original MUs or to punish the proposer by investing own MUs to subtract his/her original MUs. Participants were further asked to indicate their initials and were informed that 142 these were forwarded to the third parties. All participants of the behavioural experiment received a 4 143 € show-up fee at the end of the experiment. They were also informed that in addition all parties 144 would receive payoffs depending on one randomly chosen round of the experiment. Thus, if the third 145 146 party decided to either help the recipient or punish the dictator this decision was implemented accordingly. The additional payoffs (M = $10.05 \in$, SD = $7.26 \in$) for participants of the behavioural 147 experiment were paid four weeks later. The behavioural part of experiment was conducted in Bonn 148 EconLab via Z-tree (Fischbacher, 2007). 149

In total, 420 decisions were made by the proposers, including 63 decisions of 50/50 offer, 43 offers of 60/40 offer, 33 decisions of 70/30 offer, 57 decisions of 80/20 offer, 82 decisions of 90/10 offer and 142 offers for 100/0. Given the goal of our study and the fMRI design, we focused on the unfair offers (i.e. 60/40, 70/30, 80/20, 90/10, 100/0) and selected 160 offers to present those in the fMRI study. Among them, 120 offers were presented in the decision condition and 40 in the control condition. Each offer (i.e. 60/40, 70/30, 80/20, 90/10, 100/0) occurred 24 times in the decision condition and 4 times in the control condition.

157

158 2.3. fMRI procedure

Participants were informed about the behavioral experiment and that they would see a set of allocations made during this experiment. They were further told that they could influence the payoff of either the first or second party by investing their own endowment. Importantly, both options were costly for the participant, meaning that they had to invest one MU in order to either subtract three from the proposer or to increase three to the recipient. Prior to the scanning session, participants received an instruction which included a short comprehension test to further make sure that they understood the task.

166 The scanning session consisted of two fMRI runs, which were separated by a self-paced break. In

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each run, there were 80 trials; 60 decision trials (12 trials per offer) and 20 control trials (4 trials per 167 offer, half of them were in help/punish condition). In each trial, participants were endowed with 50 168 MU (1MU=20 Cents). In the decision condition, participants first saw the unfair monetary allocation 169 170 paired with the initials of the first and second party (Figure 1A). On the same screen they were asked whether they wanted to increase the recipient's payoff or to decrease the proposer's payoff. Once 171 they made a choice, a cue appeared under the corresponding option (the decision phase). Independent 172 of their response time the decision phase was presented for 4s. The decision phase was followed by 173 an inter-stimulus fixation point (1-3s). On the next screen participants could decide how much they 174 175 want to increase or decrease the payoffs of the other players (the transfer phase; 4s), followed by an inter-trial fixation point (3-7s). Participants could respond by pressing the button of response grips 176 with left/right index fingers in both phases of the task. In the control condition, the procedure was 177 identical except that in both phases decisions were made by the computer instead of the participants 178 lying in the scanner. The offers presented during these trials were still made by the participants of the 179 180 behavioral experiment though. Thus, participants in the scanner did not make any decisions themselves, however, these trials were relevant for the payoffs of all parties (proposer, recipient and 181 fMRI participant). Participants therefore had an incentive to keep track of the control condition trials. 182 No button presses were asked of the participants in the control condition to limit the feeling of a 183 forced choice which might lead to conflict, anger or frustration. These trials were indicated by a 184 185 white frame (Figure 1B). The display of the task and response collection was performed with Presentation 14.9 (Neurobehavioral System, Albana, Canada). Participants saw the experiment via 186 video goggles (Nordic NeuroLab, Bergen, Norway) and their responses were recorded by response 187 188 grips (Nordic NeuroLab, Bergen, Norway).

