

1  
2  
3  
4  
5  
6  
7  
8  
9  
10  
11  
12  
13  
14  
15  
16  
17  
18  
19  
20  
21  
22  
23  
24  
25  
26  
27  
28  
29  
30  
31  
32  
33  
34  
35  
36  
37  
38  
39  
40  
41  
42

Running Head: Self and Learning

Sticky Me:

Self-Relevance Slows Reinforcement Learning

Marius Golubickis<sup>1</sup> and C. Neil Macrae<sup>2</sup>

<sup>1</sup>School of Psychology, University of Plymouth, Plymouth, England, UK

<sup>2</sup>School of Psychology, University of Aberdeen, Aberdeen, Scotland, UK

Address Correspondence to:

Marius Golubickis  
School of Psychology  
University of Plymouth  
Drake Circus  
Plymouth PL4 8AA  
England, UK

Email: [marius.golubickis@plymouth.ac.uk](mailto:marius.golubickis@plymouth.ac.uk)

**Abstract**

A prominent facet of social-cognitive functioning is that self-relevant information is prioritized in perception, attention, and memory. What is not yet understood, however, is whether similar effects arise during learning. In particular, compared to other people (e.g., best friend), is information about the self acquired more rapidly? To explore this matter, here we used a probabilistic selection task in combination with computational modeling (i.e., Reinforcement Learning Drift Diffusion Model analysis) to establish how self-relevance influences learning under conditions of uncertainty (i.e., choices are based on the perceived likelihood of positive and negative outcomes). Across two experiments, a consistent pattern of effects was observed. First, learning rates for both positive and negative prediction errors were slower for self-relevant compared to friend-relevant associations. Second, self-relevant (vs. friend-relevant) learning was characterized by the exploitation (vs. exploration) of choice selections. That is, in a complex (i.e., probabilistic) decision-making environment, previously rewarded self-related outcomes were selected more often than novel — but potentially riskier — alternatives. The implications of these findings for accounts of self-function are considered.

Keywords: self, learning, self-prioritization, probabilistic selection task, reinforcement learning drift diffusion model.

## Sticky Me:

## Self-Relevance Slows Reinforcement Learning

69  
70  
71  
72  
73  
74  
75  
76  
77  
78  
79  
80  
81  
82  
83  
84  
85  
86  
87  
88  
89  
90  
91  
92  
93  
94**1. Introduction**

The self is an indispensable psychological construct, providing coherence and continuity to the narrative that underpins a personal sense of being (Baars, 1988; Baumeister, 1998; Conway, 2005; Conway & Pleydell-Pearce, 2000; Gallagher, 2000; James, 1890; Markus & Nurius, 1986; Markus & Wurf, 1987; Oakley & Halligan, 2017). As Markus and Wurf (1987, pp. 299-300) reported, “the self-concept...interprets and organizes self-relevant actions and experiences, it has motivational consequences, providing the incentives, standards, plans, rules, and scripts for behavior; and it adjusts in response to challenges from the social environment.” In other words, the self serves as a basic processing hub around which social-cognitive functioning unfolds (Humphreys & Sui, 2016; Sui & Humphreys, 2015).

In documenting how the self influences thinking and doing, a common theme runs through decades of research. Regardless of the specific outcome under investigation (e.g., attributions, memories), personal relevance biases information-processing and response selection in self-enhancing and self-prioritizing ways (e.g., Conway, 2005; Mezulis et al., 2004; Sedikides & Alicke, 2012; Sui & Humphreys, 2015; Sui & Rothstein, 2019; Symons & Johnson, 1997). Most strongly associated with the self-reference effect (SRE) in memory (Kelley et al., 2002; Maki & McCaul, 1985; Rogers et al., 1977) — whereby material enjoys a recollective benefit when processed in the context of the self compared to other people (e.g., family members, friends, celebrities) — comparable advantages also emerge when attention and decision-making are probed (e.g., Alexopoulos et al., 2012; Bargh & Pratto, 1986; Constable et al., 2019; Falbén et al., 2020; Gray et al., 2004; Golubickis et al., 2018; Shapiro et al., 1997; Sui et al., 2012, 2015). Indeed, such is the potency of self-prioritization (i.e., the self-prioritization effect [SPE], Sui et al., 2012), benefits accrue even when the stimuli paired with the self (vs. other people) are arbitrary and meaningless,

95 such as geometric shapes, abstract symbols, and colors/sounds (Golubickis et al., 2017, 2020;  
96 Schäfer et al., 2015, 2016; Sui et al., 2012; Wang et al., 2016; Woźniak & Knoblich, 2019; Yin et  
97 al., 2019).

98         Despite an extensive literature cataloguing the effects of self-relevance on core components  
99 of social cognition, important issues nevertheless remain. In particular, aside from a few notable  
100 exceptions, research has largely overlooked the extent to which the personal significance of stimuli  
101 influences a fundamental and crucial facet of daily life, the rate at which information is learned  
102 (Liao et al., 2021; Lockwood et al., 2018). That is, just as self-relevance facilitates the detection,  
103 appraisal, and memorability of stimuli (Humphreys & Sui, 2016; Sui & Humphreys, 2015, 2017;  
104 Symons & Johnson, 1997), so too it may enhance how rapidly this material is acquired. In one of  
105 the few studies to explore this matter, Lockwood et al. (2018) adopted a deterministic associative-  
106 learning task in which participants had to learn, from a pool of fractals (i.e., abstract, unfamiliar  
107 stimuli), which items belonged to various social targets (Brovelli et al., 2008; Schultz et al., 1997).<sup>1</sup>  
108 Specifically, a single fractal appeared on each experimental trial and participants had to report (i.e.,  
109 learn) whether the stimulus was owned by the self, a friend, or a stranger. Feedback was then  
110 provided indicating if the response was correct or incorrect, and the task was deterministic in that  
111 participants were told each target always possessed the same fractals. To establish the respective  
112 target-related learning rates, data were submitted to an associative learning (AL) algorithm (Sutton  
113 & Barto, 1998).

114         Lockwood et al.'s (2018) findings were revealing. Reflecting the operation of an egocentric  
115 decisional strategy (Epley & Gilovich, 2004; Golubickis et al., 2018, 2019), participants tended to  
116 report that the fractal presented on the first trial belonged to them, when in reality it was just as  
117 likely to be owned by either of the other targets. In addition, responses were faster and more  
118 accurate when learning about fractals owned-by-self compared to those that belonged to others.

---

<sup>1</sup> Forming (and probing) target-object associations through ownership is a common methodology to explore self-prioritization (Constable et al., 2011, 2014; 2019; Falbén et al., 2019, 2020; Golubickis et al., 2018, 2019, 2021).

119 Finally, learning rates were higher when acquiring knowledge about the self, although this effect  
120 was only significant when stranger comprised the target of comparison — learning rates for self and  
121 friend were comparable. The absence of a reliable difference in learning rates between self and  
122 friend is interesting as while a self-advantage has frequently been reported when the target of  
123 comparison is best friend (e.g., Ma & Han, 2010; Sui & Han, 2007; Sui et al., 2012, 2013; Zhu et  
124 al., 2007), some research has indicated that the benefits of personal-relevance can be attenuated, or  
125 even eliminated, when the self is compared with an intimate (i.e., highly familiar) other (Bower &  
126 Gilligan, 1979; Kuiper & Rogers, 1979; Symons & Johnson, 1997). Notwithstanding this  
127 observation, Lockwood et al. (2018) provided initial evidence for the biasing effects of self-  
128 relevance on aspects of associative learning.<sup>2</sup>

129 Building upon and extending prior research, here we also explored the extent to which the  
130 personal relevance (or otherwise) of material impacts learning. Our overarching objectives were to  
131 probe the characteristics of self-learning effects in a different task context (i.e., learning  
132 environment) and to establish the pathway through which these effects arise. In so doing, rather  
133 than adopting a deterministic learning paradigm, a probabilistic selection task (PST) was employed  
134 (Frank et al., 2004, 2007). We used this task for a couple of reasons. First, the PST explores  
135 reinforcement learning (RL) in uncertain (vs. certain) task environments (cf. Lockwood et al.,  
136 2018), thus examines the impact of self-relevance when knowledge is acquired under demanding  
137 decision-making conditions. It is possible, for example, that basic components of self-representation  
138 and self-function may prompt learning effects to diverge when studied in uncertain (i.e.,  
139 probabilistic) compared to certain (i.e., deterministic) task settings (Gershman & Daw, 2017).  
140 Second, in combination with recent developments in computational modeling (i.e., Reinforcement  
141 Learning Drift Diffusion Model (RL-DDM) analysis), adoption of the PST enables identification of

---

<sup>2</sup> As Lockwood et al.'s (2018) neural findings are beyond the scope of the current investigation, here we focus only on their behavioral results.

142 the latent psychological processes that underpin RL (Fontanesi et al., 2019; Pedersen & Frank,  
143 2020; Pedersen et al., 2017).

144 In the current PST, participants were presented with three different stimulus pairs (i.e., AB,  
145 CD, EF) — comprising symbols (i.e., Japanese Hiragana characters; see Frank et al., 2004, 2007)  
146 with an item in each pairing (i.e., A, C, E) representing either the self or a friend — and they were  
147 required to learn, following a series of choice selections, which of the symbols was most likely to  
148 denote each target based on feedback that was provided (see Figure 1). Critically, the feedback was  
149 probabilistic and varied for each stimulus pair (i.e., AB = 80% - 20%, CD = 70% - 30%, EF = 60%  
150 - 40%). For example, in AB trials, a choice of stimulus A led to positive feedback on 80% of the  
151 trials, whereas selecting stimulus B resulted in positive reinforcement on 20% of the trials. Thus, in  
152 this PST, learning was accomplished via choice-related feedback. Over numerous choice selections,  
153 participants learned which item in each pairing was more likely to be correct (i.e., represent self or  
154 friend; A, C, E rather than B, D, F) and the task was completed when sufficient levels of accuracy  
155 were achieved for each stimulus pair (Frank et al., 2004, 2007).

