Dynamic Representation of the Subjective Value of Information

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Abstract

To improve future decisions, people should seek information based on the value of information (*VOI*), which depends on the current evidence and the reward structure of the upcoming decision. When additional evidence is supplied, people should update *VOI* to adjust subsequent information seeking, but the neurocognitive mechanisms of this updating process remain unknown. We used a modified beads task to examine how the *VOI* is represented and updated in the human brain. We theoretically derived, and empirically verified, a normative prediction that the *VOI* depends on decision evidence and is biased by reward asymmetry. Using fMRI, we found that the subjective *VOI* is represented in right dorsolateral prefrontal cortex (DLPFC). Critically, this *VOI* representation was updated when additional evidence was supplied, showing that DLPFC dynamically tracks the up-to-date *VOI* over time. These results provide new insights into how humans adaptively seek information in the service of decision making.

1 Introduction

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Information seeking is critical for adaptive decision making. In order to improve future 3 decisions, we collect information that would help us predict their outcomes. For 4 instance, we check the weather forecast to decide whether to go out for a hike; we read 5 6 about the policies and characters of candidates to decide how to vote; and we look up 7 the number of COVID-19 cases to decide whether to have a family gathering. Recent work raises the possibility that deficits in information seeking underlie some psychiatric 8 diseases such as schizophrenia and obsessive-compulsive disorder (OCD) (Baker et 9 al., 2019; Dudley et al., 2016; Hauser et al., 2017; Ross et al., 2015). 10 11 In economic theories, information seeking should be primarily driven by information's 12 13 instrumentality, or how much the information would help the agent acquire rewards and avoid punishments in an upcoming decision. The information's instrumentality is 14 formally characterized as the value of information (VOI), defined as the improvement in 15

the expected value (EV) that the agent can achieve by making the decision based on 16 17 the information (Edwards, 1965; Howard, 1966). While this normative VOI theory does not incorporate psychological motives of curiosity, such as anticipatory utility (Caplin & 18 19 Leahy, 2001; Gottlieb & Oudeyer, 2018; Kakade & Dayan, 2002; Kidd & Hayden, 2015; 20 Kobayashi et al., 2019; Kreps & Porteus, 1978; Sharot & Sunstein, 2020), it predicts human participants' information-seeking decisions reasonably well in settings where 21 22 they acquire information at a cost (such as monetary costs or opportunity costs) that can be used to maximize rewards (Edwards & Slovic, 1965; Kobayashi & Hsu, 2019; 23 Shanteau & Anderson, 1972; Wendt, 1969; Wilson et al., 2014). The idea that 24 25 information seeking is driven by the VOI is further supported by electrophysiological and

- neuroimaging evidence that the *VOI* is encoded in reward-related regions (e.g., nucleus
 accumbens, ventromedial prefrontal cortex) as well as anterior cingulate cortex (ACC)
 and dorsolateral prefrontal cortex (DLPFC) (Blanchard et al., 2015; Bromberg-Martin &
 Hikosaka, 2009, 2011; Brydevall et al., 2018; Charpentier et al., 2018; Gruber et al.,
- 30 2014; Jepma et al., 2012; Kaanders et al., 2020; Kang et al., 2009; Kobayashi & Hsu,
- 2019; Krebs et al., 2009; Lau et al., 2020; White et al., 2019).
- 32

The notion of the VOI based on the information's instrumentality has two important 33 implications. First, the VOI should not be determined by how much the information 34 would contribute to the accuracy of prediction on the state of the world, but rather how 35 much it would help the agent maximize *rewards*. Therefore, the VOI depends on the 36 upcoming decision's reward structure, or how rewarding or punishing possible 37 38 outcomes are (for instance, the value of a weather forecast depends on how much the hiker prefers different weather conditions; those who don't mind hiking in the rain or 39 snow may not value the weather forecast as much as those who do). Second, the VOI 40 depends on decision evidence that the agent already possesses prior to information 41 seeking (Loewenstein, 1994). The VOI tends to be smaller when the agent already has 42 more evidence, because they may already know what to do and additional information 43 is less likely to influence it (e.g., a hiker may not need to check the weather forecast if 44

they have been already informed by other hikers that it is going to snow). Thus, the 45

agent needs to combine the available decision evidence with the reward structure to 46 47 assess the VOI and seek information adaptively.

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Crucially, when the decision evidence available to the agent changes, the agent should 49 dynamically update the VOI based on the most recent evidence. Situations requiring 50 such updating are ubiguitous in the real world, either because the environment 51 52 gradually supplies evidence over time (e.g., a recent weather forecast is more accurate 53 than an old one) or because the agent sequentially samples multiple pieces of information (the hiker can check multiple sources of weather forecasts). Despite its 54 importance, to the best of our knowledge, no study has examined how the human brain 55 tracks the up-to-date VOI based on the most recent decision evidence. The majority of 56 neuroimaging studies so far have focused on cases where information is not 57 instrumental for upcoming decisions, and those that have examined instrumentality-58 59 driven information seeking did not experimentally manipulate decision evidence over 60 time to characterize the neural processes of updating the VOI (Kaanders et al., 2020; Kobayashi & Hsu, 2019). 61 62 We conducted an fMRI study to examine how human information-seeking behavior is 63 sensitive to reward structure and current decision evidence, and how human brains 64 65 track the up-to-date VOI after acquiring additional evidence. Our contributions are threefold. First, we theoretically derive, and empirically demonstrate, a simple and 66 generalizable prediction for how information seeking should be biased by asymmetry in 67 reward structure. Second, we show that the right DLPFC represents the subjective VOI 68

as a function of asymmetric rewards and current evidence. Third, we show that the VOI 69

representation in the right DLPFC is dynamically updated when a new piece of evidence 70 is supplied. These results suggest that the right DLPFC plays a critical role in 71

72 information seeking in dynamic decision-making contexts by tracking the up-to-date VOI

73 over time.

Results 74

75

Experimental paradigm 76

To examine neural representations of the value of information (VOI) and its updating, 77 we adopted a variant of the beads task, an experimental paradigm widely used to study 78 79 probability judgement and information seeking (Furl & Averbeck, 2011; Hug et al., 1988; Phillips & Edwards, 1966). As in the conventional version of the beads task, participants 80 were presented with a jar containing two types of beads, one marked with a face and 81 the other marked with a house, and asked to make a bet on its bead composition by 82 observing some beads drawn from it. There were two possible compositions of the jar: 83 one that consists of 60% face beads and 40% house beads, and the other that consists 84 of 40% face beads and 60% house beads (Fig. 1A). 85

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Our variant of the beads task had three key features. First, we introduced reward 87

asymmetry, such that participants could earn more reward by correctly betting on one 88

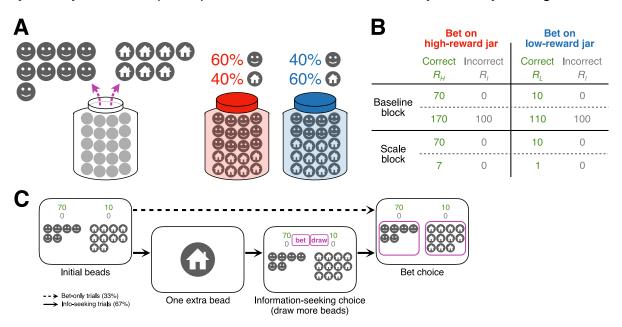


Fig. 1. Experimental paradigm. We adopted the beads task with three key modifications: asymmetry in the reward structure, initial evidence prior to information seeking, and an updating event (one extra bead). (A) Participants observed a number of beads drawn from a jar and made a bet on its composition. Each bead was marked with a face or a house. There were two possible jar compositions: 60% face beads and 40% of house beads, or 40% face beads and 60% house beads. The jars are colored here only for illustrative purposes. (B) Reward structure. Participants earned more reward points by correctly betting on one of the two jar types. The experiment consisted of two blocks, in each of which one of the two reward structures was presented in each trial. The first block involved a baseline shift and the second block involved a scale manipulation. (C) Trial sequence. In a third of the trials (bet-only trials), participants were presented with a number of beads from the jar and immediately made a bet on its type. In the remaining trials (information-seeking trials), they were presented with the initial beads, an extra bead, and then allowed to seek further information by drawing more beads from the jar before making a bet on one of the two jars. Participants could draw as many beads as they needed within five seconds, but each additional draw incurred a cost (0.1 points). The extra bead was presented to evoke updating in the value of information.

jar type (e.g., the face-majority jar) than the other (e.g., the house-majority jar) (Fig. 1B). 89 If participants were motivated to seek information to maximize rewards in the bet, their 90 91 information-seeking strategy should be sensitive not only to the current evidence (the numbers of observed beads so far) but also to the reward asymmetry (the jar type they 92 should bet on to maximize rewards). On the other hand, if participants were motivated 93 to accurately guess the jar type, their information seeking should not be sensitive to the 94 reward asymmetry. Therefore, the reward asymmetry allowed us to test whether 95 96 information seeking was driven by the instrumentality of information for future reward 97 seeking, as normatively prescribed in economic theories. 98 Second, we provided initial evidence, in the form of 20 or 30 bead draws from the jar. 99 On a subset of trials, participants could then seek more information about the jar 100 composition by drawing additional beads or elect to make a bet on the jar type (Fig. 1C). 101 The difference in the numbers of face beads and house beads was parametrically 102 manipulated to range from strong evidence favoring the low-reward jar to strong 103 evidence favoring the high-reward jar. Additional draws incurred a small constant cost 104 (0.1 points per draw) to monetarily incentivize participants to seek information only 105 when necessary. This design allowed us to empirically measure the subjective VOI, or 106

- 107 how much participants were willing to seek costly information, as a function of the
- 108 current evidence.
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Third, on the trials that allowed for information seeking, participants were presented with
one extra bead draw from the jar prior to the information-seeking phase (Fig. 1C). The
extra bead complemented the initial beads, shifting the evidence on the jar
compositions, and thus updated the *VOI* originally evaluated based on the initial beads.
We analyzed neural responses upon this extra bead event to examine how the neural
representation of the *VOI* is dynamically updated based on the up-to-date evidence over
time.