189

(Insert Figure 1 about here)

It is important to highlight the following details of the paradigm and the procedure. First, the 190 words "help" and "punish" were not used in the instructions ("increase" and "subtract" were adopted 191 192 instead) to avoid demand characteristics. Second, consistent with previous literature (Fehr & Fischbacher, 2004; Leliveld et al., 2012), the cost ratio was set to 1:3, which means that 1 MU 193 transferred from participants could either subtract 3 MU from the first party or increase 3 MU to the 194 second party. Third, in the transfer phase participants could decide to invest 0 MU. Thus every 195 decision to invest MU to either increase or decrease MUs of the others can be regarded as their 196 voluntary decision. Fourth, the position of two options (i.e. "increase" and "subtract") in the decision 197 phase were counterbalanced across trials. The default position of the amount participants could invest 198 in the transfer phase was randomly determined from 0 to 50. Finally, the first party could not lose 199 money (i.e. the minimum payoff was 0). 200

201

After scanning, participants were asked to fill in the Interpersonal Reactivity Index (IRI) scale,

used for measuring trait empathy and to make judgments about the fairness of the six different offers

203 (i.e. the offer 50/50 was also included) on a 8-point Likert scale (1=very fair, 8=very unfair). Finally,
 204 participants received a 10 € show-up fee and one randomly selected trial was paid to all three parties

205 accordingly (M = 7.0 \in , SD = 2.5 \in).

206

207 2.4. Data collection and analyses

208 The imaging data was collected via the 3-Tesla Siemens Trio platform at the Imaging Center of Life & Brain, University Hospital Bonn. For functional images, 37 axial slices (FOV = 192×192 mm², 209 matrix = 96×96, in-plane resolution = $2 \times 2 \text{ mm}^2$, thickness = 3 mm) covering the whole brain were 210 211 obtained using a T2*-weighted echo planar imaging (EPI) sequences with blood-oxygenation-level dependent (BOLD) contrast (TR = 2500 ms, TE = 30 ms, flip angle = 90 %. A high-resolution 212 structural image for each participant was acquired using 3D MRI sequences for anatomical co-213 registration and normalization (TR = 1660 ms, TE = 2.75 ms, flip angle = 9°, matrix = 320×320 , 214 $FOV = 256 \times 256 \text{ mm}^2$, slice thickness=0.8 mm). 215

Eleven participants were excluded due to the following reason: 10 of them had insufficient number 216 of trials (less than 5 trials) for one or both decision regressors (help decision: n=1; punish decision: 217 n=7; both decisions: n = 2) and one participant terminated the experiment because he or she felt 218 uncomfortable in the scanner. For the remaining 25 participants, SPM8 was used for the fMRI data 219 220 analysis (Wellcome Trust Department of Cognitive Neurology, London, UK). For each run of each participant, the first three volumes were discarded to allow the stabilization of BOLD signal. The 221 222 following preprocessing steps were applied: EPI images were first realigned to the first volume to 223 correct for head motions (< 2.5 mm) and corrected for slice timing. Then, the anatomical image was co-registered to the mean EPI image, and segmented, generating parameters for normalization to 224 MNI space. Using these parameters, all EPI data were projected onto MNI space with a $2 \times 2 \times 2$ mm³ 225 226 resolution and smoothed using an 8-mm FWHM (full width half maximum) isotropic Gaussian kernel. High-pass temporal filtering with a cut-off of 128 s was performed to remove low-frequency 227 228 drifts.

For the individual-level analyses, a general linear model (GLM) focusing on the decision-phase with five onset regressors (i.e., "help", "punish", "help_control", "punish_control", "other") convolved with the canonical hemodynamic response function (HRF) was applied. The "other" regressor included the following onsets: onsets of transfer phase and onsets of no response as well as trials in which participants transferred 0 MU in decision phase. For runs in which either "help" or "punish" condition was less than 5 trials, onsets of that condition in decision phase were also categorized into "other" condition. The six estimated head movement parameters were included in

the design matrix to account for the residual effects of head motion. For the group-level analyses, a one-sample t-test as well as a flexible factorial model was performed to test the difference and the conjunction of the activation elicited by "help" and "punish" option. Parameter estimates (contrast values) and percent signal change of the peak voxel was extracted via MarsBar (http://marsbar.sourceforge.net).

241

242 2.4.1. Correlation analysis

To investigate how trait empathy correlates with third-party decisions at the neural level, a correlation analysis was applied to compute the relationship between the individual neural contrast of "help" vs. "punish" and individual scores of empathic concern subscale of IRI (IRI_EC).