156 To identify the mechanisms underpinning learning, computational modeling was undertaken  
157 on the data. Specifically, based on recent developments, a Reinforcement Learning Drift Diffusion  
158 Model (RL-DDM) analysis was adopted (Fontanesi et al., 2019; Pedersen & Frank, 2020; Pedersen  
159 et al., 2017). Integrating sequential sampling and RL models, the RL-DDM pinpoints the  
160 psychological operations that underpin decision-making (i.e., choice selection) and how these are  
161 adjusted as learning progresses (Miletić et al., 2020; Pedersen & Frank, 2020; Ratcliff et al., 2016).  
162 This is realized through the simultaneous hierarchical Bayesian modeling of response time (RT) and  
163 choice data. A drift rate scaling parameter ( $v_{\text{scaling}}$ ) measures sensitivity to feedback and the  
164 exploration-exploitation trade-off (Cohen et al., 2007), such that higher values indicate more  
165 confident learning based on current knowledge (Pedersen et al., 2017). A learning rate parameter  
166 ( $\eta$ ) — ranging from zero to one — quantifies how quickly individuals learn, with larger values  
167 indicating utilization of current feedback (i.e., fast learning), and smaller values reflecting reduced

168 updating from recently experienced outcomes (i.e., slow learning). In this respect, either a single  
169 learning rate ( $\eta$ ) that captures all learning, or separate learning rates for negative and positive  
170 prediction errors ( $\eta^-$  &  $\eta^+$  respectively) can be estimated (Miletić et al., 2020; Pedersen et al., 2017).  
171 Finally, the model also establishes how much evidence is needed to make a decision (i.e., threshold  
172 separation,  $a$ ) and the efficiency of non-decisional processes (e.g., stimulus encoding, response  
173 execution,  $t_0$ ).

174 Central to the current inquiry is the classic exploration-exploitation trade-off that underlies  
175 learning (Cohen et al., 2007; Daw et al., 2006; Sutton & Barto, 1998). Confronted with a decision-  
176 making dilemma, learning can entail either the exploitation of options that have been optimal in the  
177 past or the exploration of alternatives that, in the long run, may prove to be more rewarding (Cohen  
178 et al., 2007). That is, one can either stick with existing knowledge or try something new. Critically,  
179 whereas exploration generally facilitates the acquisition of information, exploitation yields  
180 immediate decisional rewards, but it may impair learning (Sutton & Barto, 1998). As such, whether  
181 self-relevance enhances or reduces learning relative to a target of comparison (e.g., friend) should  
182 be reflected in decisions to explore or exploit the choice selections during RL. In this regard, an  
183 interesting possibility is that, in complex (i.e., probabilistic) task settings, people may prefer to stick  
184 (i.e., exploit) rather than switch (i.e., explore) when to-be-learned material is self-relevant, thereby  
185 prompting a slower learning rate for information pertaining to the self compared to others (cf.  
186 Lockwood et al., 2018). Several strands of evidence suggest such an outcome.

187 According to Humphreys and Sui (2015), via enhanced binding, self-reference serves as a  
188 form of associative glue for perception, attention, and memory (Cunningham et al., 2008; Rogers et  
189 al., 1977; Sui et al., 2012; Wang et al., 2016). While generally facilitating information processing  
190 and response selection, these potent self-object associations can also impede performance in certain  
191 task contexts. For example, participants find it difficult to overcome prior self-shape (vs. friend-  
192 shape) associations when given the task of forming new relations (Wang et al., 2016) and display a  
193 stubborn preference for self-relevant (vs. other-relevant) items during decision-making (Constable

194 et al., 2019; Golubickis et al., 2018, 2019; Lockwood et al., 2018). Although such sticky learning  
195 undoubtedly supports the maintenance of a stable self-concept — an essential component of social-  
196 cognitive functioning (Greenwald, 1980; Markus, 1977) — it also suggests that exploitation rather  
197 than exploration may be the preferred strategy when acquiring information pertaining to the self in  
198 uncertain (i.e., probabilistic) learning environments. That is, previously rewarded self-object  
199 associations may be selected more often than novel (but riskier) options, thereby reducing the  
200 learning rate for the acquisition of personally meaningful material. Accordingly, using a PST in  
201 conjunction with computational modeling, here we explored the possibility that self-relevance may  
202 slow RL relative to an optimal target of comparison (e.g., best friend).

203

## 204 **2. Experiment 1**

### 205 **2.1 Method**

#### 206 **2.1.1 Participants and Design**

207 Fifty participants (33 females, 17 males, 3 others;  $M_{\text{age}} = 23.04$ ,  $SD = 3.06$ ), with normal or  
208 corrected-to-normal visual acuity, took part in the research. Data collection was conducted online  
209 using Prolific Academic ([www.prolific.co](http://www.prolific.co)), with each participant receiving compensation at the rate  
210 of £7.50 (~\$10) per hour. Informed consent was obtained from participants prior to the  
211 commencement of the experiment and the protocol was reviewed and approved by the Ethics  
212 Committee at the School of Psychology, University of Plymouth. The experiment had a single  
213 factor (Correct Symbol: self or friend) repeated-measures design. To detect a significant effect, a  
214 sample of fifty participants afforded 92% power for a large effect size (i.e.,  $d = .80$ ; PANGEA,  
215 v .0.2).

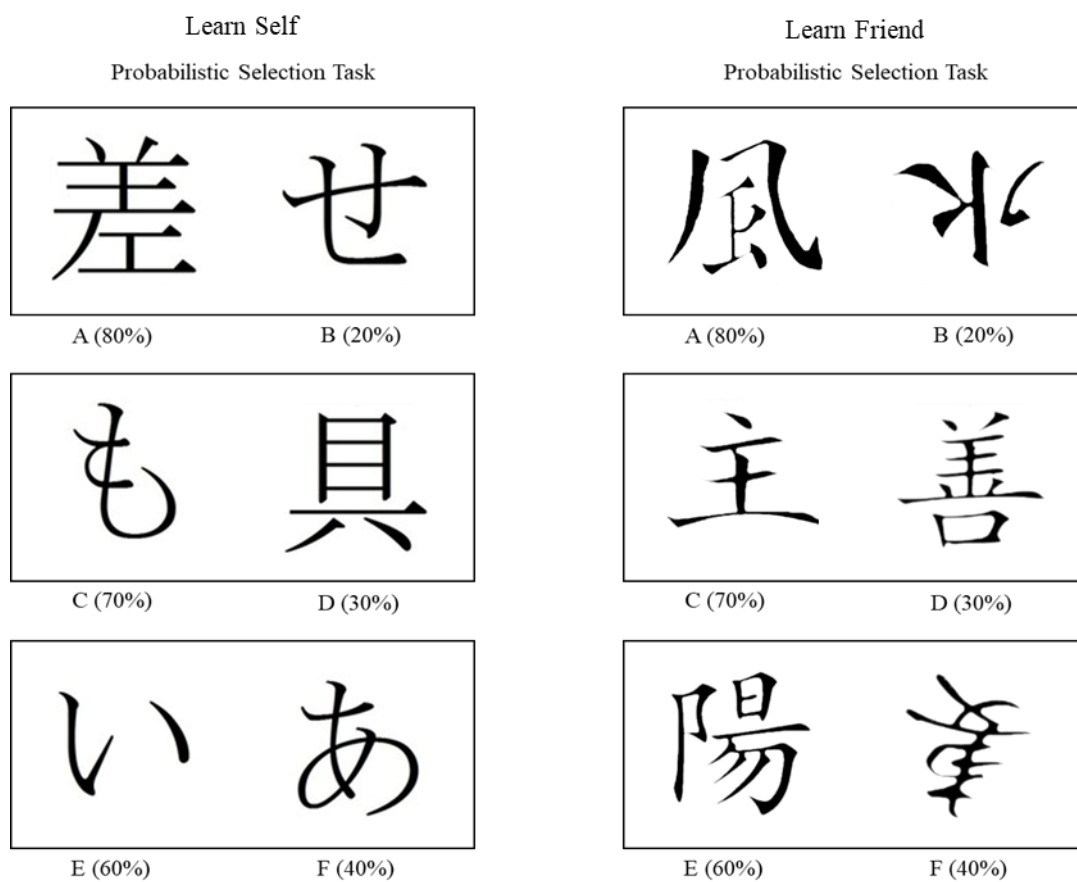
216

#### 217 **2.1.2. Stimulus Materials and Procedure**

218 Participants performed two versions of a PST (Frank et al., 2004, 2007), with each  
219 comprising a learning phase in which three pairs of symbols (denoted as AB, CD, and EF, see



220 Figure 1) were presented. Participants were instructed they were required to learn, based on  
 221 feedback provided, which symbol in each pair was most likely to represent them (i.e., self) or their  
 222 best friend. Following previous research, prior to the task, participants were requested to bring their  
 223 best friend (i.e., target of comparison) to mind (Golubickis et al., 2018). After each choice selection,  
 224 participants were informed that onscreen information would indicate whether their response was  
 225 correct or incorrect. Half of the participants were randomly assigned to perform a version of the  
 226 PST in which self-related symbols were more likely to be correct, followed by another version of  
 227 the task in which friend-related items were more likely to comprise the correct response. That is,  
 228 trial type (i.e., learning) was blocked by target. The order of the PSTs was reversed for the  
 229 remaining participants.  
 230



231

232

233 *Figure 1.* Example of the stimulus pairs (i.e., Japanese Hiragana characters) and the probabilities of  
 234 correct responses during the probabilistic selection task.