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Participants completed the task inside the scanner. In each trial, after the presentation 118 of initial beads and an extra bead, participants were allowed to draw as many additional 119 beads as they wanted within five seconds, and then made a binary bet on the jar type. 120 Additionally, to empirically elucidate participants' reward-seeking behavior in a way that 121 122 is not contaminated by information seeking, participants were asked to make a bet on the jar type without information seeking in a subset of trials (bet-only trials). Lastly, to 123 explore how information seeking is sensitive to rewards, we introduced trial-wise 124 manipulation of the reward structures. Specifically, participants earned a baseline 125 reward of 100 points, irrespective of their bet, in half of the trials in one block 126 (henceforth the baseline block), and they earned a tenth of the rewards in half of the 127 trials in the other block (henceforth the scale block). Importantly, the reward of a correct 128 bet was asymmetric across all trials and blocks (Fig. 1B). 129 130

131 Theory

132 We first derived a theoretical prediction on how agents should seek information to

- optimize their bet and maximize rewards. We obtained theoretical *VOI* under the
- assumption that the agent aims to maximize the expected value (EV) of their decision,
- 135 which they evaluate based on posterior probability of the jar type inferred in a perfectly
- 136 Bayesian manner.
- 137
- 138 The posterior of the jar type is determined by the numbers of high-reward beads (the
- majority bead in the high-reward jar, e.g., face) and low-reward beads (the majority
- bead in the low-reward jar, e.g., house) observed from the jar so far (Fig. 2A). The more

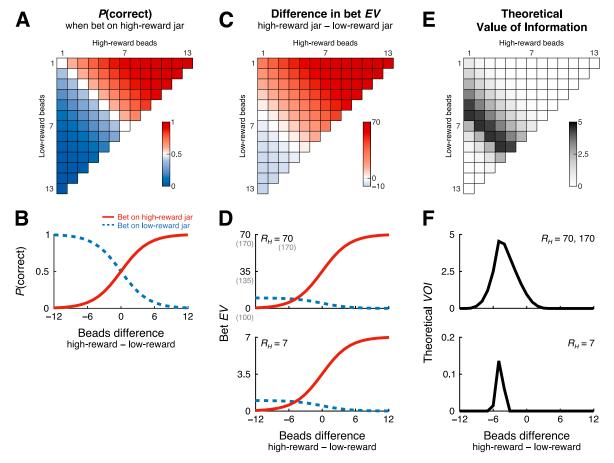


Fig. 2. Theoretical predictions. (A) The probability of the jar type (the true jar is the high-reward jar) increases with the number of observed high-reward beads and decreases with the number of observed low-reward beads. (B) The probability of the jar type is determined by the beads difference. (C) Due to the reward asymmetry, when equal numbers of high-reward and low-reward beads have been observed (the diagonal), the *EV* to bet on the high-reward jar is higher than the low-reward jar. The agent would experience the smallest *EV* difference, and hence the highest uncertainty on the bet, when more low-reward beads have been drawn (the white region). (D) The *EV* difference is smallest at the beads difference of -5 across all reward structures. *Top*: Bet *EV*s are not affected by a baseline shift in rewards. *Bottom*: The relative magnitudes of *EV*s remain the same when rewards are scaled down overall. (E) The theoretical *VOI* is highest when the uncertainty on the bet is highest (the beads difference = -5, the black region). This is because the next bead would provide evidence in favor of either jar type, resolving the uncertainty. (F) The theoretical *VOI* takes an inverted-*U* shape across all reward structures. *Top*: The *VOI* is unaffected by a baseline shift in rewards. *Bottom*: When the rewards are scaled down, the magnitude of *VOI* becomes smaller as well, but the peak location remains the same. *EV*: expected value, *VOI*: value of information.

high-reward beads have been drawn, the more likely the jar is the high-reward jar, and
vice versa. More specifically, the posterior is determined by the difference in the
numbers of observed beads (high-reward beads minus low-reward beads) (Fig. 2B; Eq.
1). When more high-reward beads have been observed than low-reward beads (the
beads difference > 0), the probability of the high-reward jar is higher than the probability
of the low-reward jar, and it increases with the beads difference. Conversely, when
more low-reward beads have been observed (the beads difference < 0), evidence

- 148 favors the low-reward jar.
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In order to evaluate the EV of a bet, the agent needs to combine the posterior on the jar 150 type with the reward structure (Fig. 2C). Due to the reward asymmetry, when the current 151 evidence does not favor either jar (the beads difference = 0; the diagonal in Fig. 2C), the 152 153 EV to bet on the high-reward jar is higher than the EV to bet on the low-reward jar. The EVs to bet on the two jars are closest to each other when more low-reward beads have 154 155 been observed (the beads difference = -5; the white region in Fig. 2C). This prediction holds across all of our reward structures (Fig. 2D); a baseline shift in rewards does not 156 affect the EV difference, and a scale manipulation in rewards multiplicatively affect both 157 EVs without changing their relative magnitudes. Therefore, if forced to bet on one of the 158 two possible jars, the EV-maximizing agent would experience the highest choice 159 uncertainty, not when equal numbers of beads have been observed, but when more 160 low-reward beads have been observed than high-reward beads. 161

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Under economic theories, the VOI, or the value of drawing an additional bead, is 163 evaluated based on how much the next bead would improve the upcoming bet on 164 average (Eq. 2). Qualitatively, the theoretical VOI tends to increase with the uncertainty 165 about which jar type to bet on, because an additional bead would provide more 166 evidence for either jar type and resolve the uncertainty over possible actions (Fig. 2E). 167 For instance, when the agent is under high uncertainty on the bet (the beads difference 168 = -5; the black region in Fig. 2E), an additional bead would help them make a bet 169 *irrespective of its type*; if the next bead is a high-reward bead, it provides additional 170 evidence in favor of the high-reward jar, whereas if it is a low-reward bead, it favors the 171 low-reward jar. The agent can improve the EV by making a bet conditional on the next 172 bead type. On the other hand, when the agent has observed more high-reward beads 173 174 than low-reward beads (e.g., the beads difference = +10; top right in Fig. 2E), or when the agent has observed many more low-reward than high-reward beads (e.g., the beads 175 difference = -10; bottom left in Fig. 2E), an additional bead would not affect the 176 subsequent bet; the agent would bet on the high-reward jar or low-reward jar, 177 respectively, no matter what the next bead would be. Therefore, the theoretical VOI 178 takes an inverted-U shape as a function of the beads difference, with its peak at a 179 negative beads difference (-5) (Fig. 2F). 180

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182 Therefore, our theoretical framework yields an important prediction that the information-

183 seeking strategy should be biased by the reward asymmetry; participants should draw 184 additional beads more frequently when more low-reward beads have been observed

than high-reward beads (the beads difference < 0). The predicted bias holds across 185 reward structures (Fig. 2F); manipulation of the reward baseline (in the baseline block) 186 187 does not affect the VOI, and manipulation of the reward scaling (in the scale block) affects the overall magnitude of the VOI but does not drastically alter its inverted-U 188 shape. This prediction might be somewhat counterintuitive, as the motivation for 189 information seeking is expected to be higher when the current evidence favors the less 190 desirable state (the low-reward jar). However, it is consistent with the widespread notion 191 192 of confirmation bias that an agent needs less evidence to bet on a desirable state than an undesirable state (e.g., Gesiarz et al., 2019). More generally, the prediction echoes 193 the general assumption that information seeking should be driven not by the motivation 194 to predict the state (which jar is the true jar?) but to maximize rewards (which jar to bet 195 on?). If, in contrast to our theoretical assumption, an agent is solely motivated to 196 accurately predict the state, they would seek information the most when the beads 197 difference is zero. Therefore, a bias in information seeking would suggest that 198 participants seek information based on its instrumentality for future reward seeking, as 199 200 normatively prescribed. To our knowledge, the bias in information seeking under the reward asymmetry is a novel theoretical prediction that has not yet been directly tested. 201 202

203 Behavior

We examined participants' information-seeking behavior, and in particular, whether it was biased due to the reward asymmetry as predicted. If participants sought to improve their subsequent bet choice and maximize rewards, the frequency of information seeking (i.e., how often they drew at least one bead) should be biased towards a negative beads difference, i.e., when more low-reward beads have been drawn than high-reward beads.

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Observed information-seeking behavior was biased in the predicted direction (Fig. 3A). 211 In both baseline and scale blocks, the frequency of drawing an additional draw was 212 highest when more low-reward beads had been drawn than high-reward beads. 213 Sensitivity to the reward asymmetry was also confirmed by the bet on the jar type in the 214 bet-only trials (Fig. 3B); the frequency of betting on the high-reward jar increased with 215 the beads difference, and the indifference point (the point at which participants were 216 equally likely to bet on either jar) was shifted towards a negative beads difference. 217 218 These results show that participants incorporated both the current evidence and reward asymmetry in reward-seeking and information-seeking choices. 219

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A notable deviation from the theoretical prediction is that participants' information 221 seeking was not sensitive to the reward scale manipulation. In our framework, the 222 223 theoretical VOI is smaller when the rewards are scaled down (even though its peak location remains the same) while it is unaffected by a reward baseline shift (Fig. 2F). 224 Thus, if our participants were perfectly sensitive to the reward structure on a trial-by-trial 225 226 basis, their information seeking should be affected by trial-wise reward manipulation in the scale block but not in the baseline block. To test this, we examined how information-227 seeking behavior differed across reward conditions and blocks. To characterize the 228

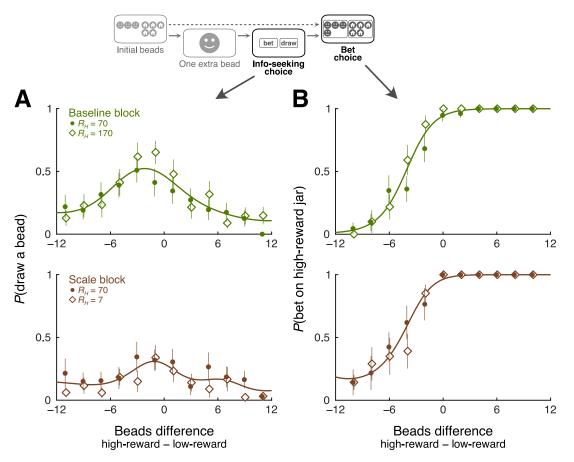


Fig. 3. Behavior. Participants' information-seeking and reward-seeking behavior was biased by the reward asymmetry as predicted. (A) Participants' information seeking, or the frequency at which they drew at least one bead, peaked when more low-reward beads had been drawn than high-reward beads. (B) In the bet-only trials, the frequency with which they bet on the high-reward jar increased with the beads difference and was biased by the reward asymmetry. Lines indicate the best-fit model, which assumed sensitivity to blocks but not to reward manipulations within blocks. Error bars: bootstrap SD resampling participants.