246

247 2.4.2. Psycho-physiological interaction (PPI) analysis

In order to test whether different networks are involved during helping and punishing respectively, 248 249 we performed a PPI analysis (Friston et al., 1997; Gitelman, Penny, Ashburner, & Friston, 2003). Specifically, the source masks were defined as two 8-mm spheres centered at the peak voxel of the 250 group-level conjunction results of the two contrasts "help" vs. "help control" and "punish" vs. 251 "punish control" within bilateral striatum based on AAL templates with the wfu pickatlas tool. The 252 seed volume of interest (VOI) for each individual was then defined as a sphere with a 6-mm-radius 253 254 centered at the peak voxel from the contrast of either "help" vs. "help control" or "punish" vs. "punish control" within these source masks. The time series of each VOI was extracted and then 255 deconvolved, multiplied with the psychological variable ("help" > "help control" or "punish" > 256 "punish control") and reconvolved with a hemodynamic response function to set up the PPI 257 258 regressor, which followed the procedure by Gitelman et al. (2003). These three regressors (i.e. the 259 PPI regressor, the VOI time-series, the psychological variable) were convolved with the canonical HRF and then entered into the regression model along with six head motion parameters. The 260 individual parameter estimates image for the PPI regressor was subsequently subjected to one-sample 261 t-tests. Finally, a group analysis was performed to identify the brain regions displaying increased 262 functional connectivity with the seed VOI during either help or punishment decisions. Besides, two 263 264 paired-samples t-tests were performed to further test the different connectivity patterns between help 265 and punishment decisions with either left or right striatum.

For all whole-brain based analyses mentioned above, the threshold of p < 0.001 uncorrected at peak voxel level with the extent threshold at k = 50 was adopted.

268

269 3. Results

270 **3.1. Behavioral results**

271 Data from 25 participants were used for behavioral analyses. A paired-samples T-test was performed 272 between help and punishment decisions in the decision condition on the behavioral factors decision rate (i.e. the ratio of help/punish decision compared in relation to all respective trials), response time 273 (ms) and transfer amount (MU). The transfer amount was significantly different between help and 274 275 punishment trials. Participants transferred more MUs when they punished the first party (M = 16.15, SD = 6.86) than when they helped the second party (M = 11.07, SD = 5.07) [95% C.I. of the 276 277 difference: -8.28 to -1.89; $t_{(24)}$ =3.266, p = 0.003, Cohen'd = -0.664]. No significant differences were detected in the rate of decisions to help or punish (help: M = 49.30%, SD = 27.28%; punish: M =278 279 42.40%, SD = 27.90%) [95% C.I. of the difference: -15.62% to 29.42%; $t_{(24)}$ =0.632, p = 0.533, Cohen'd = 0.126] and response times (help: M = 1583.15, SD = 431.63; punish: M = 1611.45, SD = 280 402.22) [95% C.I. of the difference: -207.55 to 150.93; $t_{(24)}$ =-0.326, p = 0.747, Cohen'd = -0.065] 281 282 between help and punishment trials.

283 To test whether individual differences in trait empathy correlate with the decisions to help or punish, a Pearson correlation was conducted between empathic concern subscale scores of IRI (i.e. 284 IRI_EC) and decision rate in help and punishment decisions respectively. A significant positive 285 relationship was found between IRI EC scores and *help rate* [95% C.I.: 0.06 to 0.71; r = 0.441, p =286 0.027, Fisher's $Z_r = 0.474$], whereas a negative relationship was detected between IRI_EC scores and 287 288 *punishment rate* [95% C.I.: -0.72 to -0.08; r = -0.461, p = 0.02, Fisher's $Z_r = -0.497$; Figure 2A]. To further investigate whether empathic concern has an influence on decision speed in both help and 289 punishment trials, we correlated IRI_EC and the difference in reaction times between help and 290 punishment trials (i.e. RT_help-punish), finding a negative relationship [95% C.I.: -0.69 to -0.01; r = 291 -0.406, p = 0.044, Fisher's $Z_r = -0.431$; Figure 2B]. 292

293

(Insert **Figure 2** about here)

A one-way repeated measure ANOVA on the perceived unfairness rating of the offers showed a main effect of inequity level [95% C.I.: 5.09 to 5.51; $F_{(5, 120)} = 225.967$, p < 0.001, partial $\eta^2 = 0.904$]. Post-hoc analyses revealed that ratings increased with the level of inequity of the offers (50/50: M = 1.48, SD = 1.12; 60/40: M = 3.52, SD = 1.30; 70/30: M = 5.24, SD = 0.93; 80/20: M = 6.24, SD = 0.93; 90/10: M = 7.32, SD = 0.48; 100/0: M = 8.00, SD = 0.00; *ps* < 0.05, *Bonferroni* corrected).