235           The probabilities indicating which symbol was more likely to be correct followed the  
236 standard version of the PST (Frank et al., 2004, 2007). Specifically, for the AB pair, A was 80%  
237 likely to be correct (20% for B), for the CD pair, C was 70% likely to be correct (30% for D), and  
238 finally, for the EF pair, E was 60% likely to be correct (40% for F). Over numerous choice  
239 selections, participants learned which item in each pairing was more likely to be correct (i.e., A, C,  
240 E rather than B, D, F) based on the feedback provided. The task was completed when participants  
241 reached sufficient levels of accuracy for each pairing (i.e., AB, 60% or above; CD, 55% or above;  
242 EF, 50% or above; Frank et al., 2004, 2007).

243           Each trial began with the presentation of a pair of symbols that remained on the screen until  
244 the participant made a response. After the participant selected one of the symbols, feedback (i.e.,  
245 the word ‘Correct’ in green or ‘Incorrect’ in red) was presented for 1000 ms, followed by a blank  
246 screen for 500 ms, after which the next trial commenced. Participants had to select a symbol by  
247 pressing the appropriate button on the keyboard (i.e., A for the symbol on the left side of the screen,  
248 L for the symbol on the right side of the screen). The symbols in each pair were equally likely to be  
249 presented on the left or right side of the screen. The experiment was conducted using Inquisit Web.  
250 Participants completed blocks of 60 trials in which each of the three stimulus pairs appeared  
251 randomly, equally often, until accuracy reached a satisfactory level. The maximum number of  
252 learning blocks was set to six (i.e., 360 trials in total) if the participant did not reach satisfactory  
253 levels of accuracy earlier in the task (Frank et al., 2007). On completion of the experiment,  
254 participants were debriefed and thanked.

255

## 256 **2.2 Results and Discussion**

### 257 **2.2.1. Behavioral Analysis**

258           The mean latency and accuracy of choice selections were submitted to a paired-sample  
259 (Correct Symbol: self or friend) *t*-test (two-tailed). No significant difference emerged on either

260 dependent measure (i.e., decision time:  $M_{\text{self}} = 1203$  ms vs.  $M_{\text{friend}} = 1148$  ms; learning performance:  
261  $M_{\text{self}} = 68\%$  vs.  $M_{\text{friend}} = 66\%$ ).

262

### 263 **2.1.2. Modeling Analysis**

264 To identify the processes underpinning learning, data were submitted to a RL-DDM analysis  
265 (Frank et al., 2015; Pedersen & Frank, 2020; Pedersen et al., 2017). This analysis combines the  
266 strengths of RL and sequential-sampling models (SSMs) to elucidate the operations that support  
267 task performance. Specifically, although RL models account for changes in the relative proportion  
268 of choice probabilities over the course of learning, they do not speak to concurrent differences in  
269 response latencies, a fundamental and important dimension of the available data (e.g., as learning  
270 takes place, decision times decrease). In this respect, SSMs (e.g., drift diffusion model; Ratcliff et  
271 al., 2016; Smith & Radcliff, 2004) are useful as they provide a mechanistic account of binary  
272 decision-making by explaining how choice accuracy and response latencies collectively arise from  
273 a common set of latent cognitive processes (e.g., rate of evidence accumulation, response caution).  
274 Thus, crucially, the RL-DDM extends standard RL models by explicating the processes through  
275 which learning unfolds over time (Fontanesi et al., 2019; Miletic et al., 2020; Pedersen & Frank,  
276 2020; Pedersen et al., 2017).

277 Two significant modifications characterize the RL-DDM. First, the typical choice rule for  
278 reinforcement learning (i.e., softmax) is replaced by the drift diffusion model (i.e., Wiener process,  
279 see Miletic et al. 2020; Pedersen et al., 2017). This change is important as it affords the possibility  
280 to model choice and RT data simultaneously. Second, the algorithm that captures the learning of  
281 subjective expectation values from stimuli and actions (i.e., value-based approach) is integrated into  
282 the process of evidence accumulation (i.e., drift rate). Thus, applying the delta learning rule, the  
283 model initially describes the updating of the expected Q-value for a chosen option (e.g., positively  
284 reinforced symbol A) based on the scaled by learning rate ( $\alpha$ ) reward prediction error (i.e., the

285 difference between observed and expected feedback) in the previous trial (Rescorla & Wagner  
 286 1972; Watkins & Dayan 1992, see Eq. 1):

287

$$288 \quad Q_{\text{chosen-option}}(t) = Q_{\text{chosen-option}}(t-1) + \alpha (\text{Reward}(t-1) - Q_{\text{chosen-option}}(t-1)) \quad (1)$$

289

290 Subsequently, the RL-DDM formulates the drift rate ( $v$ ) during reinforced decisions based  
 291 on the difference between the expected value of positively ( $Q_{\text{positively-reinforced}}$ ) and negatively  
 292 ( $Q_{\text{negatively-reinforced}}$ ) reinforced choices. To accommodate the manner in which this knowledge is  
 293 used, the RL-DDM allows an additional free scaling parameter to be estimated (i.e., drift rate  
 294 scaling,  $v_{\text{scaling}}$ ). This scaling parameter is similar to inverse temperature in the softmax choice rule  
 295 and reflects the level of exploration/exploitation during learning (Pedersen & Frank, 2020), such  
 296 that larger values reflect stronger exploitation of the option with the highest expected value (see Eq.  
 297 2).

298

$$299 \quad v(t) = (Q_{\text{positively-reinforced}}(t) - Q_{\text{negatively-reinforced}}(t)) * v_{\text{scaling}} \quad (2)$$

300

301 Thus, in essence, the RL-DDM assumes that evidence is gathered for each choice option  
 302 (e.g., symbol A vs. symbol B) until a critical evidential threshold is reached, at which point a  
 303 response is made. This response threshold is captured by the boundary separation ( $a$ ) parameter,  
 304 and it reflects speed-accuracy trade-offs during decision-making. For example, if a conservative (vs.  
 305 liberal) decision-making style (i.e., higher evidential requirements) is adopted, this would yield  
 306 slower but more accurate responses. At the start of the PST, participants make slow guesses as the  
 307 stimuli have not yet been reinforced, thus the difference in expected values between symbol  
 308 pairings is extremely low (i.e., slow evidence accumulation due to high uncertainty). As participants  
 309 start to receive feedback, via application of the delta learning rule (Rescorla & Wagner, 1972), the  
 310 subjective Q-values of positively/negatively reinforced stimuli increase/decrease. The speed at

311 which participants update the expected values is described by the learning rate ( $\eta$ ) parameter. On a  
312 trial-by-trial basis, this knowledge (i.e., learning which symbol is correct, Q-value) is integrated  
313 into the drift rate such that over time the difference in expected values between reinforced options  
314 (ACE vs. BDF symbol pairings) increases. The larger the difference between positively and  
315 negatively reinforced options, the easier (i.e., faster and more accurate) choice selection becomes  
316 (i.e., fast information sampling).

317 To estimate model parameters, an extension of the Bayesian hierarchical drift diffusion  
318 toolbox was adopted (Wiecki et al., 2013). Models were response-coded, such that the upper  
319 threshold corresponded to responses to stimuli that were positively reinforced (i.e., symbols  
320 corresponding to the letters A, C, & E) and the lower threshold to stimuli that were negatively  
321 reinforced (i.e., symbols corresponding to the letters B, D, & F; Pedersen & Frank, 2020). Bayesian  
322 posterior distributions were modeled using a Markov chain Monte Carlo (MCMC) with 10,000  
323 samples (including 1,000 burn), with outliers (5% of the trials) removed by the HDDM software  
324 (Ratcliff & Tuerlinckx, 2002; Wiecki et al., 2013). Two RL-DDM models were estimated for  
325 comparison (i.e., single vs. dual learning rate model). In the first model, only a single learning rate  
326 ( $\eta$ ) was allowed to vary across Correct Symbol (i.e., self vs. friend). This model examined whether  
327 there were differences in the speed of learning across the experimental conditions without taking  
328 the potential influence of different types of prediction error into consideration. In contrast, in the  
329 second model, learning rates for negative and positive prediction errors ( $\eta^-$  &  $\eta^+$ , respectively) were  
330 allowed to vary by Correct Symbol. As such, this model considered whether learning self-related or  
331 friend-related stimuli was accelerated following negative or positive prediction errors. In both  
332 models, drift rate scaling ( $v_{\text{scaling}}$ ) and boundary separation ( $a$ ) varied across Correct Symbol.

333 Model comparison was performed using the Deviance Information Criterion (DIC) as this  
334 approach is routinely adopted when comparing hierarchical Bayesian models (Spiegelhalter et al.,  
335 1998, 2002). Lower DIC values favor models with the highest likelihood and least number of  
336 parameters. This revealed better fit for the dual (DIC: 60999) compared to the single (DIC: 61059)

337 learning rate model. Examination of the posterior distributions (see Figure 2) revealed differences  
338 in learning rates for negative and positive prediction errors ( $\eta^-$  &  $\eta^+$ ), drift rate scaling ( $v_{\text{scaling}}$ ), and  
339 threshold separation ( $a$ ). Specifically, comparisons yielded very strong evidence that learning rates  
340 were faster for friend compared to self, both for negative ( $p_{\text{Bayes}}(\text{self} < \text{friend}) = .032$ ,  $\text{BF}_{10} = 30$ )  
341 and positive ( $p_{\text{Bayes}}(\text{self} < \text{friend}) < .001$ ,  $\text{BF}_{10} > 1000$ ) prediction errors.<sup>3</sup> In addition, participants  
342 integrated information more efficiently from negative than positive prediction errors, an effect that  
343 was larger for self ( $p_{\text{Bayes}}(\eta^+ < \eta^-) = .008$ ,  $\text{BF}_{10} = 125$ ) than friend ( $p_{\text{Bayes}}(\eta^+ < \eta^-) = .162$ ,  $\text{BF}_{10} = 6$ ).  
344 There was also very strong evidence that drift rate scaling ( $v_{\text{scaling}}$ ) was larger for self- than friend-  
345 related symbols ( $p_{\text{Bayes}}(\text{self} > \text{friend}) = .019$ ,  $\text{BF}_{10} = 52$ ). Finally, for boundary separation ( $a$ ), there  
346 was extremely strong evidence that more decisional information was required when selecting self-  
347 compared to friend-related responses ( $p_{\text{Bayes}}(\text{self} > \text{friend}) < .001$ ,  $\text{BF}_{10} > 1000$ ).