relationship between information seeking and the beads difference without assuming its 229 functional form, we used Gaussian Process (GP) logistic regression (Rasmussen & 230 Williams, 2006). We fit four models to participants' behavior; Model 1 assumed 231 sensitivity to the scale manipulation but not to the baseline manipulation, as normatively 232 prescribed; Model 2 assumed sensitivity to both manipulations; Model 3 assumed a 233 difference between blocks but no sensitivity to manipulation in either block; and Model 4 234 assumed no difference between blocks or reward conditions. We found that Model 3 235 outperformed other models, including Model 1, according to both leave-one-participant-236 out cross validation (LOPO CV; log likelihoods [LL] = -1216.93, -1216.15, -1214.73, 237 and -1232.15) and leave-one-trial-out cross validation (LOTO CV; LL = -1143.61, 238 -1143.76, -1142.25, and -1166.17). Therefore, participant's information-seeking 239 behavior was systematically different between blocks, even though they did not change 240 their strategy based on the reward structure on a trial-by-trial basis. 241 242

We speculate that shifting information-seeking strategies on a trial-by-trial basis was too 243 cognitively taxing for our participants, because we also manipulated the beads 244 245 difference and the trial type (information-seeking or bet-only). Despite this limitation, we observed that participants' information seeking exhibited a clear bias in both blocks. 246 Indeed, we observed that Model 3, which allowed asymmetry in information seeking, 247 performed better than another model (Model 5) that assumed symmetric information 248 seeking (baseline block LOPO CV LL = -666.82 [Model 3] vs. -679.52 [Model 5]; LOTO 249 250 CV LL = -630.87 vs. -645.10; scale block LOPO CV LL = -547.92 vs. -548.13; LOTO 251 CV LL = -511.68 vs. -512.53). Furthermore, analysis on betting choices also preferred Model 3 to Models 1 and 2 (comparison between Models 3 and 4 is equivocal; LOPO 252 CV LL = -287.01, -285.67, -283.56, and -281.19; LOTO CV LL = -267.77, -266.25, 253 -265.26, and -268.46), showing that participants were insensitive to trial-wise reward 254 255 manipulation not only in information seeking but also in reward seeking. These results are gualitatively consistent with our theoretical prediction and lend support to the 256 257 general notion that people seek information to improve their subsequent choices and 258 maximize rewards.

260 Neural representation of VOI

Next, we examined how the VOI was represented in the brain. Although previous fMRI 261 studies reported VOI representations in a set of regions including DLPFC. VMPFC, and 262 263 striatum, most of these studies focused on situations where participants obtained information that would not be useful for future decisions (i.e., information seeking for its 264 own sake), and one study that examined instrumentality-driven information seeking 265 used a one-shot paradigm that did not involve any updating (Kobayashi & Hsu, 2019). 266 Thus, it remains unknown to what extent the neural representation of VOI is 267 generalizable across tasks and decision contexts, and whether previously reported 268 regions also represent and update the VOI in our experimental paradigm. 269

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To look for brain regions that represent the *VOI*, we empirically estimated subjective *VOI* from the information-seeking behavior. We used the winning model of our GP logistic regression analysis (Model 3) to obtain the latent value function, which varied smoothly with the beads difference and differed between blocks (Fig. 4A). We then looked for regions where neural responses at the presentation of initial beads covaried with the subjective *VOI*.

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We found a cluster in the right DLPFC representing subjective VOI (Fig. 4B; clusterforming threshold p < .001, cluster-mass p < .05, whole-brain FWE corrected; peak MNI coordinate = [48, 42, 24]). Activation in this cluster peaked when more low-reward beads had been drawn in both blocks, consistent with the prediction (Fig. 4C). This cluster is the only one that survived our whole-brain statistical threshold (we also observed a cluster in the right anterior insula at a more lenient threshold, p < .10; peak MNI coordinate = [30, 24, 4]; Fig. S1).

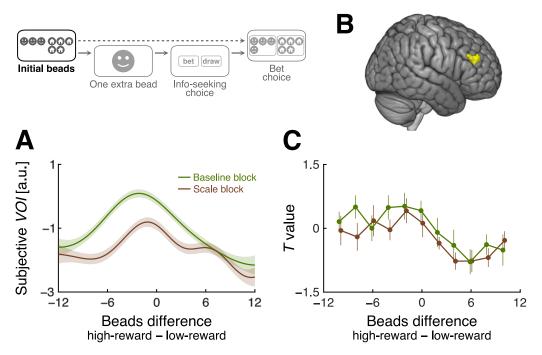


Fig. 4. Neural representation of the *VOI*. (A) The subjective *VOI* was estimated for each block based on information-seeking behavior (Fig. 3A). (B) The right DLPFC represented the subjective *VOI* (cluster-mass p < .05, whole-brain FWE corrected; the peak MNI coordinate: [48, 42, 24]). (C) As predicted, the right DLPFC activation peaks at a negative beads difference in both blocks. Error bars: SEM.

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Interestingly, the DLPFC cluster overlaps with a *VOI* cluster reported in a previous study
that examined one-shot instrumentality-driven information seeking (Kobayashi & Hsu,
2019) (Fig. S1), providing converging evidence that the right DLPFC represents the *VOI*across decision contexts, at least when information is primarily acquired based on its
instrumentality for future value-guided decisions.

291

292 Updating of VOI representation

We then turned to our final question: how is the VOI updated upon the arrival of 293 additional evidence in the brain? When the evidence available to agents changes, they 294 need to track the up-to-date VOI in order to seek information adaptively over time. 295 296 Specifically, we examined how the right DLPFC responds to the extra bead presented after the initial beads but prior to the information-seeking choice (Fig. 5A). We derived 297 the VOI updating, or the difference between the posterior and prior VOI, as a function of 298 the difference in the initial beads (the prior evidence) and the type of the extra bead (the 299 evidence that causes updating). For instance, if participants have observed many more 300 low-reward beads than high-reward beads (the beads difference < -5), an extra high-301 reward bead would positively update the VOI, as it slightly increases the uncertainty on 302 the bet, while an extra low-reward bead would negatively update the VOI, as it further 303 decreases the uncertainty on the bet. The directionality of updating is the opposite when 304 more high-reward beads have been observed (the beads difference > 0). 305

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We hypothesized that the right DLPFC tracks the up-to-date VOI over time, such that it 307 responds not only to the VOI based on the initial beads but is dynamically updated to 308 the appropriate updated VOI after observation of the extra bead. To test this, we 309 estimated the effects of the initial VOI and VOI updating on BOLD signals from the 310 region of interest (ROI) defined above (Fig. 4B). In order to avoid a strong assumption 311 about the time course of the updating process, we estimated the effects of initial VOI 312 and VOI updating across time using finite impulse response (FIR) functions aligned to 313 314 the presentation of the extra bead (Fig 5, top). We included three FIRs in a GLM, one parametrically modulated with the initial VOI, one modulated with the VOI updating, and 315 one without parametric modulation (intercept). Since the ROI was originally defined 316

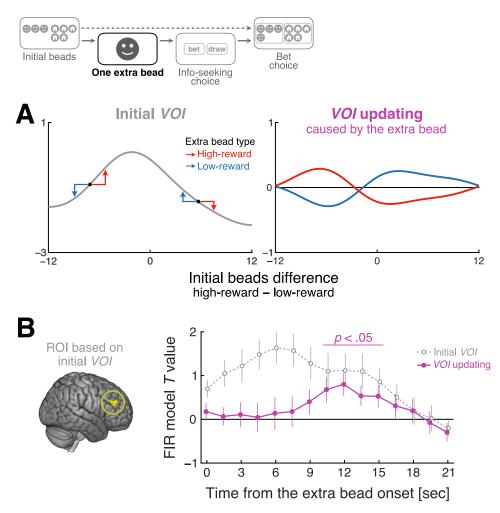


Fig. 5. Updating of the *VOI* representation. The right DLPFC tracks *VOI* as it is updated by an extra bead, presented after the initial beads but prior to information seeking. (A) The *VOI* updating was calculated as the signed difference between the *VOI* after the extra bead and the *VOI* before the extra bead. (B) Time courses of the initial *VOI* signal (grey) and the *VOI* updating signal (purple) in the right DLPFC. The right DLPFC responds not only to the initial *VOI* but also to the updating of *VOI* (temporal cluster-mass p < .05, FWE corrected). Since the region of interest was defined based on the initial *VOI* signal, estimation of the initial *VOI* signal is biased, but estimation of the updating signal is unbiased. Error bars: SEM.

based on its response to the initial *VOI* (albeit in an earlier time window), the estimated
effect of the initial *VOI* is biased, but the estimated effect of the *VOI* updating depends
critically on the exact bead that was drawn, and thus is independent of our ROI
selection process (Fig 5A).

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The estimated time courses are shown in Fig. 5B. As expected, the right DLPFC 322 represents the initial VOI early on. Importantly, the right DLFPC also positively 323 324 responded to the VOI updating (cluster-forming threshold p < .05, cluster-mass p < .05, 325 FWE corrected across time). The rise of the VOI updating signal lags behind the initial VOI signal in time, but they go back to the baseline in parallel. The estimated time 326 courses look somewhat sluggish, which presumably reflects the nature of our 327 experimental paradigm in which participants had several seconds to complete 328 329 information seeking.

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331 This evidence demonstrates that neural representations in right DLPFC shift from the

initial (*a priori*) VOI to the updated (*a posteriori*) VOI, suggesting that this brain region

333 dynamically tracks the *VOI* based on the up-to-date evidence in service of adaptive

information seeking over time.