299

300 **3.2.** Imaging findings

301 **3.2.1.** Neural correlates of third-party help and punishment

Both contrast help vs. help_control and punish vs. punish_control showed significant activation in 302 303 several regions, including bilateral striatum, supplementary motor area/mid-cingulate cortex (BA 4/6), inferior/superior parietal lobule (BA 39/40) as well as visual areas (BA 17/18/19) (Table S1 and 304 305 Figure 3). The conjunction analyses further confirmed that the bilateral striatum along with other areas mentioned above were activated by both contrasts, indicating that help- and punish-related 306 cognitive processes shared some common neural bases (Table S1 and Figure 3). Activity in the 307 bilateral striatum remained significant when controlling for motor responses due to button pressing 308 (Table S2 and Figure S1). To test the differential neural correlates between these two processes, help 309 and punishment decisions were directly contrasted, which yielded no significant difference in both 310 directions. These results remained unchanged when controlling for fairness levels and transfer 311 amounts (Table S3 and S4). 312

(Insert Figure 3 about h	ere)
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313 314

315 3.2.2. Relationship between empathic concern and brain activation during third-party 316 decisions

To determine regions in which a change of the BOLD signal to third-party decisions varied with individual difference in trait empathy, a correlation analysis was performed between the contrast of help vs. punishment and IRI_EC scores. Stronger positive correlations were detected in frontoparietal regions including left lateral prefrontal cortex (IPFC, BA 9) and left angular gyrus/inferior parietal lobule (IPL/AG, BA 7/40; **Table S5** and **Figure 4**). No negative correlations were found under the same threshold.

323

(Insert **Figure 4** about here)

324

325 **3.2.3.** Functional connectivity pattern of third-party decisions

In order to investigate whether different networks are involved in third-party help and punishment a PPI analysis was conducted. Based on our hypotheses and the results of the conjunction analyses the striatum was used as the seed region (i.e. left and right striatum). PPI analyses were conducted during help and punishment decisions, respectively (both compared with their respective control conditions). Right IPFC (BA 45/46) showed increased functional connectivity with bilateral striatum during help

decisions (**Table S6** and **Figure 5**), whereas left IPFC (BA 44/45) showed enhanced functional connectivity with both seed regions during punishment decisions (**Table S6** and **Figure 6**). Furthermore ventral medial prefrontal cortex (vmPFC; BA 10/11/32) was observed to show increased connectivity only with right striatum when participants chose to punish (**Table S6** and **Figure 6**). No significant difference in functional connectivity was found in a direct comparison of help and punishment decisions with either left or right striatum.

- 337 (Insert Figure 5 about here)
 338 (Insert Figure 6 about here)
- 339

340 4. Discussion

Our results reveal that both third-party help and third-party punishment share a common neuronal 341 basis, but that specific networks are additionally involved in the two processes. The bilateral striatum 342 was activated by both helping and punishing; functional connectivity between the bilateral striatum 343 344 and the right lateral prefrontal cortex (IPFC) was increased during help and with left IPFC and ventromedial prefrontal cortex (vmPFC) during punishment. Individual differences in empathic 345 concern correlated with people's preference to help or to punish. People with high empathic concern 346 helped more frequently, were faster in their decision and showed higher activation in fronto-parietal 347 regions during decisions to help. 348

The conjunction analysis indicated that third-party help and third-party punishment both share 349 350 some common neural bases. In line with previous findings the striatum showed increased activation during altruistic help (Genevsky et al., 2013; Harbaugh, Mayr, & Burghart, 2007; Hein et al., 2010) 351 352 as well as during altruistic punishment (de Quervain et al., 2004). Helping friends or even strangers and punishing norm violators has been associated with activity in the striatum. However, so far 353 striatal activation was only observed in third party helping and second-party punishment paradigms, 354 for example, while an investor chose to punish an untrustworthy trustee (de Quervain et al., 2004). 355 This is to our knowledge the first study investigating third party helping and punishing in the same 356 study and showing that both are associated with striatal activation. The striatum is part of the human 357 reward system, known to be activated by recognizing and evaluating rewards and learning from them 358 359 (Bhanji & Delgado, 2014). Our results are in line with literature on charitable donation and second 360 party punishment suggesting that both helping an unknown person and punishing an offender is intrinsically rewarding (Fehr & Camerer, 2007; Harbaugh et al., 2007). Here we show that punishing 361 362 an offender as a third person seems to rewarding as well.