348

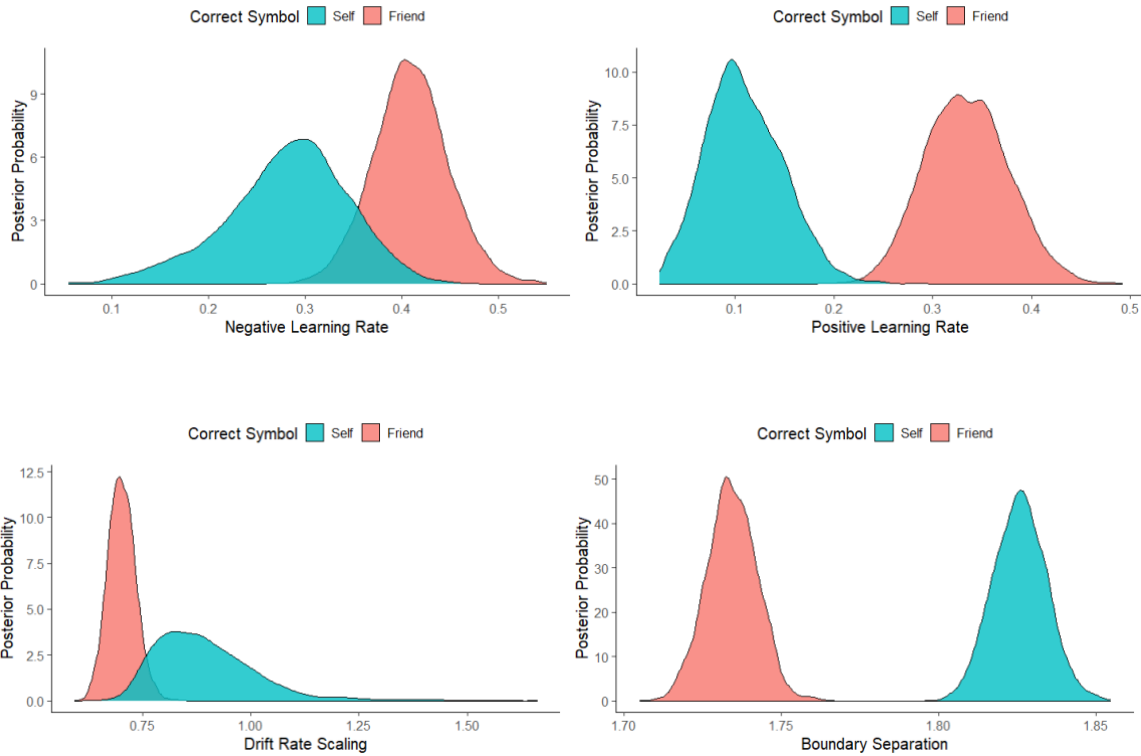
349

350

351

---

<sup>3</sup> Bayes Factors were transformed from Bayesian  $p$ -values (for details see Marsman & Wagenmakers, 2017).



352

353

354

355 *Figure 2.* Mean posterior parameter distributions as a function of Correct Symbol for negative ( $\eta^-$ )  
 356 and positive ( $\eta^+$ ) learning rates, drift rate scaling ( $v_{\text{scaling}}$ ) and boundary separation ( $a$ ).

357

358

359 These findings reveal that, in a probabilistic task context (Frank et al., 2004, 2007), self-  
 360 relevance (vs. friend-relevance) reduced the rate of learning. In addition, the RL-DDM analysis also  
 361 indicated a difference in the balance between the strategies that drive learning — exploration and  
 362 exploitation (Cohen et al., 2007; Sutton & Barto, 1998). Specifically, as indexed by the drift rate  
 363 scaling parameter ( $v_{\text{scaling}}$ ), self-relevant (vs. friend-relevant) trials were characterized by the  
 364 tendency to exploit previously rewarded outcomes rather than explore new alternatives. In other  
 365 words, self-relevance elicited a greater sensitivity to current outcomes (i.e., existing knowledge)  
 366 during learning (Pedersen et al., 2017).

367 To probe the reproducibility of these effects, in our next experiment we also explored how  
 368 self-relevance influenced learning in a PST (Frank et al., 2004, 2007), but with an important

369 methodological modification. Rather than blocking the PST (i.e., learning) by target, participants  
370 simultaneously learned about self and friend in an intermixed design as previous research has  
371 demonstrated that self-relevance exerts a greater influence on decisional processing under these  
372 conditions (Golubickis & Macrae, 2021). Replicating Experiment 1, we expected self-relevance (vs.  
373 friend-relevance) to reduce the rate of learning and favor exploitation (vs. exploration) of the choice  
374 selections.

375

## 376 **3. Experiment 2**

### 377 **3.1. Method**

#### 378 **3.1.1. Participants and Design**

379 Thirty-four participants (22 females, 10 males, 2 others;  $M_{\text{age}} = 22.97$ ,  $SD = 2.62$ ), with  
380 normal or corrected-to-normal visual acuity, took part in the research. Data collection was  
381 conducted online using Prolific Academic ([www.prolific.co](http://www.prolific.co)), with each participant receiving  
382 compensation at the rate of £7.50 (~\$10) per hour. Informed consent was obtained from participants  
383 prior to the commencement of the experiment and the protocol was reviewed and approved by the  
384 Ethics Committee at the School of Psychology, University of Plymouth. The experiment had a  
385 single factor (Correct Symbol: self or friend) repeated-measures design. To detect a significant  
386 effect, a sample of thirty-four participants afforded 80% power for a large effect size (i.e.,  $d = .80$ ;  
387 PANGEA, v .0.2).

388

#### 389 **3.1.2. Stimulus Materials and Procedure**

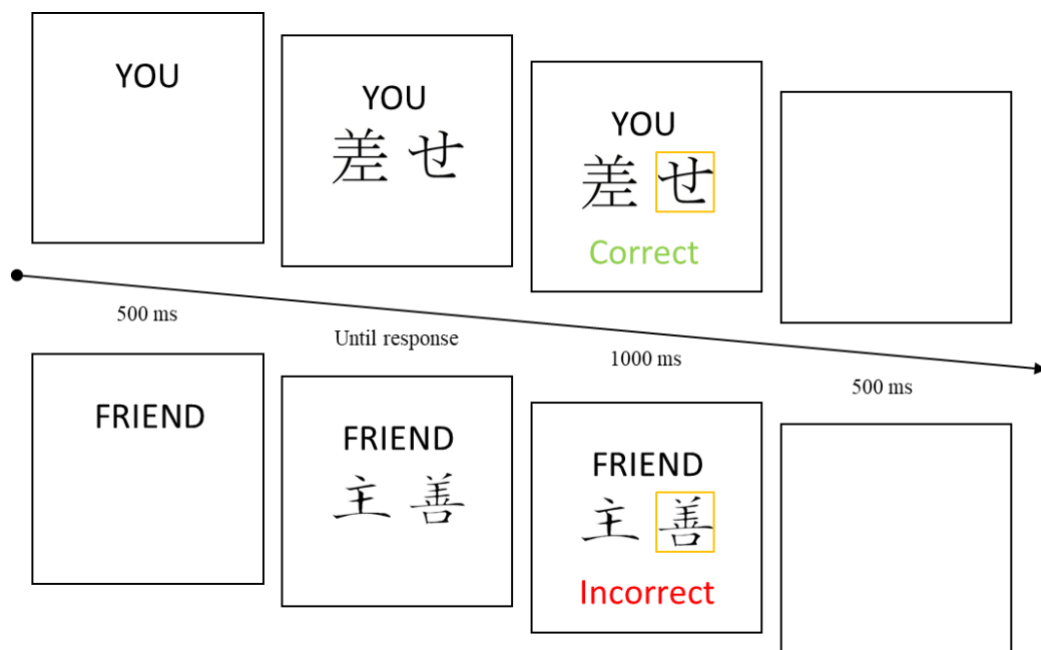
390 A modified version of the PST from Experiment 1 was adopted. Specifically, on a trial-by-  
391 trial basis, participants were required to learn which symbol in each pairing was more likely to  
392 represent self or best friend. Before the presentation of each stimulus pair, a cue (i.e., the labels  
393 “YOU” or “FRIEND”) appeared on the screen indicating the target to which the symbols pertained  
394 (see Figure 3). The cue appeared 500 ms before the symbols and remained on the screen, above the



395 stimuli, until a response was made. Participants completed blocks of 120 trials (i.e., 60 self and 60  
 396 friend) in which each stimulus pair appeared randomly, equally often, until accuracy reached a  
 397 satisfactory level. The maximum number of learning blocks was set to three (i.e., 360 trials in total)  
 398 if the participant did not reach satisfactory levels of accuracy earlier in the task (Frank et al., 2007).  
 399 In all other respects, the procedure was identical to Experiment 1.