335 Discussion

336

In order to make better decisions, we need to seek information adaptively based on 337 what we already know (up-to-date decision evidence) and what is at stake (reward 338 structure). When our knowledge is updated, we need to update the VOI accordingly to 339 340 decide whether to seek further information. Deficits in updating the VOI could lead to 341 excessive repetition of information seeking even after enough evidence is accumulated (Hauser et al., 2017), or conversely, premature jumping to conclusions without enough 342 evidence (Dudley et al., 2016; Ross et al., 2015). Despite its importance and ubiquity in 343 the real world, we know little about how people evaluate and update the VOI. In this 344 study, we used a variant of the beads task, in which decision evidence was 345 parametrically manipulated on a trial-by-trial basis, to examine how information seeking 346 is shaped by current evidence and asymmetric reward structure, and how the VOI is 347 represented and updated in the brain. 348

349 We theoretically derived, and empirically verified, the normative prediction that 350 information seeking should be biased by reward asymmetry. Participants were more 351 likely to seek information when the current evidence preferred the less rewarding state 352 353 due to high uncertainty on which state to bet. While the current study used asymmetric 354 monetary rewards, our theoretical framework can be generalized beyond economic decision making based on the notion that the people assign intrinsic values to beliefs 355 that they can hold (Kunda, 1991; Sharot & Garrett, 2016). If people are incentivized to 356 hold certain beliefs, they will be more motivated to seek information when the current 357 evidence supports the less desirable belief, even without extrinsic reward asymmetry 358 (e.g., people check the latest number of COVID-19 cases more often when it is 359 360 increasing than decreasing). It is worth noting, however, that the current study only examined reward structures where a correct bet yields asymmetric rewards but an 361 incorrect bet does not, while outcomes of an incorrect prediction could also be 362 asymmetric in some real-world scenarios (e.g., it would be more punishing to 363 364 underestimate the chance of COVID-19 transmission and end up causing a superspreader event than to overestimate it and avoid a social gathering). More 365 366 comprehensive, generalizable predictions would be obtained by expanding our findings to various reward structures. 367

368

Our theoretical and behavioral findings may provide some insight into confirmation 369 biases observed across domains. Confirmation bias is commonly framed as biases in 370 updating processes and/or decision criteria due to reward asymmetry or other factors 371 372 such as pre-commitment (Gesiarz et al., 2019; Leong et al., 2019; Luu & Stocker, 2018; Talluri et al., 2018). We showed that, even without biases in updating or decision 373 criteria, information seeking should be biased by reward asymmetry. The current study 374 was not designed to test conventional confirmation bias; our behavioral measure of 375 information seeking is not sensitive to a bias in updating, and a bias in decision criteria 376 is not distinguishable from non-neutral risk attitude in our paradigm. Future research 377 may examine how confirmation bias in updating and/or decision criteria affects 378

information seeking, and conversely, how the information seeking bias would strengthen
or weaken the effects of confirmation bias. Another exciting question for future research
would be whether people exhibit an information-seeking bias upon sampling evidence
from internal representations rather than the external world, such as episodic memory
(Shadlen & Shohamy, 2016).

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Our finding of the VOI representation in DLPFC is consistent with a previous fMRI study 385 386 on instrumentality-driven information seeking (Kobayashi & Hsu, 2019), despite a number of key differences in task design. First, our paradigm required probabilistic 387 inference on the hidden jar composition based on observable evidence, while 388 Kobayashi & Hsu (2019) provided explicit and unambiguous visual presentation of 389 outcome probability. Second, while Kobayashi & Hsu (2019) manipulated the 390 information's diagnosticity and cost on a trial-by-trial basis, the current paradigm did not 391 (the participant always drew one bead at a time, which incurred a small constant cost). 392 393 Third, and most importantly, unlike Kobayashi & Hsu (2019), the current study manipulated decision evidence available to the participant at the beginning of each trial 394 and examined its effect on information-seeking behavior and underlying neural signals. 395 Thus, the current study not only replicates but also critically extends Kobayashi & Hsu 396 (2019)'s findings by showing that DLPFC is sensitive to the current evidence and biased 397 by reward asymmetry, a key theoretical prediction of the instrumentality-driven VOI. 398 399 Along with neuroimaging evidence that DLPFC is also activated upon information seeking driven by factors other than instrumentality (Gruber et al., 2014; Kang et al., 400 2009; Jepma et al., 2012), these results suggest that DLPFC is critical for adaptive 401 information seeking across decision contexts and domains. 402

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Unlike previous studies, we did not find VOI representation in reward regions (e.g., 404 striatum or VMPFC) or ACC (Bromberg-Martin & Hikosaka, 2009, 2011; Brydevall et al, 405 2018; Charpentier et al., 2018; Gruber et al., 2014; Kaanders et al., 2020; Kang et al., 406 407 2009; Krebs et al., 2009; Lau et al., 2020; White et al., 2019). It is possible that we lacked statistical power to detect signals in these regions; indeed, we found a VOI 408 cluster in anterior insula at a liberal threshold (Fig. S1), which often coactivates with 409 ACC in task-based and resting-state fMRI (Fox et al., 2005; Menon & Uddin, 2010; 410 Seeley et al., 2007). Alternatively, the involvement of these regions could depend on 411 412 task and decision context. For instance, striatum and/or VMPFC may be more important when the information-seeking cost is larger and variable, which would demand online 413 cost-benefit analysis (Lau et al., 2020; Kobayashi & Hsu, 2019). On the other hand, 414 ACC may be more involved in evaluating uncertainty or conflict in the action space 415 (Kennerley et al., 2011; Rudebeck et al., 2008; Rushworth & Behrens, 2008; Shenhav 416 et al., 2016), which is tightly coupled with the VOI in many cases, particularly in 417 situations that involve an exploration-exploitation tradeoff (Kaanders et al., 2020; Kolling 418 et al., 2012; Shenhav et al., 2014). One possible reason that we did not observe 419 representation of the VOI in ACC, at least at the standard statistical threshold we used, 420 is that our experimental paradigm decoupled action uncertainty from the VOI 421 computation in three ways: first, information-seeking trials were intermixed with bet-only 422

trials and the participant could not tell the trial type upon the presentation of the initial 423 evidence (the epoch where we observed the VOI representation; Fig. 4); second, the 424 425 action uncertainty could not be evaluated until the presentation of the extra bead; and third, the information-seeking decision was mapped to different actions (left vs. right) 426 across trials. Further research is needed to understand the extent to which functional 427 localization of the VOI is dependent on task and decision context, and furthermore, how 428 neural representation of the VOI is related to other forms of information seeking. 429 430 including exploration and curiosity.

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Importantly, we showed that DLPFC not only represents the VOI based on the initial 432 evidence but also updates it when additional evidence is supplied, or in other words, 433 DLPFC tracks the up-to-date VOI based on the most recent evidence. Such DLPFC 434 signals may be critical for adaptive information seeking in situations where the agent 435 accumulates decision evidence over time, either because it is gradually supplied from 436 the environment or because the agent sequentially acquires multiple pieces of 437 information. DLPFC may be well suited for sustained and dynamically updated 438 representation of the VOI, as DLPFC neurons are known to exhibit sustained activity for 439 working memory retention (Funahashi et al., 1989; Fuster & Alexander, 1971; 440 Sreenivasan & D'Esposito, 2019). Critically, the VOI updating in DLPFC is distinct from 441 information prediction error (IPE) signals observed in the dopaminergic reward system 442 and habenula (Blanchard et al., 2015; Bromberg-Martin & Hikosaka, 2009, 2011; 443 Charpantier et al., 2018); IPE encodes the probabilistic delivery of information itself, 444 while the VOI updating is concerned with how the delivered information increases or 445 decreases the instrumentality of further information. Exciting open questions for future 446 research include whether VOI signals in DLPFC play a causal role in information-447 seeking behavior, and how they are adjusted when evidence acquired in the past 448 becomes less relevant in a dynamic environment (Behrens et al., 2007; McGuire et al., 449 450 2014; Nassar et al., 2019).

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Our results may have important implications for information-seeking deficits in clinical 452 populations. For instance, schizophrenia has been associated with the tendency to 453 make premature decisions without enough information seeking (Dudley et al., 2016; 454 Ross et al., 2015; but see Baker et al., 2019), which could be accompanied by DLPFC 455 456 hypoactivity (i.e., too low VOI signals) (Barch & Ceaser, 2012) and/or the lack of DLPFC's sensitivity to current decision evidence and reward asymmetry. Similarly, OCD 457 patients exhibit excessive information seeking (Hauser et al., 2017), which could be 458 caused by hyperactivity in DLPFC (i.e., too high VOI signals) (Eng et al., 2015) and/or 459 the lack of VOI updating in DLPFC. Our experimental and theoretical framework 460 provides a novel approach to characterization of key components in instrumentality-461 driven information seeking, namely the sensitivity to current decision evidence, updating 462 caused by additional evidence, and a bias due to reward asymmetry, which can be 463 readily applied in future research with typical and clinical populations. 464

465 Materials and Methods

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All procedures were approved by the Institutional Review Board at the University ofPennsylvania.

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470 Participants 15 people (11 female, 4 male, age: 18-28, mean = 21.27, standard
471 deviation = 2.79) participated in the experiment. They provided informed consent in
472 accordance with the Declaration of Helsinki.