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However, an alternative interpretation of this result cannot be ruled out completely. Participants 363 were not required to response during the computer (control) trials in order to avoid additional 364 cognitive (e.g. conflict) or affective (e.g. anger, frustration) processes. Unfortunately this paradigm 365 366 introduced a potential motor confound for the contrasts between help or punish decisions (button presses) and their corresponding control trials (no button presses). Besides its role in reward 367 processing or representation of affective value, the striatum is also frequently associated with motor-368 related functions (Witt, Laird, & Meyerand, 2008; Filevich, Kühn, & Haggard, 2012; Guitart-Masip 369 et al., 2012; Guitart-Masip et al., 2014). In a recent study on the role of the striatum in decision 370 371 making, Guitart-Masip and colleagues independently manipulated both the variables of action (i.e. "go" or "no go") and valence (i.e. "to win" or "to avoid losing") in an instrumental learning paradigm. 372 373 They found that activity in the striatum reflected primarily the action requirements, independent of the valence of decision (Guitart-Masip et al., 2012). This result suggests an involvement of the 374 striatum in motivated action during decision making. In order to control for this, we performed an 375 376 additional analysis in which the onset of the button presses were added in to the GLM as an 377 independent regressor. This analysis showed that the bilateral striatum was still strongly activated during both third-party help and punishment even after controlling for the effect of button pressing 378 (see Table S2 and Figure S1), indicating that activity in the striatum detected in the contrasts of 379 third-party altruistic decisions and control trials is not likely driven by pure motor effects only. 380 381 Rather it more likely reflects processes related to decision making, like rewarding processes as suggested by previous findings on altruistic decisions (e.g. charity donation, second-/third-party 382 punishment; de Quervain et al., 2004; Harbaugh et al., 2007; Hein et al., 2010; Strobel et al., 2011; 383 384 Genevsky et al., 2013) and on reward processing (Haber & Knutson, 2010). However, since the onset of button pressing is not a random event as it is collinear to the onset of decision trials, the analysis 385 unfortunately might not completely tease apart the effect of button pressing and that of decision 386 387 processes. Since we cannot perfectly disentangle brain activity due to decision processes and due to 388 motor processes (button press), the joint activation in striatum during third-party help and punishment decisions should be cautiously interpreted as reward-relevant processing. 389

390 Furthermore, our functional connectivity results suggest that besides the common neural basis, 391 different networks are involved in third-party help and third-party punishment. Increased functional 392 connectivity was found between the bilateral striatum and right IPFC during help decisions whereas left IPFC and the bilateral striatum showed increased functional connectivity during punishment 393 decisions. Furthermore, vmPFC showed increased connectivity with right striatum when participants 394 chose to punish. Generally, our PPI findings are consistent with the anatomical connectivity of the 395 striatum, which was found to be connected with both lateral and ventral/medial parts of the prefrontal 396 397 cortex (Haber & Knutson, 2010). Specifically, IPFC is known to be engaged in cognitive/executive control and goal-directed decisions (Miller & Cohen, 2001; Tanji & Hoshi, 2008). In the social-398