400

401



402

403 *Figure 3.* Examples of the experimental trials.

404

405

## 406 3.2. Results and Discussion

### 407 3.2.1. Behavioral Analysis

408 Four participants (3 females) failed to learn the probabilities associated with the symbols,  
 409 thus were excluded from the analyses. The mean latency and accuracy of choice selections were  
 410 submitted to a paired-sample (Correct Symbol: self or friend) *t*-test (two-tailed). The analysis of  
 411 choice latencies revealed faster responses to self-related compared to friend-related symbols,  $t(29) =$

412 2.77,  $p = .010$ ,  $d = .51$ ; respective  $M$ s: 1546 ms vs. 1689 ms). In addition, accuracy was greater for  
 413 self-related than friend-related stimuli,  $t(29) = 3.39$ ,  $p = .002$ ,  $d = .62$ ; respective  $M$ s: 70% vs. 63%).

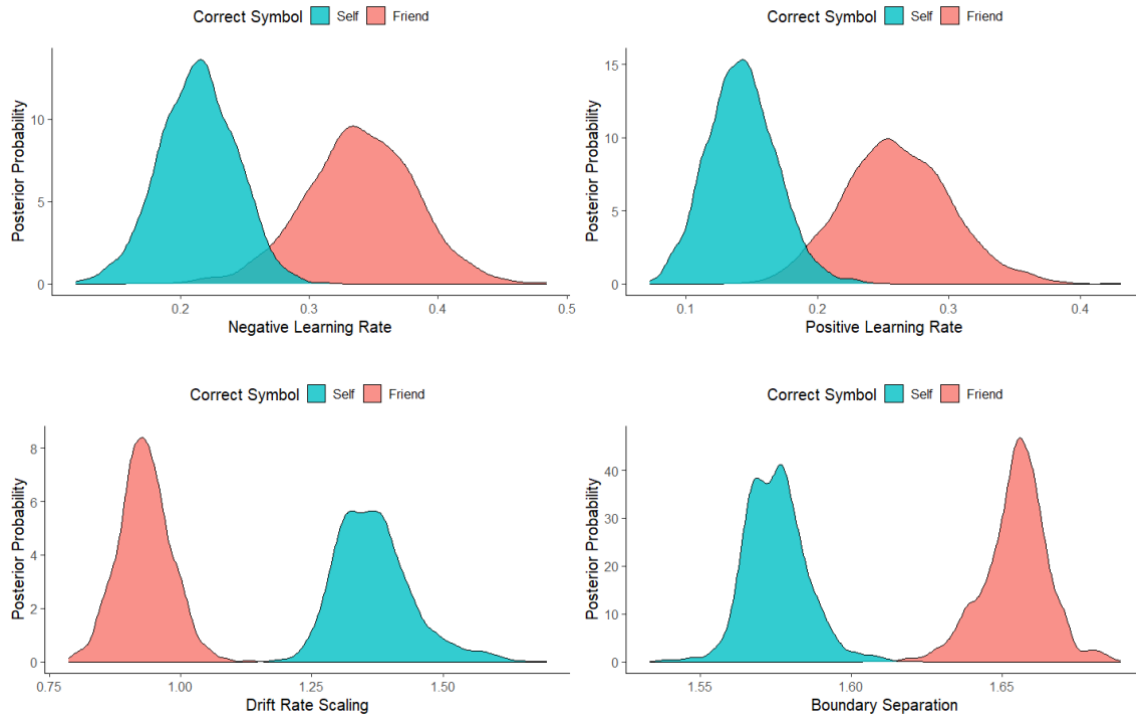
414

### 415 3.2.2. Modeling Analysis

416 To identify the processes underpinning learning, data were submitted to a RL-DDM analysis  
 417 following the same modeling procedure as Experiment 1. As previously, fit was better for the dual  
 418 (DIC: 43524) compared to the single (DIC: 43541) learning rate model. Examination of the  
 419 posterior distributions (see Figure 4) revealed differences in learning rates for negative and positive  
 420 prediction errors ( $\eta^-$  &  $\eta^+$ ), drift rate scaling ( $v_{\text{scaling}}$ ), and threshold separation ( $a$ ). Specifically,  
 421 comparisons yielded very strong evidence that learning rates were faster for friend compared to self,  
 422 both for negative ( $p_{\text{Bayes}}(\text{self} < \text{friend}) = .011$ ,  $\text{BF}_{10} = 90$ ) and positive ( $p_{\text{Bayes}}(\text{self} < \text{friend}) = .005$ ,  
 423  $\text{BF}_{10} = 199$ ) prediction errors. As in Experiment 1, participants integrated information more  
 424 efficiently from negative than positive prediction errors, an effect that was larger for self ( $p_{\text{Bayes}}(\eta^+ < \eta^-)$   
 425  $= .03$ ,  $\text{BF}_{10} = 33$ ) than friend ( $p_{\text{Bayes}}(\eta^+ < \eta^-) = .10$ ,  $\text{BF}_{10} = 10$ ). There was also extremely  
 426 strong evidence that drift rate scaling ( $v_{\text{scaling}}$ ) was larger for self-related than friend-related symbols  
 427 ( $p_{\text{Bayes}}(\text{self} > \text{friend}) < .001$ ,  $\text{BF}_{10} > 1000$ ). Finally, for boundary separation ( $a$ ), there was extremely  
 428 strong evidence that more decisional information was required when selecting friend- compared to  
 429 self-related responses ( $p_{\text{Bayes}}(\text{self} < \text{friend}) < .001$ ,  $\text{BF}_{10} > 1000$ ).

430

431



432

433

434

435 Figure 4. Mean posterior parameter distributions as a function of Correct Symbol for negative ( $\eta^-$ )  
 436 and positive ( $\eta^+$ ) learning rates, drift rate scaling ( $v_{\text{scaling}}$ ) and boundary separation ( $a$ ).

437

438

439 Using a different experimental design, these findings replicated the effects observed in  
 440 Experiment 1. First, for both negative and positive prediction errors, learning rates were slower for  
 441 self-related compared to friend-related symbols. Second, reflecting a greater reliance on existing  
 442 knowledge (i.e., sensitivity to current outcomes), self-relevant (vs. friend-relevant) trials were  
 443 characterized by the tendency to exploit previously rewarded outcomes rather than explore new  
 444 choice selections (Pedersen et al., 2017). Interestingly, unlike Experiment 1 in which response  
 445 caution was greater for self-relevant compared to friend-relevant symbols, this effect was reversed  
 446 in the current experiment. This reversal can likely be traced to task-specific differences in the  
 447 presentation of the stimulus trials during the PST (i.e., Expt. 1 - blocked by target; Expt. 2 -  
 448 intermixed; Golubickis & Macrae, 2021).

449

#### 450 **4. General Discussion**

451           Notwithstanding the acknowledged benefits that self-relevance exerts on information  
452 processing and response selection (Humphreys & Sui, 2016; Sui & Humphreys, 2015, 2017;  
453 Symons & Johnson, 1997), here we demonstrated a quite different effect. In the context of a PST,  
454 self-relevance (vs. friend-relevance) reduced the rate at which information was acquired.  
455 Specifically, whether stimuli were blocked by target (Expt. 1) or intermixed (Expt. 2), learning rates  
456 were slower for self-related compared to friend-related associations. In addition, self-relevant (vs.  
457 friend-relevant) learning was characterized by the tendency to exploit rather than explore the choice  
458 selections during the task (Cohen et al., 2007; Sutton & Barto, 1998). This indicates that, in a  
459 complex (i.e., probabilistic) decision-making setting, previously rewarded self-related outcomes  
460 were chosen more often than novel — but potentially riskier — choice selections. In other words,  
461 when learning about the self (vs. friend), participants tended to rely on their existing knowledge,  
462 thereby trading enhanced future learning for guaranteed current rewards (Pedersen et al., 2017).

463           That self-relevance has the capacity to impair performance in certain task contexts is  
464 unsurprising. Forging immediate and powerful target-object associations in working memory,  
465 personal-relevance (vs. friend-relevance) yields substantial processing benefits when responding is  
466 driven by the enhanced accessibility of these relations (Humphreys & Sui, 2016; Sui & Humphreys,  
467 2015, 2017). That is, highly accessible self-object associations — even when the stimuli in question  
468 are unfamiliar and trivial — give rise to rapid and accurate responses (e.g., Golubickis et al., 2017,  
469 2020; Schäffer et al., 2016, 2017; Stein et al., 2016; Sui et al., 2012, 2013; Woźniak & Knoblich,  
470 2019). The strength of these sticky associations, however, can also hinder performance, particularly  
471 when participants must override previous learning experiences and acquire new target-object  
472 relations (Constable & Knoblich, 2020; Wang et al., 2016). For example, Wang et al. (2016)  
473 reported that, once self-shape associations were formed, participants found it difficult to break (i.e.,  
474 undo) these relations and associate the shapes with a new target (e.g., friend). As they reported (p.

475 255), "...self-association can either enhance or disrupt processing, depending on whether new  
476 associations are assessed or whether old associations have to be discarded."

477 By enhancing the binding of target-object relations, self-relevance has obvious implications  
478 for decision-making and learning, at least in settings in which these associations are a task-relevant  
479 component of the methodology (Caughey et al., 2021; Constable et al., 2019; Falbén et al., 2019;  
480 Woźniak & Knoblich, 2021). As demonstrated here, in a PST (Frank et al., 2004, 2007), learning  
481 rates were slower when material was self-relevant (vs. friend-relevant). Several factors probably  
482 contributed to the emergence of this effect. Most notably, by shifting the balance toward  
483 exploitation rather than exploration during RL, choice selections served both to bolster the stability  
484 of the self-concept and optimize response-related rewards. A basic component of social-cognitive  
485 functioning is the possession (and maintenance) of a stable self-concept (Greenwald, 1980; Markus,  
486 1977). In this respect, favoring choice selections that previously were (correctly) associated with the  
487 self would unquestionably service this objective.