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Task design We adopted a variant of the beads task (Furl & Averbeck, 2011; Hug et 474 al., 1988; Phillips & Edwards, 1966); the participant was presented with a jar containing 475 two types of beads and asked to guess its composition (i.e., which type made up the 476 majority of the beads) by drawing some beads from the jar (Fig. 1A). Our variant had 477 three important features. First, the participant was rewarded for identifying the correct 478 jar composition, but the reward structure was asymmetric, such that the participant 479 could earn more rewards by correctly betting on one jar type than the other (Fig. 1B). 480 Second, a variable number of beads was drawn from the jar and presented to the 481 participant at the beginning of each trial, empirically manipulating the evidence available 482 to the participant before they seek information. Third, an extra bead was presented on a 483 484 subset of trials to update the initial evidence. These features allowed us to examine how the brain represents and updates the VOI based on evidence that changes over time. 485 486

The experiment consisted of two interleaved trial types, bet-only trials and information-487 seeking trials (Fig. 1C). In the bet-only trials, the participant was first presented with a 488 number of beads drawn from the jar. Each bead was depicted as a rounded picture of a 489 490 face or a house (one picture for face or house each was used throughout the experiment). Beads marked with a face were presented to the left and those marked 491 with a house to the right. The participant was told that these beads were drawn from 492 one of two jars: a face-majority jar, which consisted of 60% face beads and 40% house 493 beads, and a house-majority jar, which consisted of 60% house beads and 40% face 494 beads. Rewards for correct and incorrect bets (in points) were also presented, in green 495 496 and gray, respectively. Rewards for a bet on the face-majority jar were shown above the face beads, and rewards for a bet on the house-majority jar above the house beads. 497 Rewards for a correct bet on one jar were numerically larger than rewards for a correct 498 bet on the other jar (reward asymmetry), while an incorrect bet on either jar yielded the 499 same rewards (Fig. 1B). After the presentation of the initial beads for 3 seconds, the 500 participant was asked to make a bet. During the bet phase of the task, face and house 501 502 beads were separately outlined by magenta boxes, and the participant could press the left or right button on a response box to bet on the face- or house-majority jar, 503 respectively. Trials in which the participant did not make a bet within 3 seconds were 504 terminated and discarded from the analysis. 505

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507 In the information-seeking trials, the participant was first presented with the initial beads 508 screen (same as the bet-only trials), followed by a blank screen (0-2 seconds). Next, an

extra bead drawn from the jar was presented, either marked with a face or a house (1 509 second), which was added to the corresponding group of beads on the initial screen (0-510 511 2 seconds). The participant was then asked to decide whether to draw more beads from the jar before making a bet on its composition (information-seeking phase). Two choices 512 appeared on the screen, "draw" and "bet", and the participant pressed one button to 513 draw one more bead and another button to terminate the information-seeking phase 514 and proceed to the bet (the sides of the options were randomized across trials). The 515 516 participant was allowed to draw as many beads as they wanted within 5 seconds, and a 517 face or house bead was added to the screen every time they pressed the "draw" button. The participant was told that each draw incurred a constant small cost (0.1 points). 518 Once they pressed the "bet" button (or when 5 seconds have passed), they were 519 presented with the bet screen (same as the bet-only trials). 520 521

522 The task was programmed in Matlab (The MathWorks, Natick, MA) using MGL 523 (http://justingardner.net/mgl/) and SnowDots (http://code.google.com/p/snow-dots/) 524 extensions.

Procedure In a separate task session before scanning, participants received extensive training on the task, in which various aspects of the task were gradually introduced (betting on the jar composition, asymmetric rewards, costly draws, and multiple reward structures). During the subsequent session, participants completed the task inside the scanner. Participants made responses using an MRI-compatible button box. They were compensated based on the total points they acquired in the scanning session (500 points = \$1).

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The scanning experiment consisted of two blocks, which differed in reward structure 534 (Fig. 1B). In the first block (the baseline block), one of the two reward structures, 535 536 $(R_H, R_I, R_I) = (70, 10, 0)$ or (170, 110, 100), was randomly presented in each trial, where R_H is the reward for a correct bet on the high-reward jar, R_L is the reward for a 537 correct bet on the low-reward jar, and R_I is the reward for an incorrect bet; thus, the 538 participant earned a baseline reward of 100 points irrespective of their bet in half of the 539 trials. In the second block (the scale block), one of the two reward structures, (R_H, R_L, R_I) 540 = (70, 10, 0) or (7, 1, 0), was randomly presented in each trial; thus, the participant 541 542 earned a tenth of the rewards in half of the trials. Each block consisted of two scanning runs, one where the high-reward jar was the face-majority jar and one where the high-543 reward jar was the house-majority jar; their order was counterbalanced across 544 participants. 545

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On each trial, the participant was presented with 20 or 30 initial beads from the jar. The
difference in the number of initial beads marked with a face or house was uniformly
sampled from a discrete set of values ranging from -10 to 10 in increments of 2.
Unbeknownst to the participant, the true jar type was stochastically determined following
the Bayesian posterior conditional on the initial beads difference (see Eq. 1 below). In
the information-seeking trials, the type of the extra bead presented and of all additional

beads drawn by the participant (face or house) were stochastically determined based on
the hidden jar type. The participant was not provided with feedback on their bet
accuracy or rewards on a trial-by-trial basis. They were however informed of the total
number of points they had accumulated at the end of each run.

Theory Normative predictions about the *VOI*, or how much an optimal agent should pay for information, were derived under assumptions that the agent conducts full-Bayesian inference on the jar type, deterministically makes an optimal choice to maximize the expected value (EV), is risk neutral, and optimally seeks information based on its instrumentality, or how much it would improve the EV of the subsequent bet choice. Our theoretical framework did not consider any additional information-seeking motives, such as curiosity, savoring, dread, or uncertainty reduction.

565 Let s_H be the state where the true jar is the high-reward jar and s_L the state where it is 566 the low-reward jar. Let a_H be the action to bet on s_H and a_L the action to bet on s_L . Let 567 us further refer to the majority beads in the high-reward jar as high-reward beads and 568 the majority beads in the low-reward jar as low-reward beads (for instance, if the high-569 reward jar is the house-majority jar, a house bead is a high-reward bead and a face 570 bead is a low-reward bead; note that the beads were not directly associated with 571 rewards per se). The goal for the agent is to choose between a_H and a_L to maximize EV 572 given the current evidence (i.e., the number of high-reward beads n_H and low-reward 573 beads n_L drawn from the jar so far) and the reward structure (R_H, R_L, R_I) . 574

576 The likelihood of drawing a high-reward bead b_H or a low-reward bead b_L conditional on 577 the jar type is known to the agent:

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$$P(b_H|s_H) = P(b_L|s_L) = q$$

 $P(b_L|s_H) = P(b_H|s_L) = 1 - q$

where q = 0.6. Assuming that the agent has a flat prior on the jar type ($P(s_H) = P(s_L) = 0.5$), the posterior follows

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$$\frac{P(s_H|n_H, n_L)}{P(s_L|n_H, n_L)} = \frac{P(n_H, n_L|s_H)P(s_H)}{P(n_H, n_L|s_L)P(s_L)} = \frac{\binom{n_H + n_L}{n_H}P(b_H|s_H)^{n_H}P(b_L|s_H)^{n_L}}{\binom{n_H + n_L}{n_L}P(b_H|s_L)^{n_H}P(b_L|s_L)^{n_L}} = \left(\frac{q}{1-q}\right)^{n_H - n_L}$$

586

587 therefore

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$$P(s_H|n_H, n_L) = 1 - P(s_L|n_H, n_L) = \frac{\left(\frac{q}{1-q}\right)^{n_H - n_L}}{\left(\frac{q}{1-q}\right)^{n_H - n_L} + 1}$$
(1)

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which is a function of the beads difference, $n_H - n_L$ (e.g., the posterior is the same when $(n_H, n_L) = (5, 2)$ or (15, 12) (Fig. 2a, b).

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Given the posterior, the agent makes a choice among three options: to bet on s_H , to bet on s_L , or to seek information and draw an additional bead from the jar, which incurs a cost c_{draw} (0.1 points). The agent should decide whether to draw an additional bead based on the *VOI*, or the improvement in the bet's *EV* thanks to the next bead:

$$VOI(n_H, n_L) = EV_{draw}(n_H, n_L) - EV_{bet}(n_H, n_L)$$
(2)

where EV_{draw} is the highest EV that the agent could achieve after drawing the next bead (without considering the information-seeking cost), and EV_{bet} is the highest EV that the agent could achieve by making a bet without any further information. The agent should draw a bead if and only if the *VOI* is higher than the drawing cost c_{draw} .

606 EV_{bet} is the higher of the two bet EVs based on the current evidence, namely

$$EV_{\text{bet}}(n_H, n_L) = \max_a EV(a|n_H, n_L)$$

610 where $a \in \{a_H, a_L\}$ and

 $EV(a_{H}|n_{H}, n_{L}) = R_{H} \cdot P(s_{H}|n_{H}, n_{L}) + R_{I} \cdot P(s_{L}|n_{H}, n_{L})$ $EV(a_{L}|n_{H}, n_{L}) = R_{L} \cdot P(s_{L}|n_{H}, n_{L}) + R_{I} \cdot P(s_{H}|n_{H}, n_{L})$

Since the posterior is determined by the beads difference (Eq. 1), the bet *EV*s are also determined by the beads difference.

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In order to evaluate EV_{draw} , we have to take into account two important facets of our information-seeking paradigm: first, the content of information (the type of the next bead, b_H or b_L) is stochastic, and second, the agent can decide whether to draw yet another bead or not after observing the next bead. Therefore, we have to evaluate the likelihood of the next bead type and combine it with the EV of an optimal choice conditional on each bead type. The likelihood of the next bead type based on the current evidence is evaluated according to the posterior on the jar type:

$$P(b_H|n_H, n_L) = P(b_H|s_H)P(s_H|n_H, n_L) + P(b_H|s_L)P(s_L|n_H, n_L)$$

$$P(b_L|n_H, n_L) = P(b_L|s_H)P(s_H|n_H, n_L) + P(b_L|s_L)P(s_L|n_H, n_L)$$

If the next bead is b_H , it would update the evidence from (n_H, n_L) to $(n_H + 1, n_L)$. Then the agent can either make an optimal bet and achieve $EV_{bet}(n_H + 1, n_L)$ or pay the cost to draw another bead and achieve $EV_{draw}(n_H + 1, n_L) - c_{draw}$. Similarly, if the next bead is b_L , it would update the evidence to $(n_H, n_L + 1)$, based on which the agent can either make an optimal bet and achieve $EV_{bet}(n_H, n_L + 1)$ or draw another bead and achieve $EV_{draw}(n_H, n_L + 1) - c_{draw}$. Therefore, the highest EV that the agent can achieve after drawing an additional bead is