economic domain, especially the right IPFC was shown to be involved in the control of selfish 399 impulses (Knoch, Pascual-Leone, Meyer, Treyer, & Fehr, 2006; Ruff, Ugazio, & Fehr, 2013; Strang 400 et al., 2014). For example, disrupting the right IPFC via lower-frequency repetitive TMS caused 401 402 people to make riskier decisions (Knoch & Fehr, 2007) and to exhibit more norm violating behaviours (Strang et al., 2014). A recent TMS study found that people show more impulsive 403 behavior in an inter-temporal choice task while the left IPFC was inhibited (Figner et al., 2010). 404 Intriguingly, left IPFC showed stronger activity when participants chose to costly punish dictators as 405 a third-party compared to as a second-party, indicating that cognitive-control processes, instead of 406 407 revenge-driven motives, are involved in third-party punishment (Strobel et al., 2011). Consistently, a behavioral study showed that punishment of free-riders by cooperators is linked to self-control 408 409 abilities (Esp n, Brañas-Garza, Herrmann, & Gamella, 2012). Moreover, some studies showed an increased functional connectivity between IPFC and striatum while people controlled reward-related 410 responses to food cues (Hare, Malmaud, & Rangel, 2011) or monetary reward (Delgado, Gillis, & 411 412 Phelps, 2008). Since help and punishment are costly in our paradigm, both require control of selfish 413 impulses in order to engage in one of the two behaviors. Hence, it is possible that in our paradigm increased connectivity between IPFC and bilateral striatum during help and punishment decisions is 414 due to these control processes. Another region found to have increased connectivity with the striatum 415 is the vmPFC, which has been shown to be involved in a variety of cognitive and affective processes 416 417 including integrating emotional information (Naqvi, Shiv, & Bechara, 2006) and subjective valuation during decision making (Ruff & Fehr, 2014). Interestingly, increased activity in the vmPFC was also 418 found when choosing to costly punish an untrustworthy trustee (de Quervain et al., 2004). Our 419 findings seem to support the view of a potentially stronger involvement of the vmPFC in third-party 420 punishment rather than help. 421

422 Nevertheless, two issues limit our interpretation on the PPI results. Firstly, it is important to 423 mention that there was no significant difference between the functional connectivity during help and punishment decisions when directly contrasting the connectivity results in both decisions, which 424 weakens our inference about differential neural networks involved in each decision. This might be 425 426 due to insufficient sample size or inadequate numbers of trials in each condition. Secondly, as both 427 PPI analyses are based on the corresponding control trials (help vs. help_control or punish vs. 428 punish_control), the results are also influenced by the motor confound mentioned above. Thus, the connectivity pattern might also reflect a motor effect during both decisions compared with the pure 429 observation in the control trials. Since the PPI analyses are rather explorative, further research is 430 needed to shed more light on the network involved in third party help and punishment. 431

432 Moreover, our results demonstrate that individual differences in empathic concern influence our 433 decision to help or to punish on a behavioral as well as on a neural level. People with high levels of

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empathic concern chose to help more frequently, were faster in their decision to help and showed 434 higher activation in frontoparietal regions (i.e. left IPFC and left IPL/AG) during this decision. The 435 behavioral findings are in line with previous research (Leliveld et al., 2012), in which the authors 436 437 also reported that people with high empathic concern prefer to help instead of punishing, whereas people with low empathic concern prefer to punish instead of helping. In addition our results 438 demonstrate that high empathic people are also faster in deciding to help compared to deciding to 439 punish, whereas people with low empathic concern show the reversed pattern; they are faster in 440 deciding to punish instead of helping. Faster reaction times are often interpreted as a sign of less 441 442 conflict between the options someone has to choose from and less cognitive processing (Rand, Greene, & Nowak, 2012). According to this literature the results suggest that for high empathic 443 444 people deciding to help needs less cognitive processing. For them the decision to either help or to punish does not involve a conflict, help is the default option for them. Low empathic people also do 445 not encounter a conflict when deciding between help and punishment, their default option is to 446 punish. Whether someone helps a victim or punishes the offender hence depends on how much 447 448 empathic concern someone has. Both regions correlating with empathic concern, IPFC and IPL/AG, are considered as the core components of the frontoparietal network (FPN), which play an important 449 role in top-down cognitive control and attention (Corbetta & Shulman, 2002; Dosenbach, Fair, Cohen, 450 Schlaggar, & Petersen, 2008). Gromet and Darley (2009) argue that punishment might be the default 451 452 choice after observing injustice until people are asked to focus on the victim. Without explicit requirements to focus on the victim, such an attention shift might be influenced by individual's 453 personality trait, in this case empathic concern. Our results hint towards such an empathy-based 454 455 attention shift. However, this interpretation is inconsistent with the reaction time findings, which suggests that help is the default for people with high empathic concern. Thus the role of FPN in 456 457 mediating the relationship between empathic concerns and the two altruistic decisions still needs 458 further investigation. Future studies might shed more light on this question by adopting other 459 techniques (such as eye-tracking) to investigate the difference in fine-grained information search patterns between high and low empathic people during deciding to either help or punish. 460