488 In addition, the reward value of self-relevant (vs. friend-relevant) outcomes would similarly  
489 encourage exploitation over exploration (Cohen et al., 2007). According to Northhoff and Hayes  
490 (2011), self-referential processing is underpinned by the intrinsic reward-related properties of self-  
491 relevant stimuli (Northhoff & Hayes, 2011). Given the pivotal role of reward value during learning  
492 (Dayan & Belleine, 2002; Schultz, 1998; Sutton & Barto, 1998), exploiting formerly successful  
493 self-related outcomes would be particularly appealing (i.e., dopamine uptake), much more so than  
494 comparable friend-related responses or the exploration of novel choice selections. As such,  
495 although the precise relationship between self and reward remains a matter of continued scrutiny  
496 and debate (Sui et al., 2015; Stolte et al., 2015), during probabilistic learning this connection is  
497 likely intimate. Interestingly, in each of the reported experiments, learning was more effective  
498 following negative than positive prediction errors, an effect that was most pronounced for the self  
499 (vs. friend). It is possible that the tendency to exploit rather explore choice selections during self-

500 related learning (i.e., sticky self-symbol associations) may underpin this asymmetry. Future  
501 research should explore this possibility.

502         Although, in the current investigation, the rate of learning was slower for self-relevant  
503 compared to friend-relevant stimuli, it is unlikely this effect is immutable. Indeed, as noted earlier,  
504 Lockwood and colleagues (2018) reported that, during deterministic learning, personal (vs. other)  
505 associations were formed most rapidly, albeit only when stranger comprised the target of  
506 comparison. For a familiar target of comparison (i.e., friend), self-other learning rates did not differ  
507 significantly. These inconsistent findings potentially derive from differences in self-function across  
508 probabilistic and deterministic learning environments (Gershman & Daw, 2017). In a fully certain  
509 (i.e., deterministic) world, exploration is not a viable strategy as pursuing new choice selections  
510 following positive feedback would impair performance. In contrast, in probabilistic settings (e.g.,  
511 PSTs) feedback is accompanied by uncertainty (Frank et al., 2004, 2007), thereby moderating the  
512 balance between the competing strategies that drive choice selections (i.e., exploration-exploitation  
513 trade-off). As was observed in the current experiments, self-relevant (vs. friend-relevant) learning  
514 was characterized by the tendency to exploit rather than explore the response-related outcomes,  
515 such that potentially enhanced knowledge acquisition was traded for the certainty of immediate  
516 rewards (Cohen et al., 2007). This suggests that, depending on the characteristics of the learning  
517 environment (i.e., deterministic vs. probabilistic), self-relevance can exert quite different effects on  
518 RL.

519         Operating in this flexible way, learning mirrors the other domains in which the effects of  
520 self-relevance have been explored (e.g., attention, memory, decision-making). Inspection of a  
521 rapidly developing literature reveals the inherent malleability of self-prioritization and the divergent  
522 cognitive origins of self-bias. Specifically, whether self-prioritization facilitates or impedes  
523 performance — or indeed arises at all — is highly contingent upon the way in which self-object  
524 associations are operationalized, established, and probed (Caughey et al., 2021; Constable et al.,  
525 2019; Falbén et al., 2019, 2020; Golubickis et al., 2020, 2021; Macrae et al., 2017, 2018; Siebold et

526 al., 2015; Stein et al., 2016; Svensson et al., 2021; Wang et al., 2016; Woźniak & Knoblich, 2021).  
527 Moreover, whereas in some task contexts self-relevance influences the efficiency of stimulus  
528 processing (Golubickis et al., 2017, 2020), in others it impacts response-related operations  
529 (Constable et al., 2019; Falbén et al., 2020; Golubickis et al., 2018, 2019). A useful task for future  
530 research will therefore be to establish how this contextual-dependency modulates the acquisition of  
531 self-knowledge across learning environments that vary in important ways; including the identity  
532 and number of targets of comparison, the characteristics of the to-be-learned material, and the  
533 distribution of rewards (Haruno & Kawato, 2006; Lockwood et al., 2018; Knowlton et al., 1994).

534 Attention should also be directed to the task context in which information pertaining to the  
535 self and others is encountered. Here differences in response caution were observed across two  
536 instrumental learning experiments that differed in task structure. Specifically, whereas response  
537 caution was greater on self-relevant compared to friend-relevant trials when stimuli were blocked  
538 by target (i.e., Experiment 1), this effect was reversed when the trial types were intermixed (i.e.,  
539 Experiment 2). Relatedly, both Golubickis and Macrae (2021) and Desebrock et al. (in press) have  
540 similarly demonstrated the sensitivity of self-referential processing to the characteristics of the task  
541 environment. For example, using a shape-label matching task, Golubickis and Macrae (2021)  
542 observed a reduction in self-prioritization when stimuli were intermixed compared to blocked by  
543 target. Extending this finding, again in a shape-label matching task but using unisensory and  
544 multisensory stimuli, Desebrock et al. (in press) found that self-prioritization was greatest when  
545 trials were blocked by sensory modality. Collectively, these findings highlight the contextual  
546 dependence of self-bias, a factor that has largely been overlooked in research to date.

547 Consideration should also be given to the neural mechanisms that support the learning of  
548 material pertaining to the self and others. For example, is the acquisition of person-related  
549 knowledge underpinned by the same associative operations that drive reward-based learning in non-  
550 social contexts? Given the established role of the pre-frontal cortex (PFC) during self-referential  
551 processing (Kelley et al., 2002; Mitchell et al., 2002, 2006; Sui et al., 2013), it is interesting to note

552 that resolution of the exploration-exploitation dilemma is also associated with activation in this  
553 region (Blanchard & Gershman, 2018; Domenech et al., 2020). Specifically, whereas activity in the  
554 ventromedial PFC (vmPFC) indexes the subjective value of outcomes given the action plan that is  
555 currently in place, modulation in dorsomedial PFC (dmPFC) reflects a reduction in these values and  
556 the generation of new response-related strategies (Donoso et al., 2014). In their investigation of the  
557 neural correlates of self-learning, Lockwood et al. (2018) reported that no brain area tracked  
558 exclusively with self-bias (i.e., self-ownership effect) during a deterministic learning task.  
559 Nevertheless, vmPFC responded more strongly to self- compared to stranger-related (but not friend-  
560 related) associations. As the current experiments yielded differences in both learning rates and the  
561 drift-rate scaling parameter (i.e., exploration-exploitation trade-off) for self and friend, it would  
562 therefore be interesting to explore the neural mechanisms that underlie self/other learning during a  
563 PST. In such a task setting, distinct patterns of activation may emerge in the mPFC and other  
564 cortical regions that support learning (e.g., anterior cingulate cortex [ACC]; Kennerley et al., 2006;  
565 Holroyd & McClure, 2015).

566

## 567 **5. Conclusion**

568 Using a PST in combination with a RL-DDM analysis, here we considered how self-  
569 relevance influences instrumental learning. Across two experiments, learning rates were slower for  
570 self-related compared to friend-related associations and self-relevant (vs. friend-relevant) learning  
571 was characterized by exploitation (vs. exploration) of the choice selections. Together with related  
572 research (Lockwood et al., 2018), these findings affirm the utility of computational approaches in  
573 the investigation of core social-cognitive topics (Hackel & Amodio, 2018; Lockwood & Klein-  
574 Flügge, 2020). Continuing in this way, further research should clarify exactly when, how, and for  
575 whom self-relevance influences associative learning.

576

577



## References

- 578
- 579 Alexopoulos, T., Muller, D., Ric, F., & Marendaz, C. (2012). I, me, mine: Automatic attentional  
580 capture by self-related stimuli, *European Journal of Social Psychology*, *42*, 770-779.
- 581 Baars, B. J. (1988). *A cognitive theory of consciousness*. New York: Cambridge University Press.
- 582 Bargh, J. A., & Pratto, F. (1986). Individual construct accessibility and perceptual  
583 selection. *Journal of Experimental Social Psychology*, *22*, 293-311.
- 584 Baumeister, R. F. (1998). The self. In D. T. Gilbert, S. T. Fiske, & G. Lindzey (Eds.), *Handbook of*  
585 *social psychology* (4<sup>th</sup> ed., pp. 680-740). New York: McGraw-Hill.
- 586 Blanchard, T. C., & Gershman, S. J. (2018). Pure correlates of exploration and exploitation in the  
587 human brain. *Cognitive, Affective, & Behavioral Neuroscience*, *18*, 117-126.
- 588 Bower, G. H., & Gilligan, S. G. (1979). Remembering information related to one's self. *Journal of*  
589 *Research in Personality*, *13*, 420-432.
- 590 Brovelli, A., Laksiri, N., Nazarian, B., Meunier, M., & Boussaoud, D. (2008). Understanding the  
591 neural computations of arbitrary visuomotor learning through fMRI and associative learning  
592 theory. *Cerebral Cortex*, *18*, 1485-1495.
- 593 Caughey, S., Falbén, J. K., Tsamadi, D., Persson, L. M., Golubickis, M., & Macrae, C. N. (2021).  
594 Self-prioritization during stimulus processing is not obligatory. *Psychological Research*, *85*,  
595 503-508.
- 596 Cohen, J. D., McClure, S. M., & Yu, A. J. (2007). Should I stay or should I go? How the human  
597 brain manages the trade-off between exploitation and exploration. *Philosophical*  
598 *Transactions of the Royal Society B*, *362*, 933-942.
- 599 Constable, M. D., & Knoblich, G. (2020). Sticking together? Re-binding previous other-associated  
600 stimuli interferes with self-verification but not partner-verification. *Acta Psychologica*, *210*,  
601 103167.
- 602 Constable, M. D., Kritikos, A., & Bayliss, A. P. (2011). Grasping the concept of personal property.  
603 *Cognition*, *119*, 430-437.