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$$EV_{draw}(n_H, n_L) = P(b_H | n_H, n_L) \cdot \max[EV_{bet}(n_H + 1, n_L), EV_{draw}(n_H + 1, n_L) - c_{draw}] + P(b_L | n_H, n_L) \cdot \max[EV_{bet}(n_H, n_L + 1), EV_{draw}(n_H, n_L + 1) - c_{draw}]$$
(3)

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In Eq. 3, $EV_{draw}(n_H, n_L)$ in the left-hand side depends on $EV_{draw}(n_H + 1, n_L)$ and 639 $EV_{draw}(n_H, n_I + 1)$ in the right-hand side due to the aforementioned sequentiality of 640 information seeking. We thus solved Eq. 3 by backward recursion. Specifically, we 641 arbitrarily assumed that the agent cannot draw more than 200 beads, set 642 $EV_{draw}(n_H, n_L) = 0$ where $n_H + n_L = 200$, and used Eq. 3 to obtain $EV_{draw}(n_H, n_L)$ where 643 $n_H + n_L = 199$. We then used Eq. 3 recursively to obtain $EV_{draw}(n_H, n_L)$ for all cases 644 where $0 < n_H + n_L < 200$. Although the obtained $EV_{draw}(n_H, n_L)$ depends on $n_H + n_L$, it 645 reaches an asymptote over the course of recursion quickly (Fig. S2). We substituted the 646 asymptotic EV_{draw} to Eq. 2 and obtained the theoretical VOI as a function of the beads

- 647 asymptotic648 difference.
- 649

The *VOI* obtained for each of the three reward structures, $(R_H, R_L, R_I) = (70, 10, 0)$, (170, 110, 100), and (7, 1, 0), is shown in Fig. 2F. The baseline shift affects both EV_{draw} and EV_{bet} by the same amount, which is canceled out in Eq. 2 and does not affect the *VOI*. On the other hand, since c_{draw} was not scaled along with rewards and remained the same across conditions (0.1 points), the scale manipulation affects not only the magnitude but also shape of EV_{draw} (Eq. 3) and thus the *VOI* (Eq. 2).

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The most important prediction of this theoretical framework is that *information seeking* 657 should be biased due to the reward asymmetry. The VOI takes an inverted-U shape as 658 659 a function of the beads difference, and its peak is at a moderate negative beads difference $(n_H - n_L = -5)$. This is because the information would directly improve the 660 subsequent bet choice; when $n_H - n_L = -5$, $EV(a_H | n_H, n_L)$ is close to $EV(a_L | n_H, n_L)$, but 661 the next bead would increase their difference in either direction (if a high-reward bead 662 b_H is observed, $EV(a_H|n_H + 1, n_L) > EV(a_L|n_H + 1, n_L)$; if a low-reward bead b_L is 663 664 observed, $EV(a_H \mid n_H, n_L + 1) < EV(a_L \mid n_H, n_L + 1)$). Therefore, the agent can bet on s_H after b_H and bet on s_L after b_L , and such flexibility improves the overall EV. In contrast, 665 the VOI is effectively zero when the beads difference is positive $(n_H - n_L > 0)$, because 666 the agent would bet on s_H irrespective of the next draw. The VOI is also effectively zero 667 when low-reward beads outnumber high-reward beads by a large enough margin $(n_H - n_H)$ 668 $n_L < -7$), because the agent would bet on s_L irrespective of the next draw. 669

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This gualitative prediction, a bias in information seeking towards a negative beads 671 difference, does not depend on most of our assumptions (e.g., choice optimality, risk 672 neutrality). Information seeking would be biased as far as the agent is sensitive to the 673 rank order of rewards and the bead difference. On the other hand, if an agent is not 674 motivated to maximize rewards but to maximize the accuracy of the prediction (i.e., their 675 utility function U follows $U(R_H) = U(R_L) > U(R_I)$, they would exhibit unbiased 676 information seeking; the uncertainty about the jar type is determined by $|n_H - n_L|$ and is 677 highest when $n_H = n_L$, which is when the agent would draw beads most frequently. 678 Therefore, a bias in information seeking would suggest that information seeking is 679

680 motivated by information's instrumentality for future reward seeking.

681 682 Behavioral data analysis In order to examine information-seeking behavior, we analyzed the frequency at which participants drew at least one bead as a function of the 683 beads difference. We specifically focused on whether they drew the first bead as a 684 function of the current evidence and examined if it was biased by the reward asymmetry 685 as theoretically predicted. The relationship between information-seeking behavior and 686 687 the beads difference was analyzed using Gaussian Process (GP) logistic regression (Rasmussen & Williams, 2006). GP logistic regression estimates a latent function that 688 smoothly varies with the independent variable (the beads difference) and yields 689 likelihoods of binary choices (whether participants drew a bead in each trial), and the 690 estimated latent function can be interpreted as the subjective VOI function (the higher 691 *VOI* is, the more likely participants draw a bead). The latent function with isotropic 692 squared exponential covariance was estimated using Variational Bayes approximation. 693 as implemented in Gaussian Processes for Machine Learning toolbox, version 4.2 694 695 (https://github.com/alshedivat/gpml) (Rasmussen & Nickisch, 2010).

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To test whether information-seeking behavior systematically differed across blocks and 697 reward conditions within each block, we compared four models. Model 1 implemented 698 the theoretical prescription that information seeking is sensitive to the scale 699 700 manipulation but not to the baseline manipulation. It thus consisted of three separate 701 latent value functions, one used in all trials in the baseline block, one used in trials where $(R_H, R_L, R_I) = (70, 10, 0)$ in the scale block, and one used in trials where 702 $(R_H, R_L, R_I) = (7, 1, 0)$ in the scale block. We constructed several alternative models. 703 Model 2 postulated different value functions for reward conditions not only in the scale 704 block but also in the baseline block, one for trials where $(R_H, R_L, R_I) = (70, 10, 0)$ and 705 another for trials where $(R_H, R_I, R_I) = (170, 110, 0)$ (i.e., four value functions in total); 706 707 Model 3 postulated the lack of sensitivity to reward conditions in both blocks but a 708 separate value function for each block (i.e., two value functions in total); and Model 4 postulated one common value function for all trials in both blocks. These models were 709 compared based on leave-one-participant-out cross validation (LOPO CV) and leave-710 one-trial-out cross validation (LOTO CV). We also adopted the same analytic approach 711 712 to the bet choices, comparing the performance of Models 1-4.

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We found that Model 3 outperformed other models for both information-seeking and bet choices (see Results). To test whether information-seeking behavior was biased by the reward asymmetry, we next compared Model 3 with another model (Model 5) that assumed value functions that are symmetric with respect to the beads difference (i.e., value functions that only vary with the absolute value of beads difference). We found that Model 3 fit information-seeking behavior better than Model 5, supporting a bias in information seeking (see Results).

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The fact that Model 3 performed better than Models 1, 2, and 4 suggests that, while participants did not change their behavioral strategies based on the trial-by-trial reward

manipulation, they adapted to the different reward statistics across blocks. However, 724 725 such changes across blocks could potentially reflect time-induced behavioral changes 726 as well, such as boredom or fatigue, since all participants completed the baseline block first and the scale block second. To examine the possibility that the population-level 727 behavioral pattern was not stationary over time, we tested another model (Model 6) that 728 assumed distinct value functions between the first and second scanning runs within 729 each block (one value function for each run, four functions in total). Model 6 performed 730 731 worse than Model 3 (information-seeking choices: LOPO CV log likelihood [LL] = -1222.05 vs. -1214.73, LOTO CV LL = -1145.39 vs. -1142.25, bet choices: LOPO CV 732 = -288.55 vs. -283.56, LOTO CV LL = -266.00 vs. -265.26), suggesting that changes 733 in participants' behavior were systematically driven by reward statistics rather than time. 734 735

736 **MRI** data acquisition MRI data was collected using a Siemens (Erlangen, Germany) Trio 3T scanner with a 32-channel head coil at the University of Pennsylvania. A 3D 737 738 high-resolution anatomical image was acquired using a T1-weighted MPRAGE 739 sequence (voxel size = 0.9375 x 0.9375 x 1 mm, matrix size = 192 x 256, 160 axial slices, TI = 1100 msec, TR = 1810 msec, TE = 3.51 msec, flip angle = 9 degrees). 740 Functional images were acquired using a T2*-weighted multiband gradient echoplanar 741 imaging (EPI) sequence (voxel size = $2 \times 2 \times 2 \text{ mm}$, matrix size = 98×98 , 72 axial 742 slices with no interslice gap, 400 volumes, TR = 1500 msec, TE = 30 msec, flip angle = 743 744 45 degrees, multiband factor = 4), followed by Fieldmap images (TR = 1270 msec, TEs 745 = 5 msec and 7.46 msec, flip angle = 60 degrees).

MRI data analysis MRI data were analyzed using FSL (FMRIB Software Library,
 version 6.0) (Jenkinson et al., 2012; Smith et al., 2004). MPRAGE anatomical images
 were skull-stripped using FSL BET. EPI functional images were slice-time corrected,
 motion corrected (FSL MCFLIRT), high-pass filtered (cutoff = 90 sec), geometrically
 undistorted using Fieldmap images, registered to the MPRAGE anatomical image,
 normalized to the MNI space, and spatially smoothed (Gaussian kernel FWHM = 6 mm).

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To look for regions that represent the subjective VOI upon the initial beads presentation, 754 we ran a GLM analysis (GLM 1). The regressor of interest modeled the initial beads 755 presentation (3-second boxcar) and was parametrically modulated by the trial-by-trial 756 757 subjective VOI, which was the latent function estimated in the winning model (Model 3) of GP logistic regression on the information-seeking behavior. GLM 1 also included 758 nuisance regressors that modeled the initial beads presentation (unmodulated), the 759 extra bead presentation, and button presses. The regressors were convolved with the 760 canonical double-gamma hemodynamic response function (HRF). GLM 1 additionally 761 incorporated six head motion parameters (3 translations and 3 rotations, estimated by 762 MCFLIRT) as confound regressors. GLM 1 was run following the standard approach of 763 FSL FEAT; the GLM was first fit to BOLD signals in each run (first level) and the 764 estimated coefficients of interest were combined across runs (second level). Individual-765 level *T*-statistics were entered into the population-level inference using FSL randomise, 766 in which clusters that showed positive response to subjective VOI were defined at the 767

voxel-wise cluster-forming threshold of p < .001 and evaluated by sign-flipping permutation on cluster mass. A cluster that survived whole-brain family-wise error (FWE) corrected p < .05 is reported in Fig. 3B; another cluster that survived a more lenient threshold (p < .10) is reported in Fig. S1.