There are several limitations of this study. One constraint is the difference in motor demands 461 462 between the decision and control trials as mentioned above. Future studies should try to find a clearer 463 way to disentangle activity due to the decision process and motor responses. Another limitation is the high number of excluded participants. We were only able to use data from 25 out of 36 participants, 464 because ten participants did not show enough variability in their behavior to define all necessary 465 regressors. Since trials were sorted into different conditions according to participant's behavior in the 466 corresponding trial, sufficient numbers of trials (>25) for one condition in order to calculate a 467 contrast cannot be guaranteed. Although 25 participants is still a widely accepted sample size in the 468 469 field of cognitive neuroimaging, statistical power might explain the non-significant difference

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especially for the PPI results. Since people who exhibit either very high or very low empathic concerns have a preference for either helping or punishing, respectively, they show less variability in their decisions on the individual level. One possibility to minimize dropout rates is to increase the variability in decision behavior by only inviting participants with empathic concern score in the medium range and thereby increasing statistical power. Additionally a pre-screening could be used to exclude participants who are very selfish and are not willing to help or punish at all.

476 Taken together, by using a modified third-party decision paradigm with fMRI, our study provides first evidence for the neural basis of third-party help and punishment decisions. Both altruistic 477 decisions activated bilateral striatum, indicating that intrinsic reward processes are involved in both 478 479 third-party help and punishment decisions. Differential functional connectivity networks during 480 third-party help and punishment suggest different cognitive processes underlying both decisions. Moreover, the present study replicated previous behavioral findings on the role of empathic concern 481 in mediating people's decisions to either help or punish. Further its underlying neural correlates in 482 frontoparietal regions were detected. Despite some limitations, these findings may provide insights 483 484 for better understanding the mechanism underlying altruism and social norm enforcement.

485

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¹ Provide the doi when available, and ALL complete author names.

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611 7. Figure legends

- 612 Figure 1 Illustration of trial procedures (A) in the decision condition and (B) in the control
- 613 **condition.** ISI = inter-stimulus interval; ITI = inter-trial interval.
- 614 Figure 2 Correlation (A) between IRI_EC scores (X-axis) and average help/punish rate (Y-axis)
- and (B) between IRI_EC scores (X-axis) and the difference in RT between help and punish (Y-
- axis). IRI_EC = empathic concern subscale of interpersonal reactivity index scale; RT = response
- 617 time.
- Figure 3 Separate and conjunction mapping of regions involved in third-party help and
- 619 punishment (A) and timecourse of percent signal change in the local peak voxel of left striatum
- 620 **(B) and right striatum (C) in four conditions** (i.e. help, help_control, punish, punish_control).
- 621 Error bars: SEM.

622 Figure 4 Correlation between the contrast of help vs. punish and IRI_EC scores (A) and plots

of the positive correlation between IRI_EC scores and contrast values in local peak voxel of left

624 **IPFC (B) and that of left IPL/AG (C).** IRI_EC = empathic concern subscale of interpersonal

- reactivity index scale; IPFC = lateral prefrontal cortex; IPL = inferior parietal lobule; AG = angular
 gyrus.
- **Figure 5 Regions showing increased functional connectivity with bilateral striatum during**
- 628 third-party help decisions (compared with control conditions; A) and plots of parameter
- 629 estimates of PPI in the local peak voxel of right IPFC with left (B) /right (C) striatum in four
- 630 **conditions** (i.e. help, help_control, punish, punish_control). Abbreviations: PPI = psycho-
- 631 physiological interaction; IPFC = lateral prefrontal cortex; Stri = striatum; Error bars: SEM.
- 632 Figure 6 Regions showing increased functional connectivity with bilateral striatum during
- 633 third-party punishment decisions (compared with control conditions; A) and plots of
- 634 parameter estimates of PPI in local peak voxel of left lPFC with left (B) /right (C) striatum and
- 635 that of vmPFC with right striatum (D) in four conditions (i.e. help, help_control, punish,
- 636 punish_control). PPI = psycho-physiological interaction; lPFC = lateral prefrontal cortex; vmPFC =
- 637 ventral medial prefrontal cortex; Stri = striatum; Error bars: SEM.
- 638 8. Supplementary Material
- 639 The Supplementary Material for this article can be found in a separate document.

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