- 604 Constable, M. D., Kritikos, A., Lipp, O. V., & Bayliss, A. P. (2014). Object ownership and action:  
605 The influence of social context and choice on the physical manipulation of personal  
606 property. *Experimental Brain Research*, *232*, 3749-3761.
- 607 Constable, M. D., Welsh, T. N., Huffman, G., & Pratt, J. (2019). I before U: Temporal order  
608 judgements reveal bias for self-owned objects. *Quarterly Journal of Experimental*  
609 *Psychology*, *72*, 589-598.
- 610 Conway, M. A. (2005). Memory and the self. *Journal of Memory and Language*, *53*, 594-628.
- 611 Conway, M. A., & Pleydell-Pearce, C. W. (2000). The construction of autobiographical memories  
612 in the self-memory system. *Psychological Review*, *107*, 261-288.
- 613 Cunningham, S. J., Turk, D. J., Macdonald, L. M., & Macrae, C. N. (2008). Yours or mine?  
614 Ownership and memory. *Consciousness and Cognition*, *17*, 312-318.
- 615 Daw, N. D., O'Doherty, J. P., Dayan, P., Seymour, B., & Dolan, R. J. (2006). Cortical substrates for  
616 exploratory decisions in humans. *Nature*, *441*, 876-879.
- 617 Dayan, P., & Balleine, B. W. (2002). Reward, motivation, and reinforcement learning. *Neuron*, *36*,  
618 285-298.
- 619 Domenech, P., Rheims, S., & Koechlin, E. (2020). Neural mechanisms resolving exploitation-  
620 exploration dilemmas in the medial prefrontal cortex. *Science*, *369*, eabb0184.
- 621 Donoso, M., Collins, A. G., & Koechlin, E. (2014). Foundations of human reasoning in the  
622 prefrontal cortex. *Science*, *344*, 1481-1486.
- 623 Epley, N., & Gilovich, T. (2004). Are adjustments insufficient? *Personality and Social Psychology*  
624 *Bulletin*, *30*, 447-460.
- 625 Falbén, J. K., Golubickis, M., Balseryte, R., Persson, L. M., Tsamadi, D., Caughey, S., & Macrae,  
626 C. N. (2019). How prioritized is self-prioritization during stimulus processing? *Visual*  
627 *Cognition*, *27*, 46-51.
- 628

- 629 Falbén, J. K., Golubickis, M., Wischerath, D., Tsamadi, D., Persson, L. M., Caughey, S., Svensson,  
630 S., & Macrae, C. N. (2020). It's not always about me: The effects of prior beliefs and  
631 stimulus prevalence on self-other prioritization. *Quarterly Journal of Experimental*  
632 *Psychology*, *73*, 1466-1480.
- 633 Fontanesi, L., Gluth, S., Spektor, M. S., & Rieskamp, J. (2019). A reinforcement learning diffusion  
634 decision model for value-based decisions. *Psychonomic Bulletin and Review*, *26*, 1099-  
635 1121.
- 636 Frank, M. J., Moustafa, A. A., Haughey, H. M., Curran, T., & Hutchinson, K. E. (2007). Genetic  
637 triple dissociation reveals multiple roles for dopamine in reinforcement learning.  
638 *Proceedings in the National Academy of Sciences*, *104*, 16311-16316.
- 639 Frank, M. J., Seeberger, L. C., & O'Reilly, R. C. (2004). By carrot or by stick: Cognitive  
640 reinforcement learning in Parkinsonism. *Science*, *306*, 1940-1943.
- 641 Gallagher, S. (2000). Philosophical conceptions of the self: Implications for cognitive science.  
642 *Trends in Cognitive Sciences*, *4*, 14-21.
- 643 Gershman, S. J., & Daw, N. D. (2017). Reinforcement learning and episodic memory in humans  
644 and animals: An integrative framework. *Annual Review of Psychology*, *68*, 101-128.
- 645 Golubickis, M., & Macrae, C. N. (2021). Judging me and you: Task design modulates self-  
646 prioritization. *Acta Psychologica*, *218*, 103350.
- 647 Golubickis, M., Falbén, J. K., Sahraie, A., Visokomogilski, A., Cunningham, W. A., Sui, J., &  
648 Macrae, C. N. (2017). Self-prioritization and perceptual matching: The effects of temporal  
649 construal. *Memory & Cognition*, *45*, 1223-1239.
- 650 Golubickis, M., Falbén, J. K., Ho, N. S., Sui, J., Cunningham, W. A., & Macrae, C. N. (2020). Parts  
651 of me: Identity-relevance moderates self-prioritization. *Consciousness and Cognition*, *77*,  
652 102848.

- 653 Golubickis, M., Falbén, J. K., Cunningham, W. A., & Macrae, C. N. (2018). Exploring the self-  
654 ownership effect: Separating stimulus and response biases. *Journal of Experimental*  
655 *Psychology: Learning, Memory and Cognition*, *44*, 295-306.
- 656 Golubickis, M., Ho, N. S. P., Falbén, J. K., Mackenzie, K. M., Boschetti, A., Cunningham, W. A.,  
657 & Macrae, C. N. (2019). Mine or mother's? Exploring the self-ownership effect across  
658 cultures. *Culture and Brain*, *7*, 1-25.
- 659 Golubickis, M., Ho, N. S. P., Falbén, J. K., Schwertel, C. L., Maiuri, A., Dublas, D., Cunningham,  
660 W. A., & Macrae, C. N. (2021). Valence and ownership: Object desirability influences self-  
661 prioritization. *Psychological Research*, *85*, 91-100.
- 662 Gray, H. M., Ambady, N., Lowenthal, W. T., & Deldin, P. (2004). P300 as an index of attention to  
663 self-relevant stimuli. *Journal of Experimental Social Psychology*, *40*, 216-224.
- 664 Greenwald, A. G. (1980). The totalitarian ego: Fabrication and revision of personal identity.  
665 *American Psychologist*, *35*, 603-618.
- 666 Hackel, L. M., & Amodio, D. M. (2018). Computational neuroscience approaches to social  
667 cognition. *Current Opinion in Psychology*, *24*, 92-97.
- 668 Haruno, M., & Kawato, M. (2006). Different neural correlates of reward expectation and reward  
669 expectation error in the putamen and caudate nucleus during stimulus-action-reward  
670 association learning. *Journal of Neurophysiology*, *95*, 948-959.
- 671 Holroyd, C. B., & McClure, S. M. (2015). Hierarchical control over effortful behavior by rodent  
672 medial frontal cortex: A computational model. *Psychological Review*, *122*, 54-83.
- 673 Humphreys, G. W., & Sui, J. (2016). Attentional control and the self: The self-attention network  
674 (SAN). *Cognitive Neuroscience*, *7*, 5-17.
- 675 James, W. (1890). *The principles of psychology*. New York: Henry-Holt & Co.
- 676 Kelley, W. M., Macrae, C. N., Wyland, C. L., Caglar, S., Inati, S., & Heatherton, T. F. (2002).  
677 Finding the self? An event-related fMRI study. *Journal of Cognitive Neuroscience*, *14*, 785-  
678 794.

- 679 Kennerley, S. W., Walton, M. E., Behrens, T. E., Buckley, M. J., & Rushworth, M. F. (2006).  
680 Optimal decision making and the anterior cingulate cortex. *Nature Neuroscience*, *9*, 940-  
681 947.
- 682 Knowlton, B. J., Squire, L. R., & Gluck, M. A. (1994). Probabilistic classification learning in  
683 amnesia. *Learning & Memory*, *1*, 106-120.
- 684 Kuiper, N. A., & Rogers, T. B. (1979). Encoding of personal information: Self–other  
685 differences. *Journal of Personality and Social Psychology*, *37*, 499-514.
- 686 Liao, Z., Huang, L., & Luo, S. (2021). Intranasal oxytocin decreases self-oriented  
687 learning. *Psychopharmacology*, *238*, 461-474.
- 688 Lockwood, P. L., & Klein-Flugge, M. C. (2020). Computational modelling of social cognition and  
689 behaviour: A reinforcement learning primer. *Social Cognitive and Affective Neuroscience*,  
690 *16*, 761-771.
- 691 Lockwood, P. L., Wittmann, M. K., Apps, M. A., Klein-Flügge, M. C., Crockett, M. J., Humphreys,  
692 G. W., & Rushworth, M. F. (2018). Neural mechanisms for learning self and other  
693 ownership. *Nature Communications*, *9*, 1-11.
- 694 Ma, Y., & Han, S. (2010). Why we respond faster to the self than to others? An implicit positive  
695 association theory of self-advantage during implicit face perception. *Journal of*  
696 *Experimental Psychology: Human Perception and Performance*, *36*, 619-633.
- 697 Macrae, C. N., Visokomogilski, A., Golubickis, M., Cunningham, W. A., & Sahraie, A. (2017).  
698 Self-relevance prioritizes access to visual awareness. *Journal of Experimental Psychology:*  
699 *Human Perception and Performance*, *43*, 438-443.
- 700 Macrae, C. N., Visokomogilski, A., Golubickis, M., & Sahraie, A. (2018). Self-relevance enhances  
701 the benefits of attention on perception. *Visual Cognition*, *26*, 475-481.
- 702 Maki, R. H., & McCaul, K. D. (1985). The effects of self-reference versus other-reference on the  
703 recall of traits and nouns. *Bulletin of the Psychonomic Society*, *23*, 169-172.