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To illustrate how the cluster's activation varied as a function of the beads difference, we ran another GLM (GLM 2) using FSL FEAT, which included a regressor for each level of beads difference separately, along with the same nuisance regressors as GLM 1. *T*statistics for each regressor of interest were then averaged across runs within each block and then averaged across all voxels in the right DLPFC cluster defined as above (Fig. 3B).

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780 Lastly, to examine how the DLPFC responds to the updating of VOI, we ran another GLM (GLM 3) using FSL FEAT to estimate the time course of signals related to the 781 782 initial VOI and the VOI updating, which were derived from Model 3 of GP logistic regression. The VOI updating was calculated as the signed difference between the 783 posterior VOI, which depends both on the initial beads and the extra bead, and the prior 784 VOI, which depends only on the initial beads (Fig. 4A). GLM 3 included three sets of 785 finite impulse response (FIR) function, one unmodulated (intercept), one parametrically 786 modulated by the initial VOI, and one parametrically modulated by the VOI updating. 787 788 These FIRs were aligned to the onset of the extra bead and sampled every 1.5 seconds 789 (equal to TR) for the total duration of 21 seconds. GLM 3 also included nuisance regressors that modeled the initial beads presentation and button presses, convolved 790 with the canonical HRF, along with head motion parameters. T-statistics of 791 parametrically modulated FIR sets were averaged across all voxels in the right DLPFC 792 cluster for each participant. Population-level inference on the updating signal was 793 conducted at the cluster level across time; clusters were defined at the event-wise 794 795 cluster-forming threshold of p < .05 and evaluated by sign-flipping permutation on cluster mass, correcting for FWE across time. 796

Supplementary Figures

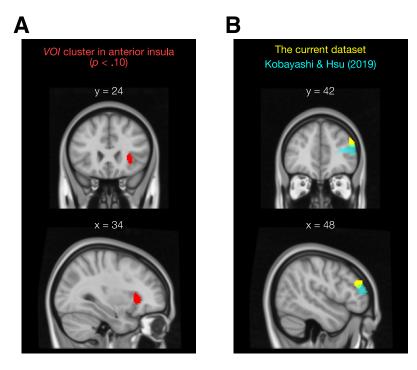


Fig. S1. (A) At a liberal threshold (cluster-forming threshold p < .001, cluster mass p < .10, corrected for whole-brain FWE), the subjective *VOI* was positively associated with activations in right anterior insula. (B) The DLPFC cluster identified in the current dataset (Fig. 4b) overlaps with a subjective *VOI* cluster reported in Kobayashi & Hsu (2019).

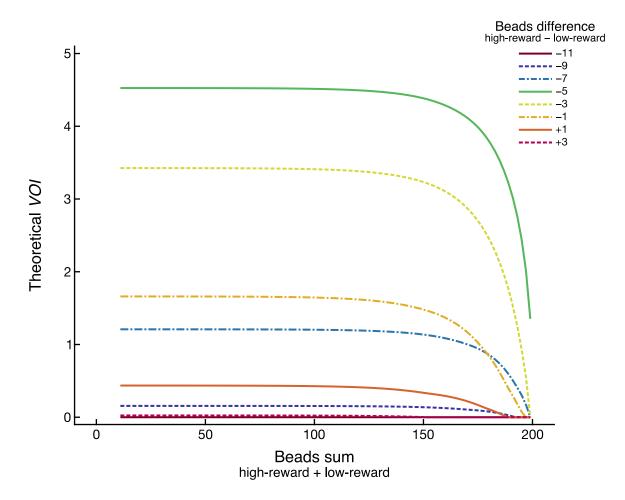


Fig. S2. The theoretical *VOI* was numerically estimated by backward recursion (up to 200 steps). The *VOI* reached an asymptote at each level of beads difference over the course of recursion. Moreover, the *VOI* was highest with a negative beads difference (–5) throughout recursion.

References

- Baker, S. C., Konova, A. B., Daw, N. D., & Horga, G. (2019). A distinct inferential mechanism for delusions in schizophrenia. *Brain*, *142*(6), 1797–1812. https://doi.org/10.1093/brain/awz051
- Barch, D. M., & Ceaser, A. (2012). Cognition in schizophrenia: core psychological and neural mechanisms. *Trends in Cognitive Sciences*, *16*(1), 27–34. https://doi.org/10.1016/j.tics.2011.11.015
- Behrens, T. E. J., Woolrich, M. W., Walton, M. E., & Rushworth, M. F. S. (2007). Learning the value of information in an uncertain world. *Nature Neuroscience*, *10*(9), 1214–1221. https://doi.org/10.1038/nn1954
- Blanchard, T. C., Hayden, B. Y., & Bromberg-Martin, E. S. (2015). Orbitofrontal cortex uses distinct codes for different choice attributes in decisions motivated by curiosity. *Neuron*, 85(3), 602–614. https://doi.org/10.1016/j.neuron.2014.12.050
- Bromberg-Martin, E. S., & Hikosaka, O. (2009). Midbrain dopamine neurons signal preference for advance information about upcoming rewards. *Neuron*, 63(1), 119–126. https://doi.org/10.1016/j.neuron.2009.06.009
- Bromberg-Martin, E. S., & Hikosaka, O. (2011). Lateral habenula neurons signal errors in the prediction of reward information. *Nature Neuroscience*, *14*(9), 1209–1216. https://doi.org/10.1038/nn.2902
- Brydevall, M., Bennett, D., Murawski, C., & Bode, S. (2018). The neural encoding of information prediction errors during non-instrumental information seeking. *Scientific Reports*, *8*(1), 6134. https://doi.org/10.1038/s41598-018-24566-x
- Caplin, A., & Leahy, J. (2001). Psychological expected utility theory and anticipatory feelings. *The Quarterly Journal of Economics*, *116*(1), 55–79. https://doi.org/10.1162/003355301556347
- Charpentier, C. J., Bromberg-Martin, E. S., & Sharot, T. (2018). Valuation of knowledge and ignorance in mesolimbic reward circuitry. *Proceedings of the National Academy of Sciences of the United States of America*, 115(31), E7255–E7264. https://doi.org/10.1073/pnas.1800547115
- Dudley, R., Taylor, P., Wickham, S., & Hutton, P. (2016). Psychosis, Delusions and the "Jumping to Conclusions" Reasoning Bias: A Systematic Review and Meta-analysis. *Schizophrenia Bulletin*, 42(3), 652–665. https://doi.org/10.1093/schbul/sbv150
- Edwards, W. (1965). Optimal strategies for seeking information: Models for statistics, choice reaction times, and human information processing. *Journal of Mathematical Psychology*, *2*(2), 312–329. https://doi.org/10.1016/0022-2496(65)90007-6
- Edwards, W., & Slovic, P. (1965). Seeking Information to Reduce the Risk of Decisions. *The American Journal of Psychology*, *78*(2), 188–197.
- Eng, G. K., Sim, K., & Chen, S.-H. A. (2015). Meta-analytic investigations of structural grey matter, executive domain-related functional activations, and white matter diffusivity in obsessive compulsive disorder: An integrative review. *Neuroscience & Biobehavioral Reviews*, *52*, 233–257. https://doi.org/10.1016/j.neubiorev.2015.03.002

- Fox, M. D., Snyder, A. Z., Vincent, J. L., Corbetta, M., Essen, D. C. V., & Raichle, M. E. (2005). The human brain is intrinsically organized into dynamic, anticorrelated functional networks. *Proceedings of the National Academy of Sciences of the United States of America*, *102*(27), 9673–9678. https://doi.org/10.1073/pnas.0504136102
- Funahashi, S., Bruce, C. J., & Goldman-Rakic, P. S. (1989). Mnemonic coding of visual space in the monkey's dorsolateral prefrontal cortex. *Journal of Neurophysiology*, *61*(2), 331–349.
- Furl, N., & Averbeck, B. B. (2011). Parietal Cortex and Insula Relate to Evidence Seeking Relevant to Reward-Related Decisions. *The Journal of Neuroscience*, *31*(48), 17572–17582. https://doi.org/10.1523/jneurosci.4236-11.2011
- Fuster, J. M., & Alexander, G. E. (1971). Neuron Activity Related to Short-Term Memory. *Science*, *173*(3997), 652–654. https://doi.org/10.1126/science.173.3997.652
- Gesiarz, F., Cahill, D., & Sharot, T. (2019). Evidence accumulation is biased by motivation: A computational account. *PLOS Computational Biology*, *15*(6), e1007089. https://doi.org/10.1371/journal.pcbi.1007089
- Gottlieb, J., & Oudeyer, P.-Y. (2018). Towards a neuroscience of active sampling and curiosity. *Nature Reviews Neuroscience*, *19*(12), 758–770. https://doi.org/10.1038/s41583-018-0078-0

Gruber, M. J., Gelman, B. D., & Ranganath, C. (2014). States of curiosity modulate hippocampusdependent learning via the dopaminergic circuit. *Neuron*, *84*(2), 486–496. https://doi.org/10.1016/j.neuron.2014.08.060