- 704 Markus, H. (1977). Self-schemata and processing information about the self. *Journal of Personality*  
705 *and Social Psychology, 35*, 63-78.
- 706 Markus, H. R., & Nurius, P. (1986). Possible selves. *American Psychologist, 41*, 954-969
- 707 Markus, H., & Wurf, E. (1987). The dynamic self-concept: A social psychological  
708 perspective. *Annual Review of Psychology, 38*, 299-337.
- 709 Marsman, M., & Wagenmakers, E. -J. (2017). Three insights from a Bayesian interpretation of the  
710 one-sided  $p$  value. *Educational and Psychological Measurement, 77*, 529-539.
- 711 Mezulis, A. H., Abramson, L. Y., Hyde, J. S., & Hankin, B. L. (2004). Is there a universal positivity  
712 bias in attributions? A meta-analytic review of individual, developmental, and cultural  
713 differences in the self-serving attributional bias. *Psychological Bulletin, 130*, 711-747.
- 714 Miletić, S., Boag, R. J., & Forstmann, B. U. (2020). Mutual benefits: Combining reinforcement  
715 learning with sequential sampling models. *Neuropsychologia, 136*, 107261.
- 716 Mitchell, J. P., Heatherton, T. F., & Macrae, C. N. (2002). Distinct neural systems subserve person  
717 and object knowledge. *Proceedings of the National Academy of Sciences, 99*, 15238-15243.
- 718 Mitchell, J. P., Macrae, C. N., & Banaji, M. R. (2006). Dissociable medial prefrontal contributions  
719 to judgments of similar and dissimilar others. *Neuron, 50*, 655-663.
- 720 Northoff, G., & Hayes, D. J. (2011). Is our self nothing but reward? *Biological Psychiatry, 69*,  
721 1019-1025.
- 722 Oakley, D. A., & Halligan, P. W. (2017). Chasing the rainbow: the non-conscious nature of  
723 being. *Frontiers in Psychology, 8*, 1924.
- 724 Pedersen, M. L., & Frank, M. J. (2020). Simultaneous hierarchical Bayesian parameter estimation  
725 for reinforcement learning and drift diffusion models: A tutorial and links to neural data.  
726 *Computational Brain & Behavior, 3*, 458-471.
- 727 Pedersen, M. L., Frank, M. J., & Biele, G. (2017). The drift diffusion model as the choice rule in  
728 reinforcement learning. *Psychonomic Bulletin & Review, 24*, 1234-1251.

- 729 Ratcliff, R., & Tuerlinckx, F. (2002). Estimating parameters of the diffusion model: Approaches  
730 dealing with contaminant reaction times and parameter variability. *Psychonomic Bulletin &*  
731 *Review*, 9, 438-481.
- 732 Ratcliff, R., Smith, P. L., Brown, S. D., & McKoon, G. (2016). Diffusion decision model: Current  
733 issues and history. *Trends in Cognitive Sciences*, 20, 260-281.
- 734 Rescorla, R. A., & Wagner, A. R. (1972). A theory of Pavlovian conditioning: Variations in the  
735 effectiveness of reinforcement and nonreinforcement. In A. H. Black & W. F. Prokasy  
736 (Eds.), *Classical conditioning II: Current research and theory* (pp. 64-99). Appleton  
737 Century Crofts.
- 738 Rogers, T. B., Kuiper, N. A., & Kirker, W. S. (1977). Self-reference and the encoding of personal  
739 information. *Journal of Personality and Social Psychology*, 35, 677-688.
- 740 Schäfer, S., Wentura, D., & Frings, C. (2015). Self-prioritization beyond perception. *Experimental*  
741 *Psychology*, 62, 415-425.
- 742 Schäfer, S., Frings, C., & Wentura, D. (2016). About the composition of self-relevance:  
743 Conjunctions not features are bound to the self. *Psychonomic Bulletin and Review*, 23, 887-  
744 982.
- 745 Schäfer, S., Wentura, D., & Frings, C. (2017). Distinctiveness effects in self-prioritization. *Visual*  
746 *Cognition*, 25, 1-13.
- 747 Schultz, W., Dayan, P., & Montague, P. R. (1997). A neural substrate of prediction and  
748 reward. *Science*, 275, 1593-1599.
- 749 Schultz, W. (1998). Predictive reward signal of dopamine neurons. *Journal of Neurophysiology*, 80,  
750 1-27.
- 751 Sedikides, C., & Alicke, M. D. (2012). Self-enhancement and self-protection motives. In R. M.  
752 Ryan (Ed.), *The Oxford handbook of human motivation* (pp. 303-322). Oxford University  
753 Press.

- 754 Shapiro, K. L., Caldwell, J., & Sorensen, R. E. (1997). Personal names and the attentional blink: A  
755 visual “cocktail party” effect. *Journal of Experimental Psychology: Human Perception and*  
756 *Performance*, 23, 504-514.
- 757 Siebold, A., Weaver, M. D., Donk, M., & van Zoest, W. (2015). Social salience does not transfer to  
758 oculomotor visual search. *Visual Cognition*, 23, 989-1019.
- 759 Spiegelhalter, D. J., Best, N. G., Carlin, B. P., & van der Linde, A. (1998). *Bayesian deviance, the*  
760 *effective number of parameters, and the comparison of arbitrarily complex models.*  
761 Research Report, 98-1009.
- 762 Spiegelhalter, D. J., Best, N. G., Carlin, B. P., & van der Linde, A. (2002). Bayesian measures of  
763 model complexity and fit. *Journal of the Royal Statistical Society: Series B*, 64, 583-639.
- 764 Stein, T., Siebold, A., & van Zoest, W. (2016). Testing the idea of privileged awareness of self-  
765 relevant information. *Journal of Experimental Psychology: Human Perception and*  
766 *Performance*, 42, 303-307.
- 767 Stolte, M., Humphreys, G., Yankouskaya, A., & Sui, J. (2017). Dissociating biases towards the self  
768 and positive emotion. *Quarterly Journal of Experimental Psychology*, 70, 1011-1022.
- 769 Sui, J., & Han, S. (2007). Self-construal priming modulates neural substrates of self-awareness.  
770 *Psychological Science*, 18, 861-866.
- 771 Sui, J., He, X., & Humphreys, G. W. (2012). Perceptual effects of social salience: Evidence from  
772 self-prioritization effects on perceptual matching. *Journal of Experimental Psychology:*  
773 *Human Perception and Performance*, 38, 1105-1117.
- 774 Sui, J., & Humphreys, G. W. (2015). The integrative self: How self-reference integrates perception  
775 and memory. *Trends in Cognitive Sciences*, 19, 719-728.
- 776 Sui, J., & Humphreys, G. W. (2017). The ubiquitous self: What the properties of self-bias tell us  
777 about the self. *Annals of the New York Academy of Sciences*, 1396, 222-235.



- 778 Sui, J., Liu, M., Mevorach, C., & Humphreys, G. W. (2015). The salient self: The left intra-parietal  
779 sulcus responds to social as well as perceptual salience after self-association. *Cerebral*  
780 *Cortex*, 25, 1060-1068.
- 781 Sui, J., & Rotshtein, P. (2019). Self-prioritization and the attentional systems. *Current Opinion in*  
782 *Psychology*, 29, 148-152.
- 783 Sui, J., Rotshtein, P., & Humphreys, G. W. (2013). Coupling social attention to the self forms a  
784 network for personal significance. *Proceedings of the National Academy of Sciences*, 110,  
785 7607-7612.
- 786 Sutton, R. S., & Barto, A. G. (1998). *Reinforcement learning: An introduction* (Vol. 135).  
787 Cambridge: MIT press.
- 788 Svensson, S. L., Golubickis, M., Maclean, H., Falbén, J. K., Persson, L. M., Tsamadi, D., ... &  
789 Macrae, C. N. (2021). More or less of me and you: Self-relevance augments the effects of  
790 item probability on stimulus prioritization. *Psychological Research*, 10.1007/s00426-021-  
791 01562-x. Advance online publication.
- 792 Symons, C. S., & Johnson, B. T. (1997). The self-reference effect in memory: A meta-analysis.  
793 *Psychological Bulletin*, 121, 371-394.
- 794 Wang, H., Humphreys, G., & Sui, J. (2016). Expanding and retracting from the self: Gains and  
795 costs in switching self-associations. *Journal of Experimental Psychology: Human*  
796 *Perception and Performance*, 42, 247-256.
- 797 Watkins, C., & Dayan, P. (1992). Q-learning. *Machine Learning*, 8, 279-292.
- 798 Wiecki, T. V., Sofer, I., & Frank, M. J. (2013). HDDM: Hierarchical Bayesian estimation of the  
799 drift-diffusion model in Python. *Frontiers in Neuroinformatics*, 7, 1-10.
- 800 Woźniak, M., & Knoblich, G. (2019). Self-prioritization of fully unfamiliar stimuli. *Quarterly*  
801 *Journal of Experimental Psychology*, 72, 2110-2120.
- 802 Woźniak, M., & Knoblich, G. (2021). Self-prioritization depends on assumed task-relevance of  
803 self-association. *Psychological Research*. 10.1007/s00426-021-01584-5. Advance online  
804 publication.

- 805 Yin, S., Sui, J., Chiu Y-C, Chen, A., & Egner, T. (2019). Automatic prioritisation of self-referential  
806 stimuli in working memory. *Psychological Science*, *30*, 415-423.
- 807 Zhu, Y., Zhang, L., Fan, J., & Han, S. (2007). Neural basis of cultural influence on self-  
808 representation. *Neuroimage*, *34*, 1310-1316.
- 809