- Hauser, T. U., Moutoussis, M., Consortium, N., Dayan, P., & Dolan, R. J. (2017). Increased decision thresholds trigger extended information gathering across the compulsivity spectrum. *Translational Psychiatry*, 7(12), 1296. https://doi.org/10.1038/s41398-017-0040-3
- Howard, R. A. (1966). Information value theory. *IEEE Transactions on Systems Science and Cybernetics*, *2*(1), 22–26. https://doi.org/10.1109/tssc.1966.300074
- Huq, S. F., Garety, P. A., & Hemsley, D. R. (1988). Probabilistic Judgements in Deluded and Non-Deluded Subjects. *Quarterly Journal of Experimental Psychology*, 40(4), 801–812. https://doi.org/10.1080/14640748808402300
- Jenkinson, M., Beckmann, C. F., Behrens, T. E. J., Woolrich, M. W., & Smith, S. M. (2012). FSL. *NeuroImage*, 62(2), 782–790. https://doi.org/10.1016/j.neuroimage.2011.09.015
- Jepma, M., Verdonschot, R. G., Steenbergen, H. van, Rombouts, S. A. R. B., & Nieuwenhuis, S. (2012). Neural mechanisms underlying the induction and relief of perceptual curiosity. *Frontiers in Behavioral Neuroscience*, *6*, 5. https://doi.org/10.3389/fnbeh.2012.00005
- Kaanders, P., Nili, H., O'Reilly, J. X., & Hunt, L. T. (2020). Medial frontal cortex activity predicts information sampling in economic choice. *BioRxiv*. https://doi.org/10.1101/2020.11.24.395814
- Kakade, S., & Dayan, P. (2002). Dopamine: generalization and bonuses. *Neural Networks*, *15*(4–6), 549–559. https://doi.org/10.1016/s0893-6080(02)00048-5

- Kang, M. J., Hsu, M., Krajbich, I. M., Loewenstein, G., McClure, S. M., Wang, J. T., & Camerer, C. F. (2009). The wick in the candle of learning: Epistemic curiosity activates reward circuitry and enhances memory. *Psychological Science*, 20(8), 963–973. https://doi.org/10.1111/j.1467-9280.2009.02402.x
- Kennerley, S. W., Behrens, T. E. J., & Wallis, J. D. (2011). Double dissociation of value computations in orbitofrontal and anterior cingulate neurons. *Nature Neuroscience*, *14*(12), 1581–1589. https://doi.org/10.1038/nn.2961
- Kidd, C., & Hayden, B. Y. (2015). The psychology and neuroscience of curiosity. *Neuron*, *88*(3), 449–460. https://doi.org/10.1016/j.neuron.2015.09.010
- Kobayashi, K., & Hsu, M. (2019). Common neural code for reward and information value. Proceedings of the National Academy of Sciences of the United States of America, 116(26), 13061–13066. https://doi.org/10.1073/pnas.1820145116
- Kobayashi, K., Ravaioli, S., Baranès, A., Woodford, M., & Gottlieb, J. (2019). Diverse motives for human curiosity. *Nature Human Behaviour*, *3*(6), 587–595. https://doi.org/10.1038/s41562-019-0589-3
- Kolling, N., Behrens, T. E. J., Mars, R. B., & Rushworth, M. F. S. (2012). Neural Mechanisms of Foraging. Science, 336(6077), 95 98. https://doi.org/10.1126/science.1216930
- Krebs, R. M., Schott, B. H., Schütze, H., & Düzel, E. (2009). The novelty exploration bonus and its attentional modulation. *Neuropsychologia*, 47(11), 2272–2281. https://doi.org/10.1016/j.neuropsychologia.2009.01.015
- Kreps, D. M., & Porteus, E. L. (1978). Temporal resolution of uncertainty and dynamic choice theory. *Econometrica*, *46*(1), 185–200. https://doi.org/10.2307/1913656
- Kunda, Z. (1991). The case for motivated reasoning. *Psychological Bulletin*, *108*(3), 480–498. https://doi.org/10.1037/0033-2909.108.3.480
- Lau, J. K. L., Ozono, H., Kuratomi, K., Komiya, A., & Murayama, K. (2020). Shared striatal activity in decisions to satisfy curiosity and hunger at the risk of electric shocks. *Nature Human Behaviour*, 4(5), 531–543. https://doi.org/10.1038/s41562-020-0848-3
- Leong, Y. C., Hughes, B. L., Wang, Y., & Zaki, J. (2019). Neurocomputational mechanisms underlying motivated seeing. *Nature Human Behaviour*, *3*(42), 962–973. https://doi.org/10.1038/s41562-019-0637-z
- Loewenstein, G. (1994). The psychology of curiosity: A review and reinterpretation. *Psychological Bulletin*, *116*(1), 75–98. https://doi.org/10.1037/0033-2909.116.1.75
- Luu, L., & Stocker, A. A. (2018). Post-decision biases reveal a self-consistency principle in perceptual inference. *ELife*, *7*, E3548. https://doi.org/10.7554/elife.33334
- McGuire, J. T., Nassar, M. R., Gold, J. I., & Kable, J. W. (2014). Functionally Dissociable Influences on Learning Rate in a Dynamic Environment. *Neuron*, *84*(4), 870 881. https://doi.org/10.1016/j.neuron.2014.10.013

- Menon, V., & Uddin, L. Q. (2010). Saliency, switching, attention and control: a network model of insula function. *Brain Structure and Function*, 214(5–6), 655–667. https://doi.org/10.1007/s00429-010-0262-0
- Nassar, M. R., McGuire, J. T., Ritz, H., & Kable, J. W. (2019). Dissociable Forms of Uncertainty-Driven Representational Change Across the Human Brain. *Journal of Neuroscience*, *39*(9), 1688–1698. https://doi.org/10.1523/jneurosci.1713-18.2018
- Phillips, L. D., & Edwards, W. (1966). Conservatism in a simple probability inference task. *Journal of Experimental Psychology*, *72*(3), 346–354. https://doi.org/10.1037/h0023653
- Rasmussen, C. E., & Nickisch, H. (2010). Gaussian Processes for Machine Learning (GPML) Toolbox. *Journal of Machine Learning Research*, *11*(100), 3011–3015.
- Rasmussen, C. E., & Williams, C. K. I. (2006). *Gaussian Processes for Machine Learning*. The MIT press. https://doi.org/10.7551/mitpress/3206.001.0001
- Ross, R. M., McKay, R., Coltheart, M., & Langdon, R. (2015). Jumping to Conclusions About the Beads Task? A Meta-analysis of Delusional Ideation and Data-Gathering. *Schizophrenia Bulletin*, 41(5), 1183–1191. https://doi.org/10.1093/schbul/sbu187
- Rudebeck, P. H., Behrens, T. E., Kennerley, S. W., Baxter, M. G., Buckley, M. J., Walton, M. E., & Rushworth, M. F. S. (2008). Frontal cortex subregions play distinct roles in choices between actions and stimuli. *The Journal of Neuroscience*, *28*(51), 13775–13785. https://doi.org/10.1523/jneurosci.3541-08.2008
- Rushworth, M. F. S., & Behrens, T. E. J. (2008). Choice, uncertainty and value in prefrontal and cingulate cortex. *Nature Neuroscience*, *11*(4), 389 397. https://doi.org/10.1038/nn2066
- Seeley, W. W., Menon, V., Schatzberg, A. F., Keller, J., Glover, G. H., Kenna, H., Reiss, A. L., & Greicius, M. D. (2007). Dissociable Intrinsic Connectivity Networks for Salience Processing and Executive Control. *The Journal of Neuroscience*, *27*(9), 2349–2356. https://doi.org/10.1523/jneurosci.5587-06.2007
- Shadlen, M. N., & Shohamy, D. (2016). Decision Making and Sequential Sampling from Memory. *Neuron*, *90*(5), 927–939. https://doi.org/10.1016/j.neuron.2016.04.036
- Shanteau, J., & Anderson, N. H. (1972). Integration theory applied to judgments of the value of information. *Journal of Experimental Psychology*, *92*(2), 266–275. https://doi.org/10.1037/h0032079
- Sharot, T., & Garrett, N. (2016). Forming Beliefs: Why Valence Matters. *Trends in Cognitive Sciences*, 20(1), 25–33. https://doi.org/10.1016/j.tics.2015.11.002
- Sharot, T., & Sunstein, C. R. (2020). How people decide what they want to know. *Nature Human Behaviour*, *4*(1), 14–19. https://doi.org/10.1038/s41562-019-0793-1
- Shenhav, A., Cohen, J. D., & Botvinick, M. M. (2016). Dorsal anterior cingulate cortex and the value of control. *Nature Neuroscience*, *19*(10), 1286–1291. https://doi.org/10.1038/nn.4384

- Shenhav, A., Straccia, M. A., Cohen, J. D., & Botvinick, M. M. (2014). Anterior cingulate engagement in a foraging context reflects choice difficulty, not foraging value. *Nature Neuroscience*, 17(9), 1249 1254. https://doi.org/10.1038/nn.3771
- Smith, S. M., Jenkinson, M., Woolrich, M. W., Beckmann, C. F., Behrens, T. E. J., Johansen-Berg, H., Bannister, P. R., Luca, M. D., Drobnjak, I., Flitney, D. E., Niazy, R. K., Saunders, J., Vickers, J., Zhang, Y., Stefano, N. D., Brady, J. M., & Matthews, P. M. (2004). Advances in functional and structural MR image analysis and implementation as FSL. *NeuroImage*, *23*, S208–S219. https://doi.org/10.1016/j.neuroimage.2004.07.051
- Sreenivasan, K. K., & D'Esposito, M. (2019). The what, where and how of delay activity. *Nature Reviews Neuroscience*, *20*(8), 466–481. https://doi.org/10.1038/s41583-019-0176-7
- Talluri, B. C., Urai, A. E., Tsetsos, K., Usher, M., & Donner, T. H. (2018). Confirmation bias through selective overweighting of choice-consistent evidence. *Current Biology*, 28(19), 3128–3135. https://doi.org/10.1016/j.cub.2018.07.052
- Wendt, D. (1969). Value of information for decisions. *Journal of Mathematical Psychology*, *6*(3), 430–443. https://doi.org/10.1016/0022-2496(69)90015-7
- White, J. K., Bromberg-Martin, E. S., Heilbronner, S. R., Zhang, K., Pai, J., Haber, S. N., & Monosov, I. E. (2019). A neural network for information seeking. *Nature Communications*, *10*(1), 5168. https://doi.org/10.1038/s41467-019-13135-z
- Wilson, R. C., Geana, A., White, J. M., Ludvig, E. A., & Cohen, J. D. (2014). Humans use directed and random exploration to solve the explore-exploit dilemma. *Journal of Experimental Psychology: General*, 143(6), 2074–2081. https://doi.org/10.1037/a0038199